### NATIONAL QUALITY FORUM

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# MEASURE APPLICATIONS PARTNERSHIP COORDINATING COMMITTEE

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# THURSDAY JANUARY 25, 2018

The Committee met at the National Quality Forum, 9th Floor Conference Room, 1030 15th Street, N.W., Washington, D.C., at 9:00 a.m., Charles Kahn III and Harold Pincus, Co-Chairs, presiding.

#### PRESENT:

CHARLES KAHN III, MPH, Co-Chair

HAROLD PINCUS, MD, Co-Chair

RICHARD ANTONELLI, MD, MS, Individual Subject Matter Expert\*

JOE BAKER, Medicare Rights Center

LEAH BINDER, MA, MGA, The Leapfrog Group

JOHN BOTT, MSSW, MBA, Consumers Union

MARY BETH BRESCH WHITE, American Nurses
Association

RAJESH DAVDA, MD, America's Health Insurance Plans

TRICIA ELLIOTT, MBA, CPHQ, The Joint Commission DAVID GIFFORD, MD, MPH, American Health Care Association

BRUCE HALL, MD, PhD, MBA, FACS, American College of Surgeons

GAIL HUNT, National Alliance for Caregiving DAVID INTROCASO, PhD, American Medical Group Association

MAUREEN KAHN, RN, MHA, MSN, Blessing Health System\*

MIRA IRONS, MD, American Board of Medical Specialties\*

- WILLIAM KRAMER, MBA, Pacific Business Group on Health
- RACHEL LA CROIX, PhD, PMP, National Association of Medicaid Directors
- SAMUEL LIN, MD, PhD, MBA, MPA, MS, American Medical Group Association
- ERIN MACKAY, MPH, National Partnership for Women & Families
- AMY MULLINS, MD, FAAFP, American Academy of Family Physicians
- SHAUN O'BRIEN, JD, AFL-CIO
- AMIR QASEEM, MD, PhD, MHA, FACP, American College of Physicians
- CHRIS QUERAM, MS, Network for Regional Healthcare Improvement
- DEREK ROBINSON, MD, MBA, FACEP, CHCQM, Health Care Service Corporation
- MARISSA SCHLAIFER, MS, RPh, Academy of Managed Care Pharmacy
- CARL SIRIO, MD, American Medical Association STEVE WOJCIK, MA, National Business Group on Health

FEDERAL GOVERNMENT LIAISONS (NON-VOTING):

- ROB ANTHONY, Office of the National Coordinator for Health Information Technology (ONC)
- MARY BARTON, MD, Agency for Healthcare

  Research and Quality (AHRQ)
- KATE GOODRICH, MD, MHS, Centers for Medicare and Medicaid Services (CMS)

## NQF STAFF:

TAROOQ AMIN, PhD, MPH, Consultant
JOHN BERNOT, Senior Director
KATHRYN BUCHANAN, Project Manager
KAREN JOHNSON, MS, Senior Director, Performance
Measures

MELISSA MARINELARENA, Senior Director
KATE McQUESTON, Project Manager
ELISA MUNTHALI, MPH, Acting Senior Vice
President, Quality Measurement
YETUNDE OGUNGBEMI, Project Analyst
ERIN O'ROURKE, Senior Director
JEAN-LUC TILLY, Senior Project Manager, Data
Analytics

### ALSO PRESENT:

BRUCE BAGLEY, MD, Clinician Workgroup Co-Chair\* ROSE DO, MD, Acumen\*

GERRI LAMB, PhD, RN, FAAN, PAC/LTC Workgroup Co-Chair\*

AMY MOYER, Clinician Workgroup Co-Chair\*

PAUL MULHAUSEN, MD, FACP, PAC/LTC Workgroup Co-Chair\*

SRI NAGAVARAPU, PhD, Acumen\*

COLLETTE PITZEN, RN, MSN, Minnesota Community
Measurement\*

JESSE ROACH, MD, Centers for Medicare and Medicaid Services (CMS)

KORYN RUBIN, American Medical Association STEPHANIE GLIER, MPH, Pacific Business Group on Health\*

CRISTIE TRAVIS, Hospital Workgroup Co-Chair\*
RON WALTERS, MD, MBA, MHA, Hospital Workgroup
Co-Chair\*

PIERRE YONG, MD, MPH, MS, Centers for Medicare and Medicaid Services (CMS)

<sup>\*</sup> present by teleconference

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1	P-R-O-C-E-E-D-I-N-G-S
2	9:16 a.m.
3	CO-CHAIR KAHN: It is the witching
4	hour. It's 9:15, so we're going to start, so if
5	everybody would take a seat and Harold and I will
6	take off here as our chairs today.
7	And I want to thank everybody for
8	coming and thank everyone for all the time and
9	effort that you put in, both on the Coordinating
10	Committee and some of you were on the workgroups,
11	or work with the workgroups, and thank you for
12	your time and effort there.
13	It's what makes NQF and our process
14	work, and also thank you to the CMS people for
15	allowing us to work with you on these really
16	important issues.
17	I think I should give it to you then,
18	and then I do the objectives. Do you have
19	anything to say?
20	CO-CHAIR PINCUS: Just I totally agree

with everything Chip says.

CO-CHAIR KAHN:

21

22

Okay, so I'll review

the objectives today. So, our primary objective is to finalize our recommendations to HHS on the measures for use in Federal programs for clinician, hospital, post-acute long-term care settings.

We're also going to consider strategic issues that span across the MAP workgroups and discuss potential improvements to the way we interact with the pre-rulemaking process.

I know that we had a lot of discussions on the phone about the different categories.

And I know from one of the workgroups there were process issues regarding -- it seems like it's a proverbial -- I mean it's the constant discussion in my experience with NQF as to what is a consensus?

So, we'll probably have some discussion about that. I guess one man's consensus is not another person's consensus -- or I should say persons, persons.

So, with that, I guess I'll turn it

over to Elisa to conduct the questions about our disclosures of interest, which we'll need to review before we begin.

MS. O'ROURKE: I actually just want to make a few housekeeping announcements before turning it to Elisa.

If you're having any trouble accessing materials from the NQF website, we do have a public SharePoint site with all of the materials for today's meeting at public.qualityforum.org.

You'll see a little folder for MAP and then MAP Coordinating Committee. So, any materials you might need you can access that way.

And for anyone on the phone, if you're having trouble getting an open line, please just let us know through the webchat, or email the MAP Coordinating Committee at qualityforum.org email, and we will work with our operator to get your line open.

So, please let us know if you're having any trouble speaking. I think, otherwise, I can turn it to Elisa to take you through

disclosures.

MS. MUNTHALI: Great, thank you, Erin, and welcome, everyone.

Good morning, my name is Elisa

Munthali, and I'm the Senior Vice President for

Quality Measurement at the National Quality

Forum.

So, I welcome you and thank you so much for serving on this Committee.

We're going to combine the disclosures of interest with introductions, and we're going to do this in two parts because there are two types of members on this Committee. There are organizational representatives and there are also subject-matter experts. And so we'll start with the organizational members.

Organizational members represent a particular organization perspective, and we expect you to come to the table sharing those views. Because of your status as an organizational member, we asked you just one question about you as an individual.

We asked you if you had an interest of \$10,000 or more in an entity that is related to the work that's in front of you. And so how we're going to do this today is go around the table. First, we'll start off with the organizational members.

It's a little easier today because we only have three subject-matter experts. Those are Harold and Chip and Rich Antonelli, who's on the phone.

And so we'll start with our organizational members. Sorry, Chip was trying to figure out: how did he become a subject-matter expert?

(Laughter.)

MS. MUNTHALI: So, we're going to start with all the organizational members.

We're going to go clockwise, so to my left, in the room first, then we'll go to the folks on the phone that are organizational members, then go to our subject-matter experts, and then welcome our federal partners.

1	And so I think, David, you might be
2	the first organizational member here?
3	MR. INTROCASO: So, I'm going to just
4	introduce myself, David Introcaso, AMGA, and I
5	have no conflicts.
6	MS. MUNTHALI: Perfect, just like
7	that.
8	MR. KRAMER: I'm Bill Kramer,
9	representing Pacific Business Group on Health.
10	No conflicts to report.
11	MS. MULLINS: Amy Mullins, American
12	Academy of Family Physicians. No conflicts.
13	MR. BAKER: I'm Joe Baker,
14	representing the Medicare Rights Center, and I
15	have no conflicts.
16	MR. HALL: Bruce Hall, representing
17	the American College of Surgeons.
18	Other than income from American
19	College of Surgeons and my healthcare hospital
20	chain, BJC Healthcare, I have no conflicts of
21	interest.
22	MS. HUNT: Gail Hunt, representing the

1	National Alliance for Caregiving, and no
2	conflicts of interest.
3	MR. BOTT: John Bott with Consumer
4	Reports, and no conflicts of interest.
5	MR. QUERAM: Morning, I'm Chris
6	Queram, representing the Network for Regional
7	Healthcare Improvement. No conflicts.
8	MR. O'BRIEN: Shaun O'Brien,
9	representing the AFL-CIO. No conflicts.
10	MS. BINDER: Leah Binder from the
11	Leapfrog Group. No conflicts.
12	MR. ROBINSON: Good morning, Derek
13	Robinson, Healthcare Service Corporation. No
14	conflicts.
15	MR. GIFFORD: David Gifford
16	representing the American Healthcare Association.
17	I am extremely conflicted. I actually have to
18	recuse myself on one of the discussions today for
19	a measure steward of a measure coming forward to
20	you all.
21	Second, if there's any discussion of
22	Medicaid-related measures, I have a family member
J	

1 involved in Medicaid so I have to recuse myself 2 personally, but I have my alternate who would step in in that case. 3 And then all my retirement accounts 4 5 are in 401Ks, I have no idea what they're 6 invested in, but I know some are in healthcare so 7 I'm sure there's some conflicts there. 8 MS. ELLIOTT: Good morning, I'm Tricia 9 Elliott, representing the Joint Commission, and I have no conflicts. 10 11 MR. SIRIO: Hi, good morning, Carl 12 Sirio, American Medical Association. 13 conflicts. 14 MS. SCHLAIFER: Marissa Schlaifer, representing the Academy of Managed Care 15 16 Pharmacy, and as a formal CVS Health employee, I 17 have CVS Health stock. 18 MR. DAVDA: Raj Davda, I'm employed by 19 Cigna Healthcare. I'm representing AHIP. I have 20 no conflicts. 21 MS. LA CROIX: Rachel La Croix, and I'm representing the National Association of 22

Medicaid Directors, and I have no conflicts. 1 2 MR. QASEEM: Amir Qaseem, American College of Physicians. I mean, you have my 3 disclosures. You guys need to decide if I have 4 5 conflicts or not. I think it should be the other way around. 6 7 (Laughter.) 8 Erin Mackay, representing MS. MACKAY: 9 the National Partnership for Women and Families. I have no conflicts. 10 11 MS. MUNTHALI: Thank you to all of our 12 organizational members in the room. So, now 13 we'll go to the phone and I'll call your name out 14 as it appears on the roster. So, do we have a representative from 15 16 the American Board of Medical Specialties? 17 MS. IRONS: Yes, this is Mira Irons. 18 I'm representing the American Board of Medical 19 Specialties, and I'm an employee of ABMS, but no conflicts otherwise. 20 Thank you, Mira. 21 MS. MUNTHALI: Do we 22 have a representative from Blessing Health

1 System? Maureen, by any chance are you on the 2 phone? Okay, we'll come back to that. Rich, are you on the phone? 3 4 you're an SME, so we'll skip you for now. 5 And finally, the National Business Group on Health? 6 7 MR. WOJCIK: Yes, hi, this is Steve 8 Wojcik, representing National Business Group on 9 Health. No conflicts. 10 MS. MUNTHALI: Great, thank you so 11 much. And so now we'll go to our subject-matter 12 experts. 13 And for everyone who's a subject-14 matter expert -- it's your co-chairs and Rich, 15 who's on the phone -- you received a lengthier 16 Disclosure of Interest form. 17 And in that form, we asked you to 18 supply us with all of the professional activities 19 that were relevant to the work in front of you. 20 So, we don't want you to recite your 21 resumes, but what we want you to do is let us 22 know if there is anything that you think is

1 important or relevant to the work of the 2 Coordinating Committee. Just a couple of reminders, you sit on 3 4 this group as an individual. You do not 5 represent your employer or anybody who may have 6 nominated you for the Committee. We're also interested in disclosures 7 8 of interest that are paid and unpaid, so things 9 that you may have volunteered for, like this Committee. 10 11 Serving on the Committee would be an 12 example of that, and we wanted to remind you that 13 just because you disclose does not mean you have a conflict of interest. 14 We do this in the spirit of 15 16 transparency and openness. And so I'll start off 17 with Chip? 18 CO-CHAIR KAHN: Well, I work at the 19 Federation of American Hospitals, and so on other 20 issues, I represent hospitals. 21 So, I guess that, in a sense, is the 22 major area. Other than that, I don't think

there's anything significant. 1 2 MR. PINCUS: Hi, Harold Pincus, I also am employed by Columbia University and also the 3 4 New York Presbyterian Hospital. 5 I'm also an adjunct staff at RAND, a consultant for Mathematica, I'm on the American 6 Psychiatric Associations Quality Counsel, and 7 8 also I'm on the NCQA Behavioral Health Advisory 9 Committee. 10 MS. MUNTHALI: Thank you both. And so 11 we'll go to the phone to Rich Antonelli? 12 are you on the phone? Okay, he may have not 13 joined us yet. 14 So, thank you so much for disclosing your conflicts of interest. We're going to now 15 16 ask our federal partners to introduce themselves. 17 So, we'll start with Kate? 18 MS. GOODRICH: Hi, everyone, Kate 19 Goodrich, Centers for Medicare and Medicaid 20 Services. I'm Chief Medical Officer and Director 21 22 of the Center for Clinical Standards and Quality.

I will just disclose that I am a 1 2 member of the NCQA Committee on performance I don't think that's a conflict but 3 measurement. I will disclose it. 4 5 In other ways, I have no disclosures. Hi, Pierre Yong, CMS as 6 MR. YONG: well; I work with Kate. And no disclosures, 7 thank you. 8 9 MS. MUNTHALI: Great, do we have any other Federal Liaisons in the room? 10 11 MR. ANTHONY: Rob Anthony, 12 representing the Office of the National Coordinator for Health Information Technology. 13 14 I have nothing to disclose. 15 Thank you very much. MS. MUNTHALI: 16 And so before I leave or we leave this 17 session, I just wanted to remind you if you have 18 a conflict or you remember something in the middle of the meeting, please speak up. 19 You can do so in real time or you can 20 21 come to any one of your chairs or approach the 22 NQF staff, or you can go directly to the staff.

1	Likewise, if you feel that any one of
2	your colleagues has a conflict or is acting in a
3	biased manner, we want you to speak up as well.
4	And you can approach the co-chairs or any of the
5	staff.
6	So, after hearing the conflicts of
7	interest that have been disclosed, do you have
8	any questions of me or of your colleagues?
9	It doesn't sound like it, so thank you
10	very much.
11	Okay, Maureen, are you on the phone
12	now?
13	MEMBER KAHN: Yes, I am. When you
14	called me, somehow I was disconnected.
15	MS. MUNTHALI: Oh, sorry about that.
16	MEMBER KAHN: So, I'm back. Not a
17	problem, but I'm back, and I have no conflicts.
18	MS. MUNTHALI: Great, thank you. And
19	just a reminder, Maureen is an organizational
20	rep. So, thank you, everyone.
21	CO-CHAIR KAHN: Boy, I was so caught
22	up in disclosures.

Okay, the NQF staff and MAP workgroup co-chairs will outline the measures and the programs evaluated by the workgroups.

That's going to be our task over today, including the top strategic issues that emerged from the year's pre-rulemaking meetings.

The MAP co-chairs will identify the pre-pulled measures, and ask the Coordinating Committee members if there are any additional individual measures that they need to be pulled for discussion before ratification of the workgroup recommendation.

So, we'll be able to cover every measure that anyone has an issue or question or problem with. And we'll emphasize that the workgroups are populated with experts relevant to their care settings, and that the Coordinating Committee should express a strong preference for the workgroups' initial recommendations, which we generally have in the past and I assume would do today.

If a Coordinating Committee member

would like to pull a measure for discussion, they must clearly identify which part of the workgroup recommendation they have disagreement with to begin discussion.

And I think the word "disagreement" -I think we can just use the word "issue", or
"question", because there are really a number of,
there are an array of, recommendations from the
workgroups for these measures. All other
measures will be considered ratified by the MAP.

So, we're obviously focusing on measures of disagreement or question, but as I said, I don't think people should be shy about bringing up a measure, even if they don't have a disagreement but they have a question.

Because this is an important process.

Obviously, CMS is here and will listen to the discussion or answer questions for us.

And part of this, I think, is clarifying, as well as potentially correcting or changing or revising.

CO-CHAIR PINCUS: Let me just add one

thing. We had a discussion about this earlier.

The term "pull" has sometimes confused people.

And in some ways there's two types of pull. If you're pulling because you disagree with it and you actually want to have a vote on it, let us know that's your intent.

On the other hand, if you just have a question about it, you don't need to pull it and have us re-vote.

So, it would be important to distinguish between, if you do want to bring a measure up for discussion, whether you really are intending for it to be a re-vote, or actually a vote, that would be outside of the consent calendar, versus just wanting to clarify something.

CO-CHAIR KAHN: Yes, I think this is important. Clearly, we -- I don't remember whether at the beginning we did, because I've been on the Coordinating Committee the whole time, whether we at one point actually went through everything.

1	We couldn't. It was hard then, now we
2	just can't go through everything. So, obviously,
3	we're going to focus on the issues that you want
4	us to focus on. But I'll just repeat: pulled
5	measures will be discussed and/or voted on.
6	So, Harold, I'll pass it on to you?
7	CO-CHAIR PINCUS: So, anyways, I'll
8	just say if you do want to pull something, quote,
9	pull, let us know what's your intent.
10	CO-CHAIR KAHN: So, now you introduce
11	
12	CO-CHAIR PINCUS: Oh, of course. So,
13	we have here Kate Goodrich, who really is
14	somebody who's really taken the lead from CMS for
15	putting all of this together, and putting
16	together the MUC list.
17	And, Kate, will you sort of bring us
18	up to date?
19	MEMBER GOODRICH: Absolutely, I don't
20	think I'll take the full 15 minutes allotted to
21	me on the agenda.
22	So, first, just want to say very, very

good to be here with you guys. I look around the room, and I've known many of you for the last six or seven years -- mostly seeing you in this room or in the old building.

And it's always really fun to be here this time of year to be talking about our measurement programs and the measures that we put in front of the Committees.

So, just very excited to be here and talk about the measures today.

I do want to emphasize -- especially for folks who are maybe new to the MAP Committee or haven't been on the Committee as long -- I want to just really be sure that I say and emphasize the importance of this work to CMS.

This is the sixth or seventh year we've been doing this. We definitely had some fits and starts the first couple of years, as we were learning together about how to do this well and what it all meant, and how we use the input.

It is now an integral part of our work that, for us, is every bit as important as the

rest of the measure development and 1 2 implementation lifecycle, including putting measures out for public comment. 3 Your input to us in getting that 4 multi-stakeholder input on measures for our 5 programs is as important as it has ever been. 6 7 This year, if you have been on the 8 Committees in the past or have been following 9 along with the work of the MAP over the past seven years, you will see that there are fewer 10 11 measures on the list that went forward to the 12 three Committees this year than ever before. And there's a couple of reasons for 13 14 that. I think you were harkening back, Chip, to the days when we had, I forget, it was like 500? 15 16 I don't know; I've probably inflated 17 it in my mind. 18 CO-CHAIR PINCUS: It was like 300. 19 MEMBER GOODRICH: But there was one 20 year where we literally had multiple hundreds of

measures, which is just insane. There were

reasons for that, but we are past that time.

21

But the reason you see fewer this year is really two main reasons, I would say. The first really just has to do with where we at CMS are in our measure-development lifecycle.

So, a lot of measures that came up over the last two or three years were a result of multi-year contracts we had to develop measures.

So, we always knew that over time, as our measures kind of went through the review process, there would come a time where we'd have fewer, just by the nature of the lifecycles that we have in our contracting cycles for measure development. So, that's one reason.

But, really, the main reason that you were seeing fewer measures this year -- and this credit goes entirely to the person to my left,
Dr. Yong -- is because Pierre and his team really thought hard about the experiences that we and you have had with this process over the last several years, the kinds of measures we brought forward, at sort of varying times in their lifecycles, and really what the MAP needed to be

able to make a robust, informed recommendation for CMS.

And also, with the development of the Meaningful Measures Framework that I'm sure you all have heard us talk about a number of times, and I know Pierre presented on a webinar to this Committee last year, we really, very genuinely looked at every single measure through that lens.

And also said, you know, it's got to be far enough along that the Committee can truly have enough information to be able to weigh in on this measure in a meaningful way.

And that actually led to a significant number of measures not going on the list this year, in great part because a lot of them just weren't ready.

We may see a number of those come through next year as their development continues.

But we felt like we tried -- I think
we experimented over the last couple of years
with bringing measures forward that maybe weren't
as far along, to get input from the MAP as to

whether or not that measure was in the right direction for us.

But what we found, I think, was that people just felt like they just didn't have enough information, and so we decided to sort of step back and take a little bit of a different approach.

I also want to acknowledge that, obviously, this is the first MAP meeting in a new administration with some new priorities.

And so CMS remains very committed to the move to paying for value, and as we do that with the new administration, we do have some new priorities.

And that is very reflected in the Meaningful Measures Framework, where our direction has been to really focus on the areas of quality of care that really matter the most.

Which really does mean not only, I think, honing the measures within each program, but for our own measure development, the signals we give to other measure-developers and other

agencies about where we think are the things that we really should be measuring.

So, for our ongoing and future development, it's really focusing on those key areas.

Another big initiative that many of you have been involved with or at least heard us talk a lot about is our Patients Over Paperwork initiative, which is really around reducing the burden on a variety of fronts, certainly not just related to quality measurement, but the burden of actually the work that's required to report on quality measures.

And that is with a recognition that for the really important areas that are part of meaningful measures, sometimes it's worth some of that burden to collect data if it's on a measure that actually can help drive improved patient outcomes.

So, we think about Patients Over
Paperwork as burden reduction in the service of
patient safety, better patient outcomes, and

program integrity.

So, it's been actually, for me personally, a tremendous opportunity to sit back and take another look at everything that we are doing across the board, and making sure that what we are asking of clinicians and providers at facility types and dialysis facilities and so forth is that it really is going to drive improvement for patients.

So, I think that when you look at the list of measures, a number that are being brought here today, but if you look at the whole list that came forward to all three committees, you will see that that is reflected in the list that came forward.

And so finally, the last thing I just want to say, since I have the mic, is just to thank the CMS staff, Pierre and his team, for the incredible amount of work that goes into preparing for these series of meetings, and the thoughtfulness with which the staff undertook preparing the Measures Under Consideration list

this year.

And then finally, I want to thank the NQF staff, because as usual, you guys have done a spectacular job at getting everybody ready.

The thoughtfulness that you all provide into this process, working collaboratively with our team, always through the spirit of continuous improvement, as we will talk about tomorrow more, is very, very much appreciated by me personally and by the administration.

So, thank you so much.

CO-CHAIR KAHN: Kate, you're going to walk us through the process?

MS. BUCHANAN: I am. So, let me just pull up the slides, one moment. Hi, everyone, my name is Kate Buchanan.

I'm a Senior Project Manager here working with the Coordinating Committee. So, we wanted to take a couple minutes and review the MAP pre-rulemaking approach. So, the approach to the analysis and selection of measures is a

three-step process.

The first is to provide a program overview, the second is to review current measures within the programs, and the third is to evaluate the Measures Under Consideration -- also known as MUCs -- for what they would add to the program measure set.

MAP workgroups must reach a decision about every measure under consideration, and as you all know, the MAP workgroups met in December of this year -- or last year, December 2017 -- to review the current MUCs.

The decision categories are standardized for consistency across measures and programs, and each decision was provided with one or more statements of rationale.

To facilitate MAP's consent calendar voting process, NQF staff working with the workgroups conducted a preliminary analysis of each measure under consideration. The preliminary analysis is intended to provide MAP members with a succinct profile of each measure,

and to serve as a starting point for MAP discussions.

Staff used an algorithm developed by the MAP measure-selection criteria to evaluate each measure. The preliminary analysis algorithm uses a series of criteria to determine if a measure receives a recommendation of support for rulemaking, conditional support for rulemaking, refine and resubmit prior to rulemaking, or a donot-support designation.

And here on this slide, you can see
the MAP measure-selection criteria, and the
measure-selection criteria are intended to assist
the MAP with identifying characteristics that are
associated with ideal measure sets used for
public reporting and payment programs.

It's important to note that the measure-selection criteria are not absolute rules. Rather, they are meant to provide general guidance on measure-selection decisions and to complement program-specific statutory and regulatory requirements.

The central focus should be on the selection of high-quality measures that optimally address the national quality strategy's three aims, fill critical measurement gaps, and increase alignment.

Although competing priorities often need to be weighed against one another, the measure-selection criteria can be used as a reference when evaluating the relative strengths and weaknesses of a program measure set, and how the addition of an individual measure would contribute to the set.

As discussed previously, there are four voting decision categories. They are: support for rulemaking, conditional support for rulemaking, refine and resubmit prior to rulemaking, and do-not-support for rulemaking.

The MAP may support a measure for rulemaking for a number of reasons, including that it may address a previously-identified gap in a program or help to promote alignment.

The MAP may conditionally support a

measure if the group thinks it is ready for rulemaking but needs NQF endorsement.

And the refine and resubmit, which the MAP implemented to allow a way to express its support for the concept of a measure, but to stipulate that it needs modifications, such as testing before implementation.

The MAP may not support a measure for rulemaking if it overlaps with existing measures, or if a different measure better addresses the needs of a program.

During the Fall 2017 workgroup meetings, workgroup members raised concerns about the refine and resubmit category.

You all, as the Coordinating

Committee, created this category with the thought

that MUCs receiving this designation will be

brought back to the MAP before implementation.

But it is important to note that the HHS Secretary has statutory authority to propose measures after considering MAP's recommendations.

And tomorrow the Coordinating

Committee will review this decision category and decide if they would recommend changes for the upcoming cycles.

CO-CHAIR PINCUS: But it's important to note that for this process today, we're going to go with the current way which it's been interpreted in the state.

MS. BUCHANAN: Yes, thank you. That's very important.

And so we thought that it would be helpful as a reminder to review the guidance that you as the Coordinating Committee made during your November 30th, 2017 meeting in reaction to the concerns raised.

And you all reiterated the intent of the decision was to support the concept of a measure but recognize a potentially significant issue that should be addressed.

The Committee suggested that this category be used judiciously -- for example, when a measure needs a substantive change. And the Committee also noted the need for workgroups to

1	clarify the suggested refinements to the
2	measures.
3	So, with that, we will now discussion
4	the voting instructions.
5	CO-CHAIR KAHN: May I ask a question
6	before you do?
7	MS. BUCHANAN: Of course.
8	CO-CHAIR KAHN: In terms of the
9	recommendations from the staff, and I didn't
10	think about this until you were just reviewing
11	it, on the consent calendar of the
12	recommendations, how many do-not-supports came
13	out of the algorithm process?
14	MS. BUCHANAN: That is a very good
15	question. I don't know if I have the number off
16	the top of my head.
17	Erin, do you happen to know? Oh, we
18	need to turn one mic off.
19	CO-CHAIR KAHN: Were there any?
20	DR. AMIN: So, just a quick question,
21	are you asking what the staff preliminary
22	recommendations were before the workgroups met or

1	after the workgroups met?
2	CO-CHAIR KAHN: Before the workgroups,
3	because the algorithm is run on all the measures.
4	DR. AMIN: Right, that's true.
5	CO-CHAIR KAHN: And then put on the
6	consent calendar that the workgroup considers, if
7	I understand the process?
8	DR. AMIN: Yes, that's correct.
9	CO-CHAIR KAHN: So, what I'm asking
10	is, in terms of that algorithm, how many do-not-
11	supports, if any, were produced from the
12	algorithm process?
13	MS. O'ROURKE: I'm looking to John in
14	the background for some of the clinician
15	measures. I think there may have been one across
16	the algorithm. So, I think that's something we
17	can look at tomorrow.
18	Katie, if you go back to the slide
19	that had the decision categories, there's some
20	information here of how you end up,
21	algorithmically, into the do-not-support.
22	And it's generally when you fail one

of the first three assessments, or when a measure fails one of the first three assessments.

And those right now really ask questions about the importance of the measure and if it's addressing a quality problem.

So, after that, we go into the other three categories for the workgroup members on the Coordinating Committee to consider if there's specific issues with the measure they'd like addressed.

So, to your point, with the algorithms that currently stand, only a few do end up in this do-not-support for rulemaking. But I think that is something that would sit, maybe, on the table for tomorrow.

CO-CHAIR KAHN: It may not be possible, but I think for our discussion tomorrow, it would be useful if it could be done easily. If it can't be done easily, then don't do it.

But if it can be done easily, if we could get some array of just the number that fell

into each of the categories from the algorithmic 1 2 process, so we can just get a sense for what your process -- now, of course, it's apples and 3 4 oranges between the different measures, but it'll 5 give us, I think, some sense and help that discussion. 6 CO-CHAIR PINCUS: 7 It would also, 8 actually, probably be a good idea if you're doing 9 that, to sort of do it as kind of a matrix. So, have the second column of what 10 11 came out of the workgroups. 12 CO-CHAIR KAHN: Right, right, yes, I'm 13 sorry, I should have said that, too. 14 So, we can try to pull that DR. AMIN: I think one of the things that I would 15 together. 16 just quickly point out is that the inputs, the 17 number of the measures and the process that CMS 18 has used into the algorithm has also changed, 19 meaning the number of measures that are going 20 through the algorithm.

opinion, precision about what CMS is recommending

Meaning there's a lot more, in my

21

or even putting forward to the MAP has sort of 1 2 changed over the last years as well. So, I just want to keep that in mind 3 as we think about and look at the context of how 4 5 the algorithm is working. Yes, I guess I'm just 6 CO-CHAIR KAHN: 7 looking at this slice in time, and the question 8 is what is -- and maybe the algorithm works fine. 9 And I think, actually, comparing it to what came out of the discussion will just tell us 10 something, I'm not sure what, but I think it'll 11 12 give us some information, I think, that we can then delve into when we look at the categories. 13 14 Because how it does it in the algorithm I think tells us a lot about your thought processes. 15 16 Okay, thanks. 17 MS. BUCHANAN: Great. Thank you very 18 much. So, we will now review the voting 19 instructions. 20 MEMBER QASEEM: Can I ask a quick 21 question? Because I might have missed it. 22 conditional support that we do, can you just

remind me what happens? Because I still was just thinking about there were a lot of conditional supports we approved last time around as well, and I'm not sure the Committee ever hears back what happened to what did CMS decide to do? Or even if they agreed or didn't agree? Was there any change?

I mean, I was trying to figure out if there was going to be feedback. But if this is not the right time to talk about it, maybe we can talk about it later.

MS. O'ROURKE: Sure, so I think maybe that's something we could also get some input on the feedback loop for tomorrow.

At the workgroup's Fall web meeting, we did implement a feedback process where CMS brings back information about some of the measures that were under consideration on prior lists, and gives an update about where they are in their development, how MAP's input has been taken, which ones have gone in for endorsement.

It's not comprehensive, given,

essentially, the accumulation of measures since the beginning of MUC makes it a little impossible to go back over every MUC list.

But I think that might be something you could also put on the table for tomorrow, to find out what information would be most valuable to you all but you'd like to see come back, and then how we can collaborate with CMS, what NQF can track internally and bring back to the workgroups.

MEMBER QASEEM: Yes, so what I'm looking for is very simple because I know everyone is busy and no one will have time to go through it.

So, we have let's say 30 measures this time and there's going to be two columns where we say MAP said that this is conditional support only, and whether it was agreed by CMS or not.

Just check yes or check no so we can start getting a feel for percentage of measures we are saying are conditional, and what percent of them are being changed or not being changed by

1 CMS.

So, we can start having that kind of discussion. Because at this point in time at least, I have no feel for what's going on.

We make these recommendations, I do know what happens of course, but it'll be more transparency, bringing transparency to the process, so we can start figuring out that the work we are doing, a lot of us are spending a lot of time.

It's a very important group. But the time and effort we spend, what happens to that time and effort?

MS. O'ROURKE: Okay, so just to make sure I'm understanding what you're saying, going through the rules for the previous year and just tracking which measures ended up being proposed and finalized, and if not supported or did-not-support.

CO-CHAIR PINCUS: Kate, and then Carl, and then Marissa.

MEMBER GOODRICH: Sorry, just to

respond to that, so I agree that's helpful; we 1 2 actually did used to do this in MAP, where the next year, we would do exactly --3 4 CO-CHAIR PINCUS: Yes, but usually 5 it's before the MUC list gets presented. Usually it's in the earlier meeting. 6 I don't remember. 7 MEMBER GOODRICH: 8 You guys would remember better than I. 9 think that's a great idea. We should definitely do that, and then 10 as Erin said, we have committed to the feedback 11 12 loop and to figuring out the best way to do that. 13 So, we've started some of that this 14 year; I want to build on that. So, I think what you're talking about is actually not that hard 15 16 to do and would be helpful. 17 CO-CHAIR PINCUS: Carl and then 18 Marissa? 19 MEMBER SIRIO: Yes, I was going to 20 make a similar point, and I just want to put a 21 little bit sharper of a finality to it, which is

I do think we need to spend some considerable

time tomorrow.

I was reminded this morning that, ultimately, our recommendations are just that, they're recommendations, and CMS can do as it sees fit.

so, it seemed to me that for, actually, all three of the categories under support -- so conditional, refined, and do-not-support -- we not only need some kind of feedback loop, but some way to, in fact, re-look at the measure potentially the following year to have a dialog with respect to a decision that may have been contrary to something we recommended, one, to understand what CMS is doing and why, and two, potentially to persuade you of the fact that there may be an alternative way to look at a measure.

And in addition, it would seem to me, to the point, we spend a lot of time with a process where the ball has moved over time, which is fine, as we evolve this.

But it seems to me that in the process

over the last year or two, we've lost a bit of that feedback that at least gives me comfort that, in fact, the input is adhered to, given the thought and the work that goes into it.

So, I would ask Chip and Harold that tomorrow, when we have the process discussion, that we spend a fair amount of time thinking about how we will revise this process in a way that at least makes some of us feel a little closer to the end result, as opposed to what, really, can be viewed as an intermediate step, which is we make the recommendations and stuff happens, and then it is what it is.

CO-CHAIR PINCUS: Yes, I think that's an excellent idea to get more of the sense of the flow of decisionmaking that occurs.

And I think also, tomorrow we're going to have some discussion also about issues around the removal of measures.

So, Marissa?

MEMBER SCHLAIFER: Not just to create extra work, but probably to create extra work,

rather than just doing a list of the recommendations where we had conditional support and then whether or not they went forward, I think it would be important, just because sometimes the conditions do get met, that what the condition was and whether that got met.

Because otherwise it might not be as meaningful.

MEMBER GIFFORD: Yes, I think it's been an evolution of this process, but I think it's become clear that, you know, as everyone's realizing, none of our recommendations are binding to the Secretary.

But what is binding is the comments we make have to be addressed in rules. And I think we have put way too much emphasis on the vote and not on what our comments are. Because that's where there's binding aspect on the Secretary, and there's also a real tension often here, because CMS is in a position where Congress has mandated they do something, and Congress has also mandated that they get recommendations from us on it, and there's often a conflict there.

And so CMS has to figure out all that, 1 2 and so their comments often might be we accept it but we have to move forward because Congress has 3 4 said we have to do something. And we've seen 5 that a number of times here. And I think there's always been --6 sometimes a misperception that our 7 8 recommendations and vote is somehow binding to 9 CMS, and it's clearly not. And I think we've seen that as 10 11 Congress has been more aggressive with 12 measurement development. 13 So, in tomorrow's discussion, I think 14 it would be helpful to talk about not just the vote or reaching a consensus, but it's more 15 16 almost the recommendations. 17 In many ways, it doesn't even really 18 matter what the vote is, it's more the 19 recommendations. 20 CO-CHAIR PINCUS: That's something 21 that Kate and her colleagues at CMS have really

emphasized all the way through, that a discussion

1	is at least as important, if not more important,
2	than the actual voting.
3	MS. BUCHANAN: Okay, wonderful; thank
4	you. And so we'll now review the voting
5	instructions.
6	So, here we have several key voting
7	principles. So, the MAP has to establish a
8	consensus of greater than 60 percent for a motion
9	to pass.
LO	And importantly, abstentions do not
L1	count into the denominator, but if you are
L <b>2</b>	abstaining from a vote, we ask that you let us
L3	know so that helps us.
L <b>4</b>	CO-CHAIR KAHN: Can I ask a question
L5	here?
L6	MS. BUCHANAN: Yes.
L <b>7</b>	CO-CHAIR KAHN: I can't remember.
L8	This rule, I guess, we approved, but is that a
L9	MAP rule or is that a broader rule in terms of
20	the way the 60 percent is treated?
21	MS. BUCHANAN: So, I think Erin can
22	MS. O'ROURKE: Sure, so when we

brought 60 percent to the Coordinating Committee to approve, it actually ties back across NQF and we use 60 percent in the endorsement work.

And correct me if I'm wrong, I believe that come out of our consensus taskforces that we did on the CDP side, and then we adopted it on the MAP side.

CO-CHAIR KAHN: Now, on the MAP side, do we have the power, considering what you just described, when we consider this tomorrow to say 60 or greater rather than greater than 60? Or is that not even within our power because of the broad rule in NQF?

MS. O'ROURKE: I think we'd certainly be interested in your feedback and input, and then maybe seeing how that would work across NQF.

I think the balance of ensuring that the MAP is comfortable with your process, but also that it's easier understandable to everyone. And it's confusing to have one standard for CDP and another for MAP.

So, I think if we could just get the

feedback on the table, and then I'll iterate with everyone in the fall for what works for next year and how to best operationalize a definition of consensus across NQF.

MEMBER KRAMER: I might be able to offer some context on this since I was part of the NQF insights development process work that was done several years ago, which has evolved to this.

This actually is expressed as a voting rule and it was developed as part of a consensus-development process. And in fact, the 60 percent was not intended to be a voting decision rule either in NQF's decisionmaking process or in MAP's.

What was intended was, in both NQF's decision-making and MAP's decisionmaking, was intended to be a consensus process, which is not based on voting. And the 60 percent was used as an intermediate process.

After some discussion, if there did not appear to be broad agreement or consensus,

there would be a straw vote in which they said,
okay, let's just do a straw vote and see how we
feel.

If it didn't reach a 60 percent

If it didn't reach a 60 percent threshold, something like we need to have more discussion here to try to reach consensus. It was not intended to be the end-point in the decisionmaking process.

Now, I think that's a complicated process, it takes longer, it's not as efficient, and it's not as easy to explain.

But I think we've in some ways devolved to a more simplistic voting rule, which is basically a super-majority vote.

And if that's how it's been applied in a number of NQF Committees and in MAP, I think a lively discussion tomorrow, a worthwhile discussion, would be: How do we actually want to use this going forward?

I, both to my personal preference and
I think principles of how NQF and MAP are
supposed to work, and the historical context of

how this came about, would lead us back to a more robust consensus development process, which might use straw voting along the way, but not dumb it down to a super-majority of 60 percent voting as the way we make all our decisions.

CO-CHAIR KAHN: We'll have this discussion tomorrow. I would say I think people should think about it overnight because on the one hand, I totally understand, I mean I understand now better the history of this, and I understand why this is a consensus group.

On the other hand, the trouble is consensus takes a lot of skill as well as a will, and the trouble is on some of these matters -- and potential for agreement -- and on some of these matters, it doesn't exist or you've got taskforces with a lot of work to do.

And they don't necessarily have the right makeup of leadership, frankly, to figure out how to get everybody on the same wavelength.

So, I think tomorrow we need to discuss from a process standpoint, and maybe we

come up with a hybrid.

Because I think that one of the problems we have is that these processes, even when there's 60 percent, we don't want to come out of it with dissidence rather than consensus.

And I think we have some dissidence from the process we had, and so we've got to figure something out, at least for MAP. But that's for tomorrow.

MS. BUCHANAN: Excellent. So, for right now, we have greater than 60 percent, which is what we'll be implementing today and did want to ask if people are abstaining from votes to please let us know. This will help us calculate the scores.

And a couple of the other principles are that every measure under consideration receives a decision either individually or as part of a consent calendar.

And in their December in-person meetings, the workgroups had to reach a decision on each measure under consideration. There were

no split decisions allowed.

This is a process issue that the CC can discuss tomorrow if it's something they want to explore further for future cycles.

Continuing on, how we will run today is that staff will provide an overview of the process for establishing consensus at the start of each in-person meeting, which is what I'm doing right now.

And then staff from our workgroups, as well as the workgroup co-chairs who provide an overview of their discussions and decisions, and then the MAP CC can begin identifying measures for either further discussion or measures they would like to submit a new motion on.

The discussion guide is available on public.qualityforum.org, and it is organized by setting-specific workgroup.

So, PAC/LTC, clinician, and hospital, and did want to just take a quick pulse check to make sure that everyone had access to the discussion guide, because that will be very

important in guiding our decisions.

And so it's public.qualityforum.org.

And if you have any difficulties, please email

MAPcoordinatingcommittee@qualityforum.org and we
can help you get set up.

And so within the discussion guide, each measure has the measure specifications, a summary of the workgroup decision and deliberations. So, this is why it will be so important as we go through today.

So, the voting procedure. We will now walk through the actual steps of a voting process.

The first is that the workgroup staff and co-chairs will present their recommendations from their December 2017 in-person meetings.

And I apologize, this is a text-heavy slide, but we'll go through the important parts.

So, second, the Coordinating cochairs, Harold and Chip, will ask if there are any members that identified any Measures Under Consideration that they would like to discuss further. And during our pre-work, we requested that members identify measures.

The co-chairs can start with those measures previously identified, and then we can open it up to measures that members would like to pull that were not identified. There is no limit to the number of measures a member can identify.

The co-chairs will ask each member to clarify if they are pulling a measure for further discussion or because they would like to submit a new motion on it. Measures identified for further discussion will focus on clarifying questions.

If during the course of discussion, a Committee member determines that the discussion has shown a need for a new vote, the Committee member can put forth a new motion.

Some reasons why a member may want to put forth a new voting measure is if the member disagrees with the workgroup's recommendation, or if there's new information available that would change the algorithm.

Please note we will only focus on one measure discussion or one motion at a time.

One moment.

And once all the measures the

Committee would like to discuss are removed from

the consent calendar, the co-chair will ask if

there was any objection to accepting the

workgroup recommendation on the MUCs remaining on

the consent calendar.

And so if a measure is not removed from the consent calendar, we will still vote on it in accepting it.

So, now we will talk about the instances in which a Committee member wants to put forth a new measure or a new motion on a measure.

In the instances when a Committee member identified a need for a new motion, the member should describe their perspective on the use of the measure and how it differs from the recommendation from the workgroup.

If a motion is up for conditional

support or a refine-and-resubmit, the Committee member making the motion should clarify and announce the conditions, or suggest refinements.

We have tapped some members to be lead discussants for the different program areas and what we ask is that they can be some of the first reactors.

So, if a Committee member identifies a measure either for further discussion or to put forth a new motion, the people who are identified as lead discussants, we ask them to open up the conversation.

And they should state their own point of view, regardless of whether or not it agrees with the workgroup recommendation.

Following input from the lead discussants, the co-chair will then open it up for discussion among the Committee, and we ask that in the interest of time, people refrain from repeating points that have already been identified.

After the discussion, the Committee

member who made the motion has the option to withdraw. Otherwise, the Committee will vote.

And we want to emphasize again if the motion is for conditional support or refine-and-resubmit, we will ask you to list out what the conditions or the recommendations are. And this will be able to inform the discussion as well as the vote.

So, tallying the votes. So, before we begin voting, which we will be doing later on, my colleague, Yetunde, will do a quick run-through of our clickers, but everyone should have a blue clicker in front of them.

For those members participating on the phone, we ask that you email your vote to mapcoordinatingcommittee@qualityforum.org, and we will actually tally your vote.

If the motion receives greater than 60 percent, the motion will pass. If the motion does not receive greater than 60 percent, the cochairs will resume discussion to develop another motion.

If no motion put forward from the Committee achieves greater than 60 percent, the workgroup's original recommendation will stand. And abstentions are discouraged but will not count to the denominator. So, as we said earlier, please announce if you are abstaining from a vote.

I now just kind of want to take some time to talk about the commenting guidelines.

So, comments from both public periods, public commenting periods, are available in the discussion guide. They are linked with each measure.

There will be an opportunity for the public to comment either in the room or over the phone, prior to the discussion for each setting-specific workgroup. Commenters are asked to make any comments on MUCs or opportunities to improve the current measure set at this time.

We ask that you limit your comments to two minutes, and then we will additional have a global public commenting period at the end of

each day.

Here, you can see our calendar of where we are with our activities, and we are coming towards the end of this MAP cycle.

recommendations will be sent to CMS on February 1st. We will then be releasing a series of reports for each of the setting-specific workgroups. And the clinician one will be released on March 15th; the hospital PAC/LTC programs, that report will be released on February 15th. And then as I said earlier, the actual recommendations will be released on February 1st.

And before we go into questions on this, I did just want to take a quick opportunity to talk about the MAP Rural Health Workgroup feedback on the Measures Under Consideration.

So, as many of you know, the MAP Rural Health Workgroup is a new MAP workgroup that was established this year.

And during their December 2017 web

meeting, the Rural Health Workgroup engaged in a high-level discussion of the MUC list using the perspective of the appropriateness for their patients in their communities.

And there were several observations that they had that we would just like you to keep in mind as we go through the three program areas.

The MAP Rural Health Workgroup members emphasize the importance of considering the low case-volume challenge for rural providers for several measures on the MUC list.

For example, applicability to rural providers may be challenging for the cancer readmission measure.

On the other hand, they noted that the shingles vaccination measure would likely be resistant to the low case-volume challenge.

Workgroup members identified the topic areas of diabetes care, vascular care, opioidrelated care and events, HIV screening, prostate screening, and simple pneumonia hospitalization, as areas of interest and rural relevancy to them.

1	I will now turn it over to I believe
2	Harold to facilitate the discussion? Chip,
3	sorry. Chip?
4	CO-CHAIR KAHN: Any other questions
5	about the voting process?
6	MEMBER HALL: Just a minor
7	clarification, what's our total denominator on
8	this Committee and our quorum?
9	It seems like having a quorum, but I
10	suppose if there were abstentions, we could lose
11	it.
12	CO-CHAIR KAHN: How many people do we
13	have on the phone in total here?
14	MS. O'ROURKE: I'm going to punt that
15	one to Kate and Yetunde to get us our numbers.
16	CO-CHAIR PINCUS: And also, Rich, are
17	you on the phone yet?
18	MS. BUCHANAN: So, we'll do a quick
19	calculation because I know Rick hasn't joined us
20	yet, but we'll get you that number very soon.
21	CO-CHAIR KAHN: I've got to believe
22	we're

1	MS. BUCHANAN: Without Rich, we have
2	32 right now.
3	CO-CHAIR KAHN: We have well over a
4	quorum?
5	MS. BUCHANAN: Yes.
6	CO-CHAIR KAHN: That was the question,
7	whether we had a quorum? Yes, I don't think the
8	quorum's a problem.
9	MEMBER O'BRIEN: I just wanted to
10	clarify, for this year, for pulled measure, the
11	Committee member is making a motion for a
12	different recommendation, so is the vote,
13	essentially, this year going to be a yes or not
14	vote, an up or down vote on the motion?
15	Okay. Thank you.
16	CO-CHAIR KAHN: Wait, and I guess if
17	I understand it, for us to change the
18	recommendation, it's got to be in excess of 60
19	percent.
20	So, it's not just having the vote;
21	it's not just the majority. It has to be, in the
22	sense, the super-majority.

Even though I understand what Bill described as a process, we still have the formal

MEMBER O'BRIEN: Right, I just wanted to clarify that we weren't loading on all the categories of potential recommendations.

CO-CHAIR PINCUS: No, no.

Now, if that motion doesn't pass, then we go back to discussion and see if somebody wants to make a motion for a different category.

CO-CHAIR KAHN: I think, and we spent time talking about the problems that came up in the Hospital Taskforce, and I'm not sure this really matches with coming to a consensus, but I think we have to be linear in terms of the process.

It's sort of Robert's rules that we have motions and let's say somebody puts a motion to remove and it gets 59 percent, whatever the number, it doesn't get to the 60, then the floor is open and after the vote, somebody can bring another motion.

And I think if people stop bringing motions, then it reverts back to -- and we don't get more than 60 -- it reverts back to the recommendation of the task force where they may have been above 60, or to the base consent calendar.

MEMBER O'BRIEN: Right, that sounds like the right approach to me.

CO-CHAIR PINCUS: And of course, all the discussion about all the different motions would then be input to CMS.

CO-CHAIR KAHN: Right, and I think tomorrow we really need to talk about what we want to do, because I think Bill's raised an important point, what we define as a consensus process.

But at the end of the day, I'm concerned because of just lack of time and effort, the best we can probably do with some of these is have the substantive input.

I mean the text of why we were concerned as much as the vote, because it's going

1 to be difficult to get to a vote, frequently, of 2 more than 60. And just to echo what 3 MS. O'ROURKE: 4 Chip was saying, and Harold, and we do capture 5 all of your input for the various reports Kate went through. 6 So, as we're going around, please feel free 7 8 to express any opinions you have about the 9 measure. 10 We don't only pass on the vote to CMS; 11 in the report, we document how the group came to 12 that, what the opinions of people who supported that decision were. 13 14 Also, if there were any minority opinions who disagreed with what MAP put forward, 15 16 we put all of that in the text of report that 17 goes to CMS. 18 So, to Chip's point earlier that they 19 weigh those heavily, so please share your input, and we do include all of it in the report. 20 21 So, while I know there's some concerns 22 about the voting process and if that really lets

	everyone truly reflect their opinion, we do
2	capture all of your discussion, and that goes
3	along with the vote.
4	It's not just a 61 percent of the
5	Coordinating Committee said this. It's a nuanced
6	report of the decision.
7	So, I'll just let you know that that
8	is heard.
9	CO-CHAIR KAHN: Okay, any more
10	questions? We're 15 minutes ahead.
11	MS. O'ROURKE: Should we do the voting
12	test before we start PAC public comment so that
13	we're all ready to go? Did you have a test
14	CO-CHAIR KAHN: So, how does the
15	voting test work? Who's in charge of the voting
16	test?
17	MS. O'ROURKE: Just to make sure that
18	your clickers are working and everyone knows how
19	to use them before we get into
20	CO-CHAIR KAHN: So, everybody should
21	pick it up.
22	MS. O'ROURKE: One second, it ties to

a slide that we need to bring up. 1 2 Apologies, I've put Yetunde on the spot but this, I think, was going to happen after 3 Jean-Luc did some of the presentation on PAC/LTC. 4 5 But to that we respect that we told 6 the public the comment period would start at 10:30 a.m. and we're running a little -- I 7 8 thought we could get this test vote out of the 9 way and make sure everyone understands the clickers and is comfortable. 10 11 Oh, well, then we could go. 12 want to skip the test vote or just do it since I 13 brought it up? Okay, never mind. 14 Okay, so let's transition to the 15 PAC/LTC. CO-CHAIR KAHN: So, now we're going to 16 17 the public comment on PAC/LTC, and I'll ask 18 whether anyone would like to make a public comment on the PAC/LTC programs. 19 And remember the guidelines for public 20 21 comment; they're up on the screen.

comments to just PAC/LTC recommendations, limit

comments to no more than two minutes if possible, 1 2 make any comments on MUCs or opportunities to improve the current PAC/LTC measure set at this 3 4 time. 5 So, why don't I do this? Obviously, I'm hoping no one is going to come at 10:30 a.m. 6 7 that isn't here already, so do we have any 8 comments in the room from anyone? 9 Okay, not seeing any comments in the 10 room, let me go to the phones. And the 11 operator's supposed to say something now. 12 At this time, if you'd like OPERATOR: 13 to make a comment, please press Star then the 14 number 1. Okay, any comments? 15 CO-CHAIR KAHN: 16 OPERATOR: There are no public 17 comments at this time. 18 CO-CHAIR KAHN: Okay, well, we are 19 ahead of schedule but I think we should go on. Ι 20 think we gave people ample opportunity. 21 And so I guess now we're going to prerulemaking recommendations on the PAC/LTC, and 22

Leah and Gail Hunt are the primary lead 1 2 discussants. We're going to discuss the key themes 3 from the workgroup meeting, review and finalize 4 the broader guidance, and review and finalize 5 workgroup measure recommendations. 6 So, I think I'm turning it over to 7 Jean-Luc to coordinate this. 8 9 MR. TILLY: So, good morning, everyone, I'm Jean-Luc Tilly. I'm joined today 10 on the phone by both my co-chairs for the PAC/LTC 11 12 Committee, Gerri Lamb and Paul Mulhausen. 13 So, we actually just had one measure 14 under consideration this past cycle for the SNF QRP, the CoreQ short-stay discharge, which we'll 15 16 get into a little bit later. This slide kind of summarizes the 17 18 overall themes, and we'll dive into each of these 19 in turn. 20 But broadly speaking, we're talking 21 most about making a move to high-value measures 22 and trying to give some guidance to CMS on future measure development for these programs.

And it just says record the short-stay discharge, so let me just move one more.

All right, so of course, most PAC programs are affected by the IMPACT Act, which requires PAC providers to provide Center S patient assessment data, and guides the adaption of new measures in those programs.

CO-CHAIR PINCUS: Talk a little bit more into the mic.

MR. TILLY: So, the workgroup encouraged continued alignment of those measures and recognized that the measure under consideration in the CoreQ was consistent with those principles.

The MAP Workgroup did note several opportunities to continue to address quality in IMPACT settings.

So, here, we're thinking primarily of patient-reported outcome measures that were emphasized, as well as measures of transfer of information of care preferences, and finally,

medication management.

Now, see, I have care preferences beyond the end of life, but presumably someone else is in charge of that. We meant more care preferences beyond just end-of-life care. So, shared accountability across the care continuum is also a big theme.

So, of course, we mean partnerships between providers, between hospitals and PAC/LTC providers, but then also the inter-operability of health IT technology.

So, now looking at considerations for specific programs in turn, first, for SNF, a number of opportunities for new measurement were advanced.

So, that includes measures of shared accountability between stints in hospitals, measures around transfer, measures around advanced directives.

And of course, one measure was submitted for this program.

So, turning now to the IRF QRP, MAP

had some similar notes about the transfer of information, and then a setting-specific consideration around opioid use in IRF facilities. There's a special consideration there.

And then the workgroup also thought about some of the infection measures that are used in IRF settings have a relatively low incidence rate relative to other settings, and thought those measures could be re-examined.

And finally, for the LTEC QRP, the workgroup cited measures addressing mental and behavioral health and particularly for depression, which is maybe a bigger problem around setting than in others.

For the Home Health QRP, the workgroup highlighted the importance of social determinants of health, which can have a much greater impact in that setting.

The workgroup noted that measures around the maintenance of ADL capability could have particular value in this setting.

So, not just improvement but also maintaining a relatively good performance.

And then, finally, for the hospice program, the last program, a program not covered by the IMPACT Act, the workgroup cited several -- including medication management, caregiver support, bereavement, and then finally, safety and functional status measures.

So, turning now to the CoreQ, our measure under consideration, the workgroup supported the measure.

The measure addresses an important

CAHP; it's a patient-reported outcome but it was,

actually recently in 2017, endorsed by the

Person-and-Family-Centered Care Standing

Committee.

The workgroup did note that the measure introduces the possibility of some burden for SNF, and actually, the one public comment we received very closely mirrored the recommendation.

It supported the measure but they

1 noted that what seemed like the timing of 2 implementation in 2019 seemed maybe too soon for that setting. 3 So, I think at this point, I'll turn 4 5 it over to my co-chairs, Gerri and Paul, who are both on the line. 6 7 I think they might have a few things 8 they'd like to add. 9 MS. LAMB: Good morning, this is Gerri Lamb. Thanks, Jean-Luc, and I just would like to 10 11 acknowledge the NQF team. 12 We had a very, I think, productive, robust discussion in December. 13 And Erin and Jean-Luc and Miranda 14 15 really put a lot of time and effort into setting 16 context and helping us really look at the gaps in 17 how we could move forward on some of these gap 18 areas. 19 So, we also had discussions about how 20 the measures, the post-acute measures, fit into 21 IMPACT, MACRA, had a great discussion about

22

attribution issues.

So, as you all know, we had one MUC measure so we spent a lot of time looking at how to move this field forward.

And as Jean-Luc gave you the overview, some really key areas, key discussion points, were related to transition and information exchange.

And with this one MUC measure, that came up as well in terms of the feasibility of EHRs to accommodate this.

And I think people felt that this whole field was moving ahead, that patient experience was critically important, and this was sufficient time for the EHRs to catch up.

We heard a lot about beginning to bring in functional performance, and also some of the national issues like opioid use.

So, really, I think great meeting, lots of discussion concerned about the gap areas and being able to get measures that will move the field forward.

Paul, comments?

DR. MULHAUSEN: Yes, thanks Gerri. 1 2 This is Paul Mulhausen, and I'm the sort of rookie co-chair with Gerri. It's a pleasure to 3 4 be here. I don't know that I have a lot to add 5 to context; that's already been shared with you 6 7 by Jean-Luc and Gerri. 8 I do want to reflect a little bit 9 about the deliberations around the one measure that we had on our MUC list, which is the survey, 10 the patient experience of care survey or the 11 12 survey that we had talked about. And I think the deliberations were 13 14 heavily influenced by the observation that this was an NOF-endorsed measure. 15 16 It was a measure that was put forth by 17 the industry, it was put forward by the American 18 Healthcare Association, and most of the testing 19 and development was done by them. And an occasional member of our 20 Committee raised concerns about that. 21

People already perform fairly well

with this measure. It has an 80 percent satisfaction, good score.

And although that was framed as there's lots of room for improvement, the deliberations included some observations that people already perform fairly well in this measure.

Gerri and Jean-Luc have already highlighted that we have a deficit of patient-reported, or resident-reported, or family-reported, measures and this was an important gap.

And you may wonder at this point, whatever happened to the nursing home CAHPS? Which was my question.

So, although we had nursing home CAHPS for the patient experience of care, that is no longer endorsed by the NQF.

So, having this CoreQ measure as a reflection of the patient's or resident's experience of care ends up being very important at this particular moment.

And then the only other sort of issue 1 2 around the measure in our deliberations was actually how to most effectively implement it. 3 You've heard Jean-Luc and Gerri 4 5 discuss sort of the burden issue of having to send out survey mailings to get the results. 6 7 And then there's also the actual 8 testing and validation has all been done around 9 time-limited survey processes. And I think CMS is going to have to 10 11

And I think CMS is going to have to think very hard about how to actually implement this measure as part of a rolling quality-reporting system.

So, in balance, everybody's excited about the measure because it does fill that gap of patient experience, but there were some reservations that were primarily swayed by the observation that it was a very well-vetted NQF endorsed measure.

So, anyway, thanks for having me on the phone today, appreciate the opportunity to share what I think happened during our meeting.

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And I, like Gerri, have to thank 1 2 everybody at NQF who helped us coordinate and support a very robust and engaging conversation 3 4 at our face-to- face meeting. 5 So, thank you. Great, thanks, Paul. 6 CO-CHAIR KAHN: Okay, are there other comments from -- I guess 7 8 we have discussed this but are there any other 9 comments from --10 MS. O'ROURKE: So, no one's pulled it but if we still want to ask our lead discussants, 11 12 since we do only have one measure, if they have 13 any comments? 14 MEMBER HUNT: It's Gail Hunt. I would just echo what's been said. 15 16 I think it's critically important to 17 get the patient-reported viewpoint of this, and 18 this is, obviously, a short-stay discharge 19 measure and not for the longer-term people who 20 are in the hospital. 21 But I just wanted to say that I think it's really important and we should be supportive 22

of that because it does represent the patient-1 2 reported outcome that we don't often hear from. CO-CHAIR KAHN: Just to ask a 3 4 clarification, so this is, I guess, beside the 5 testing and validation, this particular instrument is done by mail you said? 6 7 MS. O'ROURKE: I believe, correct me 8 if I'm wrong, there's several modes of 9 administration. Mail is also in the EHR, am I 10 incorrect? 11 MEMBER GIFFORD: No, it's currently 12 mail or telephone. 13 CO-CHAIR KAHN: You know, I'm going to 14 just make a comment. And I assume it's mail or 15 telephone because of the rules of CMS, or the way 16 you did it? 17 MEMBER GIFFORD: Kate may want to 18 comment on the issue about rulemaking, what to 19 do. 20 I mean this is actually a suite of 21 five measures we have for long-stay and AL, but 22 traditionally, our long-stay measures are not put

in rule.

So, all of our long-stay measures, none of them are ever put in rule, a five-star is not in rule.

Only the measures related to shortstay are put in the rule, and so I believe that's why this is coming forward here and why you're not seeing some of the longer issues.

As far as how they propose to acquire the administration of this, we do not know because they have not yet, that we're aware, put it in any rulemaking.

It would be within the CoreQ program and, of course, there's a lot of details and I think we would support as well a lot of details to work out how this would roll out.

But we put it forth in the call.

Understanding the MUC list, it's not just

measures that CMS puts forth.

They put a call out to the general public right about now, right? And anyone can put measures in.

1 They then review whether they are 2 worthy of putting into future rulemaking. CO-CHAIR KAHN: I just meant in terms 3 4 of the response. 5 Because in terms of the feedback, the vehicles for feedback from the patient are 6 7 limited, and I guess I'm sort of reacting here to 8 HCAHPS more than this. 9 MEMBER GIFFORD: Yes, it's either telephone or mail. 10 11 Yes, which I just CO-CHAIR KAHN: 12 think is -- my mother's 91 and my father's 92, 13 and they're very capable people and they're in 14 and out of institutions and they spend more time on the computer than they do on the phone. 15 16 And I just don't understand it. Yes, so we don't 17 MEMBER GOODRICH: 18 have any kind of restrictions, or philosophical 19 restrictions either, around the mode for these 20 surveys. 21 As long as they have been tested and 22 found to be validated, you have to go through

that rigorous testing process.

I will just say for everybody, we are conducting testing now for several of our CAHPS family of surveys, web-based administration as well, but that does have to go through the rigorous testing process.

But we know folks are eager to move in that direction.

CO-CHAIR KAHN: I'm with you, but it really needs to be done because I'm sure it can be validated and the response rates are not sufficient, and that will really help the response rate.

MEMBER KAHN: This is Maureen. I echo a lot of the comments made and I think it's important, as someone who also has an again parent, and I lived through this survey process with the parent, they are being overwhelmed with surveys from physician offices, hospitals, and now another center.

And I think we need to make sure as we go through the testing that it's clear where it's

coming from and what they're being asked 1 2 questions on. Because I know it takes a lot of time; 3 4 these are not quick surveys. 5 CO-CHAIR KAHN: That's a good point. But we're sort of in a dilemma because if you're 6 7 going to be patient-centered, you're going to be 8 patient-centered. 9 MEMBER KAHN: I don't disagree. 10 CO-CHAIR KAHN: And everybody's 11 getting the same message, too. MEMBER KAHN: Yes, I just think there 12 13 has to be clarity on what they're being asked to 14 comment on. 15 CO-CHAIR KAHN: I understand. 16 MEMBER KAHN: Because I just went 17 through an experience with a rehab hospital, five 18 days in a rehab hospital, and so she has now 19 received I think four different surveys that I'm 20 now working with her to help her because of her 21 vision and the complexity. 22 What are you evaluating?

1	DR. MULHAUSEN: This is Paul
2	Mulhausen. May I make an observation?
3	CO-CHAIR KAHN: Yes, and then after
4	that
5	DR. MULHAUSEN: All right, so this
6	particular CoreQ survey is very simple, certainly
7	relative to the CAHPS surveys.
8	It's four questions; it's basically a
9	satisfaction survey ranging from very unsatisfied
10	to very satisfied.
11	And so I think in terms of something
12	that's quite simple to complete, you would find
13	this meets the bill for that.
14	MEMBER KAHN: Fantastic.
15	CO-CHAIR KAHN: Okay, Leah?
16	MEMBER BINDER: I just wanted to make
17	one comment about the comment, the public
18	comment, that was received on this measure and
19	about the potential burden of this measure.
20	I'm not familiar with short-stay, I'm
21	not familiar with this particular area of the
22	healthcare system so I don't pretend to know the

level of burden. 1 2 But I would say that it's my recommendation to CMS not to slow down the 3 4 implementation of this measure in any way. It's 5 a really urgent thing. I think it's very important to start 6 7 to get more patient feedback and have that inform 8 the ongoing operations and quality that's 9 delivered at these organizations. 10 So, I do think that it's a very 11 important measure. I congratulate everyone for 12 putting this forward and I do strongly support 13 it. 14 CO-CHAIR KAHN: Okay, so Harold has a 15 comment, and then we don't really have any 16 actions, it's just the discussion on the measure? 17 Okay, so Harold will comment, and does 18 anybody else have -- do we have any comments on 19 this measure? 20 Okay, then, Harold, why don't you 21 comment and then we'll go --22 CO-CHAIR PINCUS: So, I had two

questions, one for CMS and one sort of more generally to the workgroup.

The one for the workgroup is I noticed that for the hospital long-term care program, a recommended priority was behavioral health measures to be developed.

But I didn't see that as a recommendation for the Home Health or the hospice one, and I'm curious, was there some distinction there, and concern?

Because I think that, in some ways, those may be as important or more important in terms of behavioral-health criteria.

And then the second question I had,

Kate, to you was the comment about sort of the

no-longer endorsement or continuation of the

CAHPS long-term care measure, just to get some

sense of where is CMS in terms of the

continuation of CAHPS and other forms of patientreported outcomes?

MEMBER GOODRICH: So, Pierre maybe needs to help me here, I'm not sure. The

nursing-home CAHPS, I'm not sure we've ever even 1 2 used that. As long as I've been at CMS, we 3 haven't used it, and I don't know if that's 4 5 because it never got NQF endorsement, or lost NQF endorsement, I'd have to go back. 6 7 I know we've got Alan Levitt on the 8 phone and he may be able to answer that question. 9 But that --But not specifically. 10 MR. LEVITT: 11 MEMBER GOODRICH: Right. 12 CO-CHAIR PINCUS: But just the broader 13 program? 14 MEMBER GOODRICH: Right, no, CAHPS and patient experience are as important to us as they 15 16 have ever been, so the fact that we don't have one in nursing homes, and I know we've been in a 17 18 lot of communication with Giff. 19 And the AHCA team on this measure have 20 been very supportive of this measure, to have 21 something out there that can be used in this 22 setting.

1 But certainly, as it goes with CAHPS, 2 still a very high priority for us, that has not changed. 3 CO-CHAIR PINCUS: And is there 4 5 anything going on with regards to behavioralhealth-specific CAHPS? 6 Do you know the 7 MEMBER GOODRICH: 8 answer to that? Not that I'm aware of, but we should talk about that. 9 I mean in my role as 10 CO-CHAIR PINCUS: 11 being the person for behavioral health. 12 But also, the other issue had to do 13 with, specifically for the health home and the 14 hospice, there's not a specific recommendation 15 around development of behavioral and health-16 related measures. 17 MEMBER GOODRICH: Recommendation by 18 the MAP? 19 CO-CHAIR PINCUS: In the priority. 20 Yes, by the MAP, and actually, it 21 wasn't so much directed at you but it was directed towards the co-chairs of the workgroup 22

1	in terms of did that come up?
2	MS. LAMB: This is Gerri. I think we
3	were dealing with recommendations at two levels.
4	One went across all of the post-acute
5	services, and some were specific.
6	The behavioral health, mental health,
7	went across all of them, and then when we did the
8	drill-down, I think IRF and the long-term care
9	hospital went into it in more detail.
LO	Home Health and also the IRF were also
L1	looking at substance abuse as a subset of
L2	behavioral health, but I don't think that it
L3	wasn't a priority.
L <b>4</b>	It went across, but there were
L5	different drill-downs.
L6	CO-CHAIR PINCUS: Thank you.
L7	CO-CHAIR KAHN: Okay, so I think we've
L8	completed our discussion of this particular
L9	measure but I'm sorry, David?
20	MEMBER INTROCASO: Thank you. I'll
21	pick up on the spirt of the discussion as much as
22	the voting on this.

So, my first comment on this measure, 1 2 when you see these measures, I always find myself asking, it took this long to get to this measure? 3 So, that's to say we're strongly 4 5 supportive of this measure, but I would like for context and to make note of the importance of 6 7 this measure as it relates to spending variation. We know it's mostly in post-acute and 8 9 we know as it comes to P-for-P, I don't think we have P-for-V, but as it relates to pay-for-10 performance programs, the bundles and ACOs, we 11 12 know the savings is on the post-acute side. 13 So, I would encourage, since Kate's 14 well aware I'm a big person on measuring for value, I would say, beyond just collecting this 15 16 measure and data about it, is to correlate how 17 are beneficiaries in a P-for-P model scoring on 18 this measure? Because, obviously, P-for-P providers are trying to save on spending to win 19 on their financial benchmarks. 20 21 So, if they're having particularly

these patients, who would be more likely short-

1 stay patients in an ACO possibly, are they more 2 dissatisfied than just SNF patients in fee-forservice? 3 4 Thank you. CO-CHAIR KAHN: Any other comments? 5 I guess no one wants to pull anything else, so 6 7 any just general comments like David's about this 8 area? 9 Okay, we do need to have a vote on the 10 consent calendar, so what I'm going to ask is 11 that we get an explanation of the voting now in a 12 moment, and that we then have our vote. 13 I, then, am going to suggest, because 14 we're doing well on our time, that we have a ten-15 minute break post the voting and that we'll come 16 back right before 11:00 a.m. 17 And we'll take the attribution 18 discussion from tomorrow and staff will present 19 the terms of that discussion. 20 MEMBER GIFFORD: Sorry, I'm 21 misunderstanding. 22 So, I was staying silent because I

have to abstain from the voting on this so I want 1 2 to make sure we drop the denominator. CO-CHAIR KAHN: We were about to. 3 MEMBER GIFFORD: I had more general 4 comments about post-acute measures and I didn't 5 know if we were going to take that up after this 6 7 vote. That's why I've been staying silent 8 9 through that discussion. Why don't you make 10 CO-CHAIR KAHN: those comments now before we vote, and then we'll 11 12 -- so, that may move our timing a little bit. 13 MEMBER GIFFORD: Sorry, they're not 14 going to be that long. I do think that the one feedback I 15 16 think would be worth giving to CMS on these post-17 acute settings and these measures, as pointed 18 out, post-acute setting is where a lot of really 19 sick, high-cost and high-acuity -- and people are 20 going through that; 40 percent of Medicare 21 beneficiaries use SNF or Home Healthcare, and

it's the big driver of costs.

Most of the measures still seem to be very provider-centric and with new payment models and everything else, I would encourage you think about how these measures are developed across.

And I know the workgroup mentioned this, but they sort of just talked about cross-cutting themes of transitions, but really -- and I think IMPACT Act was beginning to get at that but I don't see -- there's still a very provider-centric sort of siloed nature of all the measures and the data collection.

And I think as we evolve in the healthcare system, it would be very helpful, and CMS could help drive that thought process about how measures are really at the patient and episodic-care measures, rather than triggered at the provider level.

CO-CHAIR KAHN: Other comments about that? That could probably be true for other areas, I mean not just that setting, too. We'll have a discussion of the consent calendar.

MS. OGUNGBEMI: So, when the Committee

has concluded their discussion and the motion is 1 2 set, I will announce that voting has opened. I will also announce the motion so we 3 4 are clear what we are voting on. Once voting is open, I ask that you 5 point your blue remote towards me because I have 6 the device that captures the votes here in the 7 8 corner. 9 And you make your choice by pressing 10 either one or two. This is a binary vote, so we're either 11 12 voting on yes, we support the motion that's being 13 set, or, no, we disagree with the motion that's 14 being set. Our N right now is 28, but if people 15 16 are abstaining from voting, I ask that you make 17 that known once I say voting is open so I can re-18 calculate the N. 19 When voting is closed, I will announce 20 the result. 21 Additionally, Federal Liaisons 22 not have remotes because they are non-voting

1	participants. So, if you do have a remote and
2	you're a Federal Liaison, please, you can give
3	that back to me, thank you.
4	And also, we will be proxy-voting for
5	our members on the phone.
6	So, I think what we've discussed is
7	that people will email their votes in, and I
8	think it might be easier to submit them through
9	the chat on the web platform.
10	So, email them into the Coordinating
11	Committee inbox, and that's
12	MAPcoordinatingcommittee@qualityforum.org and we
13	will do our best to capture your votes that way.
14	CO-CHAIR KAHN: We have 33 members of
15	the coordinating Committee, so we're well within
16	
17	MS. OGUNGBEMI: Right, and four of
18	them are non-voting participants.
19	CO-CHAIR KAHN: Okay, so in terms of
20	the quorum question, we have a quorum.
21	MS. OGUNGBEMI: Yes.
22	CO-CHAIR KAHN: I wanted to make sure.

1 So, our test vote right now is a 2 recommendation for the SNF QRP, and again, this is just a test because -- yes, well, this is just 3 4 a test, right? We're just testing? 5 Okay, just a test for now, and then we will vote really on the next slide. 6 So, we are 7 voting on the recommendation for the SNF QRP. 8 Let's say that the motion is to 9 support, and this is MUC 17 258, CoreQ Short-Stay 10 Discharge Measure; yes, support, no, do not 11 support the motion. 12 Voting is open. Yes, it's right there 13 on the screen. 14 CO-CHAIR KAHN: How long do we leave it open? 15 MS. OGUNGBEMI: Until we get our N. 16 17 And just give us one second, thank you. We can 18 take a break and come back? 19 CO-CHAIR KAHN: So, the voting 20 software is not working. We have 39 votes so 21 there's some gremlins some place or somebody pressed their button too many times. 22

So, in order to just keep things 1 2 moving while they're fixing the software, let's vote the old-fashioned way, and let's see a show 3 of hands for all those -- I'm sorry? 4 5 Yes, this counts, and Cliff is going to recuse himself. Cliff will completely recuse 6 7 himself from the room, so he can't even see how 8 people voted. 9 So, anyway, all those in favor of -yes, that was the test. We're now going to vote 10 on the consent calendar. 11 12 For those who are in favor of the 13 consent calendar going forward as the 14 recommendations of the Coordinating Committee, as well as, obviously, the workgroup, all those in 15 16 favor, raise your hands. 17 And I guess we need to get a count, so 18 raise your hands high and leave them up until 19 staff tells us that the time for voting has 20 closed. 21 Okay, and then on the phone, anyone 22 should send in their thing, and then are there

1	any noes from those who are eligible to vote?
2	Okay, so we've reached a consensus.
3	I'm sorry, question?
4	MEMBER HALL: I'm sorry, you were
5	saying there were 22 votes? We had just been
6	told a different number.
7	MS. BUCHANAN: Yes, so I want to
8	provide some clarification. So, we have 26
9	people who can vote today, 4 are on the phone, 22
10	are in the room.
11	Three of the people on the phone have
12	stepped away. One person in the room has
13	abstained so now we have an N of 22, so that's
14	the change.
15	So, we can update this throughout
16	because it will be a little challenging as
17	people, especially participating remotely, step
18	out during various times.
19	So, we have 22 voting people, and we
20	have 22 yes, zero no, and 1 abstention in support
21	of the consent calendar for the PAC/LTC program.
22	CO-CHAIR KAHN: A question.

1	DR. AMIN: The threshold will continue
2	to change as the denominator changes. So, Kate
3	will monitor that as we go through the day.
4	MS. MUNTHALI: So, it's 75 percent.
5	So, what we'll do for every vote is
6	announce the quorum and announce the end so you
7	aware of and I think that will help to clarify
8	everything.
9	CO-CHAIR KAHN: So, we had a quorum in
10	the case of the last vote, right?
11	MEMBER HALL: So, to be clear, 22
12	votes is 75 percent of the Committee? What's the
13	actual denominator of the full Committee?
14	CO-CHAIR KAHN: Well, that was the
15	issue, was that was the number that I'd asked
16	for. The problem was we have nine voting members
17	on the Committee as well as voting members.
18	So, in terms of the voting members on
19	the Committee, what's the threshold?
20	MS. BUCHANAN: So, we have, of voting
21	members on the Committee, 29.
22	Because we have four Federal Liaisons

1	and so 22 divided by so, it gives us just over
2	75 percent so we just have quorum.
3	CO-CHAIR KAHN: Okay, so I'm a liar
4	then. So, that 29 doesn't change from vote to
5	vote because that's the N.
6	MEMBER HALL: 22 is our quorum, that
7	won't change.
8	CO-CHAIR KAHN: Yes, that won't
9	change.
10	MEMBER HALL: We have to have 60
11	percent of every vote.
12	CO-CHAIR KAHN: Yes.
13	MS. O'ROURKE: Yes, we need 22 people
14	to participate.
15	(Laughter.)
16	CO-CHAIR KAHN: That is a technical
17	question. A recusal the person is present at
18	the meeting but he or she doesn't count towards
19	the quorum if they have to recuse.
20	MS. O'ROURKE: Yes, okay. And we will
21	monitor these numbers and announce them so
22	everyone is clear on what we need.

1	CO-CHAIR KAHN: I know, my wife keeps
2	calling me, and it actually was my wife.
3	Okay, so now we've moved a little in
4	time, so why don't we say we will reconvene at
5	ten after the hour?
6	And we really need you to be back here
7	at ten after to leave enough time for the
8	attribution discussion.
9	MS. O'ROURKE: And just to clarify,
10	we're going to move up the attribution
11	conversation from tomorrow morning, and do this
12	now so that we can maintain the time for public
13	comment for MIPs and the shared-savings programs.
14	So, we did let the public know that
15	that would be happening this afternoon.
16	CO-CHAIR KAHN: Yes, I think he said
17	that. You know, I think that was it.
18	(Whereupon, the above-entitled matter
19	went off the record at 10:55 a.m. and resumed at
20	11:12 a.m.)
21	MS. O'ROURKE: So, if you could go one
22	more slide?

1	CO-CHAIR KAHN: Okay, if we could get
2	everybody back to their seats?
3	CO-CHAIR PINCUS: So, because there
4	was relatively limited discussion on the PAC/LTC
5	workgroup, we're moving tomorrow's discussion of
6	attribution, some of the general concepts, to
7	right now.
8	Attribution has always been the issue
9	that hovers over sort of everything we do. Yes,
LO	what?
L1	DR. AMIN: I'm just trying to figure
L <b>2</b>	out who's managing the slides. So, can I go?
L3	CO-CHAIR PINCUS: Yes.
L <b>4</b>	DR. AMIN: Okay, great, thank you very
L5	much, everyone. For those that are following
L6	along on the slides, we're on Slide 83.
L7	So, we're going to be moving around a
L8	little bit; this was a discussion that was slated
L9	for tomorrow.
20	So, just to give you guys an
21	orientation about this discussion, we've been
22	working on the concept of attribution as a

measurement science project over the last about year and a half, in two phases of work.

The first phase of work, we completed and we're sort of starting a second phase of work.

The discussion around attribution has come up several times, both in our CDP consensus-development process, but also our work in the MAP.

And so if we move to the next slide, the various components of the legislation are related to the IMPACT Act and MACRA, and demonstrated a continued focus on value-based purchasing to drive improvements in quality and cost by realigning incentives.

And as we start to implement these various pay-for-performance programs, there needs to be some discussion around knowing who can be held responsible for the results of the quality and efficiency measures used to judge performance.

And we noticed through the endorsement

process, and again, through the MAP process, that as we increasingly assess quality using outcome measures rather than process and structural measures, the ability to actually judge who is reasonably responsible for those outcomes has posed some challenges.

So, the concept of attribution is really defined as the methodology used to assign patients and their quality outcomes to providers or clinicians, and to help identify the patient relationship that can be used to establish this accountability.

As we move to a system that's away from fee-for-service to alternative payment models, there needs to be more of -- this issue around patient outcomes and who is ultimately responsible becomes even more important, especially as we move to a health system that's built on shared accountability.

So, we moved ahead with an initial project, and if you could move to the next slide, which really looked to basically answer several

questions, which is as we think about the concept of attribution, what is the definition of attribution?

What are the elements of attribution, and can we basically define some best practices about how one should consider the development of attribution models?

We engaged in an environmental scan with our colleagues at the University of Michigan, Andy Ryan and his group, to conduct this environmental scan of 163 models that were in use, or proposed in use.

And that sort of gives you the breakdown of how many, where they fall, and really characterized what these attribution models -- and basically characterizes these attribution models using program characteristics and also measure characteristics.

If you could move to the next slide?

Thank you. So, the Commission paper had some

very interesting findings that are relevant to

the MAP, which is why we're bringing this

1	discussion
2	CO-CHAIR PINCUS: And do we have
3	access to the Commission paper?
4	DR. AMIN: Absolutely, we can send it
5	along to the Panel if you'd like to take a look
6	at it as we're talking.
7	CO-CHAIR PINCUS: Yes, that would be
8	great.
9	DR. AMIN: Kate, maybe I could ask you
10	to pull that up? Thank you. So, the Commission
11	paper had several important findings.
12	First, the best practices for
13	developing attribution models have not yet been
14	determined, and existing models are largely built
15	off of previously-used approaches.
16	And the tradeoffs in the development
17	of attribution models should be explored, and
18	maybe, potentially more importantly, be
19	transparent.
20	In the attribution models that are out
21	being used or that have been proposed, there's

really not a standard definition for the

attribution models or the elements of an 1 2 attribution model. One might think about this as sort of 3 attribution model specifications. 4 And there really is a lack of 5 standardization across the models, which really 6 limits the ability to evaluate them. 7 CO-CHAIR PINCUS: Can I interrupt you 8 9 for a minute? What do you mean when you say a 10 model? 11 DR. AMIN: And I'll get into that, 12 Harold. I think that was actually one of the 13 challenges we came up with, which was what 14 elements need to be made transparent? Which is, essentially, the question 15 16 you're asking, wasn't really well-defined in what 17 we found. And so I'll get into that a little bit, 18 what we ultimately landed with. 19 Additionally, some of the challenges are that we need greater standardization in order 20 21 to really be able to do comparisons between the models, and really be able to then have a 22

conversation about best practices.

And then, again, the limited consistency across them makes evidence and the ability to build evidence challenging.

And then finally, I think this last point, which has really led to the second phase of our work, is the authors of the Commission paper noted a lack of transparency on how the results are attributed, and no way to appeal the results of the attribution model that, quote, unquote, and I use this very loosely, wrongly assign responsibility.

And so to address these challenges, we convened a multi-stakeholder committee to develop guiding principles, make recommendations, and create the Attribution Model Selection Guide, with the hope that these products would be able to allow for greater standardization, transparency, and multi-stakeholder buy-in, and allow for the evaluation of such attribution models in the future, and lay the groundwork to hopefully develop a more robust evidence base.

And so while we're doing this work, 1 2 and again, Harold, I'll get to your question, there were several sort of guiding principles 3 that emerged as part of this work, which, again 4 5 noting that this was an area of measurement science that was very developmental, the multi-6 stakeholder group really felt strongly that we 7 needed to lay out some guiding principles. 8 9 And maybe, Erin, I can ask you to start walking through this and then I can jump 10 back in as we get to some of the later slides? 11 12 MS. O'ROURKE: Sure, absolutely. 13 So, as part of this preamble, the 14 expert Panel wanted to acknowledge the complex multi-dimensional challenges to implementing 15 16 attribution models, recognizing that the model 17 may change depending on its purpose and what data 18 is available. 19 The Panel used the national quality 20 strategy as their north star.

advancing the NQS, and recognized that

They grounded their principles in

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attribution plays a critical role towards making progress on healthcare goals.

They recognize that attribution refers to both the attribution of patients for accountability purposes, as well as the attribution of the results of a performance measure.

Going back to what you were saying from the Commission paper, they highlighted that right now there's really no gold standard for designing or selecting a model, and that it's crucial to understand the goals of each potential use case.

They highlighted that some key criteria for selecting an attribution model are actionability, accuracy, fairness, and transparency.

So, on this slide, you can see the guiding principles that the Panel came up with: a model should fairly and accurately assign accountability, attribution models are an essential part of measure development,

implementation, and policy and program design. 1 2 The considered choices among available data are fundamental in the design of an 3 attribution model. 4 These models should be regularly 5 reviewed and updated, emphasizing these models 6 should be transparent as well as consistently 7 8 applied. 9 And the model should align with the 10 stated goals and the purpose of the program. Thanks, Erin, I can jump 11 DR. AMIN: 12 back in here. So, the Attribution Model Selection 13 14 Guide, I think one of the things that's important as we began this work was that there was a 15 16 tension between the desire from our stakeholders, 17 generally through the comments that we have had 18 from the MAP process and the CDP process, that 19 attribution models should be fit for purpose and the science related to the attribution. 20 21 And there was really sort of a desire for this, a desire as we set up this group, from 22

stakeholders that we should sort of clarify which attribution models should really be used in a given circumstance.

And really, we landed with the fact that there's really not enough evidence to support the development of such rules at this time.

And so the goal of the attribution model selection guide was really to sort of get back to, basically, Harold's question, which was to define the minimum elements that should be shared with accountable entities, and basically define the minimum specifications of what an attribution model should be, and make those transparent.

And so the elements that we sort of landed on was, and this would basically be the definition of what an attribute model is, and the elements that one should consider when developing an attribution model.

The first is what is the context and goal of the accountability program?

And again, there's certain subelements here that should be considered by those
program-implementers and measure-developers
related to the outcomes and results of the
program, whether the goals of the program are
aspirational or evidence-based, and the
accountability mechanism of the program.

The second is how do the measures relate in the context in which they're being used?

And these really look at some of the measure-specification elements related to inclusion/exclusion criteria, and really, the sample size of whether or not there's enough individuals to draw fair conclusions.

And I know this might generate some conversation around what is a sufficient number of individuals to draw fair conclusions? Which, again, the testing elements here have really triggered a second phase of work.

I'll also draw sort of a connection between this element and many of the public

comments that we received in this Committee's work related to whether or not there's a sufficient sample size at various levels of analysis.

And we hear this a lot, related to the clinician programs around whether or not we can take measures that have been developed for other levels of analysis, and whether they're appropriate to be attributed to individual providers.

Again, I think that's just an example.

There's other care settings that we've seen this

discussion sort of play out through.

Third is who are the entities receiving the attribution? What are the units eligible for attribution? What's the conceptual rationale between the accountable unit?

And whether or not they can meaningfully influence the outcome, again, issues of sample size and whether or not multiple units are actually used to be held for that outcome.

And how is the attribution performed,

the data, how the services are assigned, and the details of the algorithm and methodology.

And so as we look forward to the recommendations of the attribution models, the recommendations are built on the principles and are intended to be applied broadly to developing, selecting, and implementing attribution models in the context of both public and private-sector accountability programs.

I just wanted to sort of reiterate that the environmental stand that was evaluated really very much focused on private-sector programs as well, to inform the work of this group.

And thirdly, recognize the current state of the science and what is able to be achieved now, and the ideal future state of attribution models.

And then also stress the importance of aspirational attribution models in order to drive a greater accountability for our patients as we move the field forward.

And so Erin, maybe I can ask you to 1 2 also just sort of review some of the factors as one selects an attribution model? 3 Sure, so the final MS. O'ROURKE: 4 5 product that the Committee put out in this first phase of work was a series of recommendations. 6 7 The first was to use the attribution 8 model selection guide that Taroon just ran 9 through to evaluate these factors. When making this recommendation, they 10 emphasized right now there is no gold standard. 11 12 Different approaches may be more 13 appropriate depending on the situation and what 14 you're trying to achieve with an attribution 15 model. 16 They emphasize that the model choice 17 should be dictated by the context in which it 18 will be used and supported by evidence. 19 They highlighted that developers and 20 program-implementers need to be transparent about 21 the potential tradeoffs between the 22 accountability mechanism, the room for

improvement, the level of influence that the entity you're holding accountable might have over the outcome you're measuring, as well as the scientific properties of the performance measure you're considering for use.

The second recommendation they made was that models should be tested.

Attribution models offer quality initiative programs to be subject to some degree of testing for things like goodness of fit, scientific rigor, and potential unintended consequences.

They did highlight the degree may vary based on the stakes. Perhaps for public reporting or payment, it's higher than a program that's for internal quality improvement.

Using a mandatory accountability

program, the Panel emphasized that the model

should be subject to testing demonstrates there's

an adequate sample size, that outliers are

appropriately dealt with through either

exclusions or risk adjustment to allow fair

comparisons, and that the data source is sufficiently accurate.

The third recommendation is that attribution models should be subject to a multi-stakeholder review.

The Panel really built on the current lack of evidence, noting that without a gold standard, stakeholder perspectives are key in this area and recommended that a process be developed that allows the stakeholders to weigh in and come to consensus about what might be the best way to attribute performance.

The Panel emphasized that attribution model selection and implementation in both the public and private sectors should use a multi-stakeholder review process to determine the best model for their purposes.

The next recommendation, attribution models should attribute care to entities who can influence care and outcomes.

The Panel recognized that currently, there are models that could unfairly assign

results to an entity -- that was the word they used for providers, just to clarify -- who may have little control or influence over the patient's outcomes, and that for a model to be fair and meaningful, the accountable entity needs to be able to influence the outcomes that they're being held accountable for, either directly or through collaborating with other groups.

And then the Panel highlighted that as we continue to move to team-based care, our facilities become more integrated.

I think, to some of the points that just came up in the PAC discussion as we're moving to an ACO world and trying to really emphasize team-based care, attribution models need to reflect what an accountable entity is able to influence, rather than necessarily directly control.

Then, finally, the Panel recommended that attribution models used in mandatory public reporting or payment programs should meet a set of minimum criteria.

2 transparent, clearly articulated, reproducible methods of attribution to identify an accountable 3 entity that is able to meaningfully influence the 4 5 measured outcomes, that you have an adequate sample size, that outliers are excluded, or you 6 7 used risk adjustment to fairly compare results, 8 that the model has undergone sufficient testing 9 with scientific rigor at the level of 10 accountability being measured, that the model

They highlighted the need for

And the Panel finally recommended there's a need for an adjudication process for attribution models that are open to the public, that allow for a timely and meaningful appeal by the entity being measured when they may be attributed a patient that they've never seen, or they feel they don't actually have the ability to

demonstrates accurate enough data sources to

support our policies, the measure-developer or

the program-implementer demonstrates that their

data is sufficiently accurate to support the

model.

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influence their outcomes.

So, we sort of wanted to bring up the fact that we have a second phase of work, and then the purpose of introducing it back to the MAP Coordinating Committee is the fact that, obviously, these issues related to attribution are very relevant to the work that we do.

So, we're looking for input.

I just wanted to close out by just describing the second phase of work that we're currently in the middle of, which is to expand on the first piece of work that received a lot of interest in terms of public comments and input from you all, and then, obviously, the stakeholders of NQF, as well as our colleagues from CMS.

And so the second phase of work to continue to develop additional guidance in this area of measurement was to really pressure-test the Attribution Model Selection Guide against different use cases, both, again, private-sector and public-sector use cases in which we sort of

looked at some advanced APM models and then also some work by our colleagues at health partners in Minnesota.

So, pressure-test the Attribution

Model Selection Guide and then consider

attribution guidance as it relates to several

important subcategories related to team-based

care, attributing complex patients and special

populations, thinking about data integrity and

data collection, and then other unintended

consequences that should be monitored as we

deploy attribution models.

And so these are the areas that we're continuing to explore in the second phase of work.

And so with that, I think the main question that we've raised to the group is as we continue to developed additional guidance in this area, is there feedback that you all have based on the work that we've done together, that we should consider as we deploy this next phase of work relating to attribution?

So, I'll stop there and welcome 1 2 comments and feedback from the Coordinating Committee. 3 4 CO-CHAIR PINCUS: I just want to kick 5 off the discussion by just raising a couple of issues, then we'll go down one side and up the 6 7 other. 8 So, Taroon and Erin, I have a good idea of what are the issues in attribution but I 9 still don't understand what the unit of analysis 10 is in terms of what is a model? 11 12 And when I think about attribution and 13 how to apply it, I think about it in terms of, 14 number one, what are the changes we want to achieve or what are the goals we want to achieve? 15 16 Number two, who do we expect, who 17 needs to lead and participate in achieving those 18 goals? 19 And number three, what incentives or 20 supports will induce them to engage in that role? 21 Those are the ways I think about it.

And what you've laid out is a lot of

the issues you need to think about in terms of appropriately identifying and motivating groups, entities, to achieve those goals.

But I still don't know what the unit of analysis is.

Like when you said over 160 different
-- are you talking about programs, or are you
talking about actually conceptual frameworks?

DR. AMIN: Yes, so, Erin, feel free to jump in here.

So, when we looked at what was out there, it was really the programs, and what we were trying to extract from those programs is what are the elements that define what those models are?

And so I think some of the questions that you're sort of raising, Harold, and maybe I could just go back a little bit to at least the way that we were thinking about this, it's that the elements that really need to be -- because we didn't see this very clearly laid out for all those 160 programs that were out there.

And I think just the first piece of work was just to take those 160 and say what do you actually need to make transparent as you're deploying any attribution model?

And these are the things that we need to be able to see so that we can actually at least just understand what you're doing, what each of these programs are.

And so I think you've laid out several of these considerations.

I think where the Expert Panel sort of landed as a first piece of work was that given the heterogeneity of what we were seeing out there in terms of the programs, we wanted to at least lay out the elements that need to be very clearly specified and made transparent.

Now, I think the question is then, how do you look at that and make some sort of subjective evaluation of that? And I think we're still not there yet.

This was really just laying out, at least if you sort of think about it as the

specifications of when you talk about an 1 2 attribution model, these are the elements that should be made transparent. 3 4 CO-CHAIR PINCUS: No, I think these 5 are important issues. Two other quick points, then we'll go around. 6 7 One is I think one thing that's not 8 there, what I think needs to be included, is a 9 theory of change. So, what is the conceptual 10 approach in terms of how you go about achieving 11 it? 12 What are the different steps that 13 would have to take place in terms of some causal 14 pathway that would theoretically impact the goal? And the other thing is we've done a 15 16 lot of work around integration of mental health, 17 behavioral health, and general medical care, and 18 of the issues we've talked about is shared 19 accountability or shared responsibility. 20 So, for example, if I'm a 21 psychiatrist who has a patient with schizophrenia

and diabetes, which is not uncommon, I'm

responsible not just for the outcomes for 1 2 schizophrenia, but also for the outcomes for diabetes. 3 4 And my colleague, who's their 5 internist or diabetologist who is treating their 6 diabetes, also has responsibility and 7 accountability for the diabetes outcomes, but 8 also for the schizophrenia outcomes. 9 Which means that we have to talk to 10 each other, and so that's one concept to think 11 about in terms of the way in which accountability 12 is shared. 13 So, why don't I stop there and call on 14 Andy? 15 CO-CHAIR KAHN: I just want to make 16 sure it's clearer. 17 I know you mentioned something 18 concrete. Can you walkthrough one of these concrete programs and then present the problem, 19 before we have the discussion? 20 21 I'm just sort of simple-minded, I'm 22 having trouble visualizing it exactly, so what

would be a good example? Readmissions?

I don't know, just think of an example and then walk us through, just so we all know what the terms are that you're discussing.

MS. O'ROURKE: Sure, and I'll take some liberties.

One example the Committee played with in their meeting was the use of the Medicare-spending-for-beneficiary measure, and the hospital, value-based purchasing program.

so, I think not to necessarily put words in CMS's mouth, but maybe you would say the goal of this program is to improve care in acutecare hospitals, use a cost measure that is the cost for the hospital episode, plus 30 days after to try to get some shared accountability around cost savings, ask the hospital to innovate who they're partnering with, examining what happened to the patient after discharge, might be the type of things we'd ask under this first box.

Thinking about how the measures relate to the context in which they're being used, that

measure is developed for a facility-level analysis in an acute-care hospital.

But when it was being endorsed, some question around this 30-day post-discharge window is that something the hospital can reasonably influence is a fair time period post-discharge to consider costs.

So, I think that's the type of questions we'd ask someone to think about in this second box.

DR. AMIN: Can I just jump in on that last one? So, I think this is actually really helpful.

So, as we look at that measure of the 30-day spending, part of what you would look at there is also how much of the variation in that measure is related to what's going on in the 30 days, versus hospital performance, if you're attributing it to a hospital, in the context of a hospital, value-based purchasing program.

And so that would be another sort of discussion that we would expect to have, as the

measures relate to the program that's being used. 1 2 MS. O'ROURKE: So, then thinking about this third question about accountability 3 4 entities, obviously, the program attributes to the hospital. 5 But the spending within the measure 6 can be driven by post-acute providers and 7 clinicians in the community. 8 9 So, that kind of dissonance for those 10 30 days, who is responsible for the cost versus who is being attributed, and in the program, the 11 12 performance would be held to the hospital. 13 And then, finally, how was the 14 attribution performed? Asking some questions about the data. 15 16 So, in this example, claims data, does 17 everyone have access to it? What are the 18 services, the cost for the inpatient episode plus 19 30 days? 20 Asking some questions about the 21 details of the algorithm, and I think we've already covered a little bit. 22

Okay, thank you. 1 CO-CHAIR KAHN: 2 But I think just in terms of your example, you really have to be careful with 3 proportionality because let's say you have an 4 episode of illness over the 30 days without a 5 readmission, a readmission throws everything off. 6 The hospital's paid a DRG so assuming 7 8 that the DRG is the correct DRG for that ailment, 9 it could end up taking the weight of the proportionality of the total spend. 10 11 But if there's no readmission, then if 12 there's higher than normal spend post, that may 13 or may not be something the hospital could or 14 could not affect. So, it's a very difficult thing to 15 16 assess. I'm sorry, I just wanted to get the 17 example out there. 18 Amy's got some questions? 19 So, I think that we MEMBER MULLINS: 20 really need to -- building on that, are we 21 attributing to measures or are we attributing to

hospitals, or are we attributing to clinicians?

1 Because I think that when I was 2 hearing the explanation, I could kind of hear you saying all of those three different things. 3 4 So, I think we need to be really clear 5 about who or what we are attributing to. need to ask ourselves why we do attribution to 6 begin with? 7 8 We do attribution because physicians 9 need to know who their patients are so they can do quality improvement work. If I don't know who 10 my patients are, I can't improve their quality of 11 12 care. 13 So, if we want to do quality 14 improvement, we have to know who we are taking 15 care of. 16 We know who we're taking care of because we have an attribution model that tells 17 18 us who belongs to us. 19 So, I think, at its root, that is what 20 attribution is for, is to be able to do quality 21 improvement. 22 And if you make it too complicated,

clinicians aren't going to be able to understand it or be able to utilize it, or work within that system, to do the quality improvement work that we're all so desperately trying to do.

So, 130 models, too many, most of them too complicated to understand and too complicated to use, so we need to have something that's simple, something that is the same, something that's harmonized across payers.

How many times do we say this about other things that we're working on? But I think we also need to work in patient choice.

So, the AAFP just submitted an APCAPM proposal to PTAC that was accepted for testing, and we have to come up with an attribution model.

So, the first step in our attribution model was why don't you ask the patient who their physician is and attribute them to that person?

I mean, it seems quite simple when you think about it, but that's probably who the patient's physician is. It's whoever they say their physician is.

So, I think that step doesn't need to 1 2 be missed, is ask the patient who they need to be attributed to. But I think we've got to tease 3 out this work attribution into a measure. 4 5 That's not where attribution belongs; attribution belongs at the clinician level and 6 7 not at the facility level and not at the measure 8 level. 9 CO-CHAIR KAHN: Bruce? MEMBER HALL: Well, I certainly second 10 11 much of what's already been said so I can keep it 12 short. 13 I certainly enjoyed this paper when it 14 was first released, and as a measure-developer formally serving on multiple NQF Panels --15 16 CO-CHAIR PINCUS: A little bit closer 17 to the mic? 18 MEMBER HALL: -- I think it's an 19 important work. There are some gold standards about attribution that I think should be included 20 in the deliberations. 21

So, for instance, sometimes whoever

got paid for the service, there shouldn't be any 1 2 question about attributing to them. But there's another gold standard, 3 which Harold alluded to, and that is based on 4 5 theory of change, who can actually effect a 6 change. And that may be totally different than 7 8 whoever delivered a service or got paid for 9 service, and I think that should be considered as well. 10 11 But I think the one area where the NOF 12 can be more strict is that I think we should 13 stand firmly against use of measures that are not 14 consistent with their development attribution. 15 And I know there's kind of a grey 16 history in the past about stating what the intended use of a measure is. 17 18 But when the development and testing 19 and science behind a measure is based on 20 attribution at a particular level, I don't think 21 anyone should conflict with that.

So, if all the testing shows

reliability at the individual level, that's one thing. That doesn't mean you can go use it for institutional or vice-versa.

And I think the NQF can stand pretty firmly on that, when use of a measure conflicts with the development attribution that was tested.

CO-CHAIR KAHN: Leah?

MEMBER BINDER: I want to suggest an addition to the model, or an enhancement to the model perhaps, that there needs to be monitoring of attribution at later stages once data has actually been collected on a measure.

It is one thing to theoretically surmise who would be responsible for what outcome.

But I think when we're actually looking at the data, often, we recognize, especially when we see variation, that for example, a high performer in a particular measure may have a different way of achieving the outcome, if it's an outcome measure.

And so that can update an attribution

model in a way that will also drive overall 1 2 quality improvement because we begin to identify best practices for achieving a given outcome by 3 looking at the variation in the outcomes. 4 5 And it's kind of like healthcare's a team sport; it's kind of like a baseball team. 6 7 Who is responsible for winning the game? Was it the batter who hit the winning 8 9 run, or was it really the whole team because they had some participation at some point? 10 11 And then once we see a whole bunch of 12 games, we can start to analyze and begin to understand which role had which impact. 13 14 So, I think the attribution model should be an evolving phenomenon and not 15 16 something we see as just a theoretical part of 17 the launch of a measure. 18 CO-CHAIR KAHN: Rob? 19 MEMBER ANTHONY: I just wanted to 20 speak very firmly with my ONC hat in place and ask that as we move into this and we think about 21

it with this project, to think about the role

health IT occupies in this area or, I think more 1 2 importantly, could occupy in this area. I think not only where it can 3 4 facilitate that attribution model currently, but 5 perhaps in the way health IT can be used to innovate in this area and think about how we do 6 7 electronic measurement, ECQMs very much now are 8 modeled on the same way that we do chart 9 abstraction. 10 I do think as you start to look at 11 attribution across multiple systems and multiple 12 settings, you open up some new ways of looking at how that measurement is done. 13 14 I realize that's not necessarily the central driving part of it, but it very well 15 16 could be as we move forward. 17 CO-CHAIR KAHN: Giff? 18 MEMBER GIFFORD: I reiterate Bruce's 19 point that the testing is an important issue. 20 But I'd say one of the things I think 21 is for the Committee to really consider is the 22 mindset we bring, and that you don't

inadvertently, through attribution, reinforce the existing siloed nature of healthcare issues.

The tension clearly is that the

payment models are siloed, and so you could have measurements that are not. But the payment is siloed, so how do you do it?

But I will say the mindset that, generally, most of us bring to the table is very clinically-oriented and we forget about the systems.

And so taking the hospital example, what launched all this was Steve Jenk's article looking at CHF discharges and showing that a large proportion of them had no follow-up care at all in the next 30 days, because it wasn't arranged by the hospitals.

And then as soon as the hospitals were starting to have attribution to rehospitalizations after discharge, they started to change their practice.

We're seeing the same thing in the nursing-home settings and elsewhere.

So, I caution going too far down this 1 2 attribution level to basically have us pull back from this idea of systems and episodes of care, 3 4 and going there. 5 But I understand and appreciate that the payment systems don't really lend itself to 6 7 doing that at the same time. 8 So, I give a lot of credit to a really 9 thoughtful, I think, process going forward. Because I would make sure that that's out there. 10 11 Don't let the dominant view of the 12 current practice reinforce attribution, because I 13 think we will not make progress then. 14 CO-CHAIR PINCUS: Other comments, 15 questions? Oh, Dave, sorry. 16 MEMBER INTROCASO: So, I haven't read 17 this document but I'm extremely interested in doing so, and AMJ would have a great deal of say 18 19 about this. 20 This is an extremely important subject 21 so I applaud your willingness to take it on, very complicated. You'll invariably or inevitably 22

miff off some people or several people. 1 2 In the physician world, this is all about largely P-for-P programs, principally, 3 MSSP. 4 There is a great deal of argument 5 about retrospective versus prospective 6 7 assignment, attribution. There's no consensus relative around 8 9 how you do this. But I will say just a generic point, 10 11 which is I don't see how you get to pay-for-value 12 unless you can intelligently, correctly, 13 appropriately, whatever --- attribution. 14 I can tell you the physicians could go on for hours about their complaints in this 15 16 regard, but this work is sorely needed. 17 I will say I didn't see in these any 18 reference to MIP's or MACRA's patient 19 relationship category. 20 So, maybe as a question, I'm assuming 21 you looked at the Title 1 of MACRA and you talked to CMS about how far down the road they are on 22

that?

DR. AMIN: Yes, so, Erin, correct me if I'm wrong, the conversation did come up in our first phase of work but it was really new in terms of what -- we were sort of interested to see how it would be deployed.

I don't know if there was anything else to say about that.

Yes, I think when the first paper came out, it was something.

They acknowledged the potential source for the future, but I think in this new phase of work, we can explore what's happened since then, and maybe see if Kate and Pierre wouldn't mind connecting us with their staff?

DR. YONG: Sure, so for folks who aren't familiar with these patient relationship codes, we had sought public comment and then developed a new set of codes, which would be included as part of the claims submission process, in which the provider would self-identify their relationship with the patient for

which that claim is being submitted, for that service.

So, it ranges from whether it's an acute or chronic, and then how close, whether they're the referring physician or they're the primary doctor, whether they a short-term relationship or a long-term relationship with that patient.

So, we are starting to collect that data beginning in 2018 on a voluntary basis from clinicians. Ultimately, in the long term, we'd like to see how the data that comes in examine that.

But certainly, there is the potential, as David mentioned, for use of that data once we have a firm sense of the data that's coming in, to potentially include that as part of the attribution methodologies that we use for the variety of potentially quality and cost measures that we have in the program.

CO-CHAIR PINCUS: Amir?

MEMBER QASEEM: I just want to

reiterate two points that Amy and Bruce made. 1 2 One is I think that the attribution models, they need to be easier understood and 3 meaningful to patient's families and clinicians. 4 5 And I think that was one of the guidance principles. It seemed like it 6 7 disappeared in this, and at least in the matrix. 8 I quickly tried to look at this and I didn't see 9 that. And what Bruce brought up, that whole 10 11 issue of these measures are getting the whole 12 reliability, validity is happening at a certain 13 level. 14 But then they're getting applied at a 15 different level, and it continues to be a 16 problem. And I'm not entirely sure what is the 17 reason. 18 So, when you did this environmental 19 scan, I think that would have been interesting to 20 hear about why does that issue continue to 21 happen? 22 And the measures are getting tested at health-plan level, but then clinicians are getting attributed or just being applied over there.

So, I'd like to hear a little bit more and I think that's something that needs to be looked at.

I think you guys have done a very good job when you did the guiding principles. I really like them, but then again, they were not even at 100,000-foot level, they were very, very high up there.

And I think once you start applying them, they started making more and more sense and I like what you had on your Slide 18, in terms of what should happen before attribution models are used in mandatory public reporting or payment programs.

And many do meet these criteria.

I'm not really entirely sure, still, of what happened. Is this something that you have now put out as an official statement and now that's what we're hoping to follow?

What exactly does that slide mean?

Because I think that's the key and I'm really

glad that you're going to continue following.

You have the project where you're going to do this white paper and all that going forward, but if you already have that minimum criteria, and of course we can debate those too, it'll be good to get that going first before we proceed to whatever is the next step of what you're trying to achieve.

Because you already have a good model in place.

And the final point I want to make is
I think since NQF does believe in harmonization
of efforts, there's a lot of good work that's
happening.

Other groups, PCPI comes to my mind;

I'd hate to have NQF come up with something and

PCPI have something -- I think if we can speak

with a single voice together, that would carry

more weight if we go do any of the pairs.

Just more comments and a couple of

questions in there.

CO-CHAIR PINCUS: Just one other point just to add to that, I think in relation to the PCPI and other groups, but there's also, I don't know if it still exists, but the Healthcare Payment, Learning, and Action Network also should be in the loop in this.

Chip, did you have a comment?

CO-CHAIR KAHN: Yes, I thought

something that Cliff said was very profound and important, and it's covered but I don't know, frankly, how to deal with it here.

And part of the dilemma in terms of this aspirational aspect, which I think is what Cliff was talking about, that, on its face, as the system works now, a measure may break the system because the entity, the hospital, has nothing to do with post-acute traditionally.

So, how can you say you're responsible for it?

Well, once you make them responsible for it, then something changes, even though you

don't receive it anymore for -- actually, you

could receive less money than you did before.

And you could then get into the fairness issue.

The trouble is that, part of the

problem is, having been in this position myself,

problem is, having been in this position myself, policy people have notions and aspirations that may not be realistic.

And their attitude is, well, I'm just going to break the current system, screw them.

And it may not actually reflect the reality of what people can do.

On the other hand, then you've got this issue of stretch, which is, well, but if you don't stretch them, they're not going to do it.

And the distance between screw them and stretch is a really difficult value judgment that's really hard to make, and actually, is not applicable in all situations.

Because in some of them, by

definition, it's going to be they're going to get

screwed, and then by definition in others, they

should have been thinking about it and now

	they'll start thinking about it.
2	And actually, it makes a difference to
3	the patient.
4	So, I guess what Cliff brought up
5	really troubles me because I know that the gut
6	reaction of most is if I can't do something about
7	it directly, then don't bother me with that
8	measure, it's not fair.
9	But we've got to figure out how to
10	deal with that.
11	Also, I thought Amy's point was really
12	important, except I think of myself and I still
13	haven't done anything about my blood pressure.
14	And both my cardiologist and my
15	internist both talk to me about it separately,
16	and both are responsible for it and
17	CO-CHAIR PINCUS: Have you talked to
18	your psychiatrist?
19	(Laughter.)
20	CO-CHAIR KAHN: No, I haven't done
21	that yet.
22	And so frankly, yes, they both take

the blood pressure so they pass the process measure, and yes, they talk to me and I said, well, you know, it's because you make me nervous in the room and that's why the blood pressure is what it is.

And I'll go home and do my own blood pressure, and then if I sit down long enough at home, I can get a blood pressure I like until I'm fine. But this is a realistic issue.

If you ask me, so who is responsible for your blood pressure? Well, first, actually it should be me, but I don't know whether it's my cardiologist, and I'm a relatively educated patient.

So, I think that's a very difficult one and then when you get to the seniors who have -- I guess I may be almost a senior, I guess I am a senior.

I don't have a lot of doctors. I'm not sure they could answer that question. If I ask my mother about a couple of her ailments, I'm not sure she would know who to answer.

So, that's a very difficult one too.

Those are just two points.

Let me just add to that. I think in addition to the screw them and stretch, I think there's also a third thing, which is let's make up another entity that's accountable.

And that, in some cases, may be a way to fix the system, but it also comes with it certain startup costs and sort of other kinds of issues that sort of come up down the line.

So, in each of these cases, I don't think there's one great solution and I think what you're developing is a way to evaluate the options in a systematic way.

DR. AMIN: And I would even maybe take a step back to say maybe it's just to make it transparent at least, and then to evaluate, hopefully, at some point.

I know we're out of time. I'm not going to try summarize everything that was said, but I think that there's some key things here that relate to what we were thinking and take it

a step further.

I think the first is this concept of what's the theory of change that we're expecting, and to make that transparent and at least to have that conversation.

I think several times that we've -whether it's through the MAP process or through
the CDP process, it's not always necessarily
clear what we're expecting when we're trying to
do the stretch in the conversations that we've
had.

Even in the initial conversations,

Giff, that you brought up around readmissions, we
went through the conversation around how much can
a hospital realistically be held accountable?

And then moving from one readmission to multiple different settings, the question was what really can each setting do to really handle these readmissions?

And so I think if we maybe started with some more conversation about this and made it more transparent, it might have at least

added, contributed to the dialogue a little bit 1 2 more. I think, Bruce, you sort of reinforced 3 that conversation as well. 4 The discussion around data is one 5 that's been perplexing quite a bit because many 6 7 of these models are really built on retrospective and administrative claims data. 8 9 And so that in itself, going back to the comment that was made around whether it's 10 retrospective or prospective, really, the data 11 12 elements really does make a meaningful difference 13 around what you can actually attribute, and more 14 importantly, offer providers, clinicians, physicians, various different entities. 15 16 If you had the ability to sort of 17 prospectively understand your patient population, 18 your ability to do quality improvement, 19 obviously, would be enhanced. 20 So, many of these things we'll take 21 back and consider and we appreciate the

conversation, obviously, since we deal with this

every time we come into the conversation about 1 2 the various measures as they're used in the 3 programs. 4 We, collectively as a Committee, see 5 these challenges. So, I know this was a very big topic to throw at everyone. 6 7 We would welcome comments once you 8 have an opportunity to review the paper as well 9 via email. Feel free to send those along and 10 we'll be sure to consider those as we think about 11 12 the next phase of work. 13 CO-CHAIR PINCUS: Thank you. Is there 14 one additional point before we go to lunch? 15 MEMBER GIFFORD: I was just going to 16 say you can take immunization measure, it's 17 potential for a population measure down to the 18 individual. 19 Good immunization measures for 20 influenza, good population measure. 21 But until we started holding 22 individuals accountable, a lot of providers said

I don't want to be attributable to it.

But then they design the measures in a way that people were acceptable, then it really helped drive up immunization rates.

And so there is a balance between these two and how to do it, and to Bruce's point, about how you can't just take a population measure and hold an individual position accountable for it without changing the specs.

And so it might be a measure to think about and how do you balance that tension between stretching and screwing somebody?

Chip's measures terminology.

DR. AMIN: This point came up several times. Bruce, you made it and, Amir, you made it as well. The question is what are we doing with this?

It's a 10,000-foot project. The way that we were sort of thinking about this is when we're talking about attribution, let's at least develop the boundaries of what we're talking about, make it transparent.

And now what we're starting to think about is how do we make this actionable, either through the CDP process or the MAP process so that we can actually -- these are minimal thresholds.

Amir, to your point, what do we do with them? Are we going to implement this and require this?

We're not necessarily sure yet, which is part of the reason why we're having this conversation here and in our endorsement processes, to see if this is going to be a future requirement?

Because right now, the endorsement process is sort of use-agnostic.

It asks the question about level of analysis, which is important and what I think the basic elements of what Bruce is making clear, but it doesn't necessarily go as far as what's being asked for here.

And so all of that will be sort of what we're striving to with at least this next

1 phase of work, and then potentially future 2 activities going forward. Erin, anything else to add? 3 4 MS. O'ROURKE: Yes, I think it's 5 probably more a conversation for tomorrow when we're going to ask you to take a look at the 6 7 preliminary analysis algorithm. 8 But we wanted to bring up this topic 9 and at least get you all thinking of are there ways to do that algorithm or other things that 10 11 staff do that we could bring this type of 12 consideration to mind. I think, Amir, to your point of how 13 could we make this actionable for the MAP 14 Committee, is there anything the Coordinating 15 16 Committee would like to add into the algorithm 17 about attribution? 18 Anything that staff should be calling 19 out about some of these issues? 20 So, something to think about overnight 21 as we bring you back tomorrow to take a look at 22 that algorithm, that we do prepare about each

measure is there a role for attribution questions 1 2 there? And what would you like, as the 3 4 Coordinating Committee, the workers to have in 5 front of them when they're making their initial recommendations to you? 6 7 CO-CHAIR KAHN: Okay, that was a great 8 discussion, and I appreciate everybody's input. 9 And it was a good opening for tomorrow, but now let's come back to today. 10 11 And I know everybody's anxious to get 12 lunch, so we'll take between now and 12:30 for 13 lunch. 14 Then at 12:30, I guess we come back, we'll reconvene, I guess we open up the phones 15 16 for public comment on the clinician programs, and 17 then we'll get into a full discussion of the 18 recommendations of the clinician programs. 19 And I have a feeling that we won't get 20 off as easy as we did earlier today from our 21 discussion on the post-acute area.

So, everybody go get lunch and we'll

1 reconvene at 12:30 p.m. 2 (Whereupon, the above-entitled matter went off the record at 12:05 p.m. and resumed at 3 4 12:37 p.m.) 5 CO-CHAIR KAHN: Okay. So we're about enter the clinician review which will begin 6 7 formally with the opportunity for public comment. 8 But before we do that, staff wanted to clarify 9 some of the voting issues, so Erin will -- I guess Erin will take the lead on that and do that 10 11 for a moment and then we'll open the mics. I'll 12 say a few other things and we'll open the mics for public comment. 13 14 MS. O'ROURKE: Apologies. We have a visual aid that we are pulling up momentarily. 15 16 I just wanted to, before we take any 17 move votes, clarify some of the confusion from 18 the morning to hopefully avoid any process 19 concerns going forward. 20 So I misspoke on the rules about 21 quorum. For MAP; and this across MAP and CDP, NQF defines quorum as 66 percent of the eligible 22

voters, however to pass a motion we have defined consensus as greater than 60 percent of the eligible voters. So those are the thresholds to keep in your mind. Sixty-six percent is quorum; greater than sixty percent is consensus. So we wanted to break this down for you so you could just be tracking exactly what numbers we need here.

There are 32 members of the MAP

Coordinating Committee. Four of you are federal

liaisons, so you are not eligible voters; so you

come off the top if you're a fed, leaving us with

28 eligible voters. For our 66 percent threshold

of eligible voters that comes to 19 people are

needed to achieve quorum. If we do not have

quorum, we won't take a vote in the room. We

will follow up with you via SurveyMonkey after

the fact.

If you need to recuse yourself from a vote, you will be removed from the denominator.

Going along with the federal liaisons you are not eligible to vote. We define that as if you have

a conflict of interest, if you participated in the development of the measure, similarly to how Elisa described it this morning. We don't think anyone else has one for the rest of the day, so -- but please let us know if you feel like you might have a conflict and feel you need to be recused from the denominator of eligible voters.

Abstentions are included in the denominator for quorum, so abstaining is if you just choose not to -- you could vote on a measure, you choose not to for whatever reason. So we've just played out some scenarios for you for quorum. No number of abstentions will change what we need for quorum. Nineteen is our magic number. If we do have any more recusals, then you'll see people drop out. Our N would change from 28 to 27, so we would need 18 people around the table to have a quorum.

For consensus this is we are looking for greater than 60 percent of you to pass a motion. So again, we have 28 eligible voters.

We would need 17 people to hit a 60 percent

threshold for consensus. Here again recusals are removed from the denominator. For consensus we also remove abstentions from the denominator.

So playing out some scenarios, if we have one abstaining, we would need 17 people to pass a motion. If there are two abstentions, we need 16 people. Three, fifteen and then so on and so forth.

So let me just pause and make sure people are okay with this. Any questions? Any process -- we'll obviously have a session on fixing the process tomorrow, but before we get into some of the heavier voting I just want to pause and make sure people are comfortable.

John?

MEMBER BOTT: So this will come up in the hospital thing. So I was on a TEP for the hospital-wide mortality measure, one of the CMS TEPs. I don't think that's a conflict the way I read the paperwork. I wasn't a measure developer, but we were -- the TEP provided advisory -- had an advisory role to developing

1	the measure. So does that is that a conflict
2	where you shouldn't be commenting or is that
3	okay?
4	MS. MUNTHALI: Yes, that would be a
5	conflict, so we'd ask you to recuse yourself.
6	MEMBER BOTT: Oh.
7	MS. MUNTHALI: Yes.
8	MEMBER BOTT: Okay. Well, the
9	hospital session is later, right?
10	MS. MUNTHALI: Yes.
11	MEMBER BOTT: We're on the clinic one?
12	MS. MUNTHALI: So for that measure,
13	whatever measure is in front of you in which
14	you
15	MEMBER BOTT: Oh, wow.
16	MS. MUNTHALI: participated in the
17	measure development process, even in an advisory
18	role, you would be conflicted on that.
19	MEMBER BOTT: Yes, okay. Is there a
20	break before then? There are some kind of
21	logistical things to discuss. Maybe it's just
22	best one on one.

MS. MUNTHALI: Yes, we do have a
break.
MEMBER BOTT: Okay.
MS. MUNTHALI: So we can talk further.
MEMBER BOTT: Great. Thanks.
CO-CHAIR KAHN: We're going to do the
clinician first.
MEMBER BOTT: Yes, okay.
CO-CHAIR KAHN: And then we'll do the
hospital.
MEMBER BOTT: Okay. Thanks.
MS. MUNTHALI: And also just to let
you know, since the other measures are not
competing, you could vote on the other measures.
It's just that one measure in which you had
involvement in.
MEMBER BOTT: Okay. Thanks.
CO-CHAIR KAHN: Is everything clear on
the voting? I mean, it is what it is, so it's
I mean, it's the rules we work under, unless they
change, but they're not going to change today.
(No audible response.)

CO-CHAIR KAHN: Okay. So we'll now proceed. And let me remind those in the room or those on the phone who would like to make public comment regarding the Clinician Programs Task

Force that you should limit your comments to the clinician programs recommendations, that we need you to limit it to no more than two minutes and make comments on the MUC list or opportunities to improve the current clinician measures that are in the set at this time.

So I'll look back in the back of the room, and I see that we have a speaker. And could you identify yourself and then make your point? Thank you.

MS. RUBIN: Koryn Rubin, American
Medical Association. Just two general kind of
comments related to hopefully improving the MAP
process, one specific to the Clinician Workgroup
and the other that could probably be applied
broadly across all the workgroups.

First, in terms of the Clinician Workgroup there's been ongoing concern and

comment that's been made that there's a lack of clinician expertise on the Clinician Workgroup.

And what really gets discussed and evaluated in depth is based on whether that clinician or that physician specialty has a seat at the table, otherwise a lot of items really get overlooked and are not considered as part of the discussion when the public makes comment.

So for example, on the Clinician
Workgroup there are only four clinician
specialties represented on there, and as these
measures get -- become more complex, there really
is a need for more physician expertise and input
into what gets discussed and put forward by the
MAP.

Second is the need for consistency in terms of testing data. If CMS is going to require as part of proposing a measure to be placed on the Measure Under Consideration list, then all testing data like they require of the private sector needs to be available and put into place at the time of proposal. We know of

instances of many physician specialties proposing 1 2 measures on the MUC list last summer which was due July 1, and they were rejected because they 3 didn't have their testing data available. 4 5 And some said they could have it available by August and were told that that was 6 7 not acceptable, however, CMS was able to bring 8 forward measures that lacked testing data and in 9 some instances, particularly with the cost measures, only an oral update was provided and 10 the testing data has yet to be released today to 11 12 the public for evaluation. Thank you. 13 CO-CHAIR KAHN: Any other public 14 comments on the floor? (No audible response.) 15 16 CO-CHAIR KAHN: Any comments from the 17 group about the comments that were made? 18 thoughts? 19 (No audible response.) 20 CO-CHAIR KAHN: Does CMS have any 21 thoughts about what AMA said? Yes, Amir? 22 MEMBER QASEEM: I think actually --

thanks so much, Koryn, for bringing up some of these issues.

So I do actually have a follow-up questions for NOF. It was a little bit surprising for me. I was -- Multi-stakeholder's I absolutely think it's sound. We need them. But I got little concerned about hearing that there are only four clinicians in Clinician Workgroup. With that, I'm going to look at these recommendations a whole different light versus my -- because right now we're voting just yes and no keeping in mind that there is enough expertise. So four out of how many members are we talking about are in that group that are clinicians?

MS. MUNTHALI: So I think what we're trying to do is confirm, number one, the number of clinicians that are on the group, but I think the challenge we have is that we're trying to bring multi-stakeholders to together. So different perspective, not just clinicians, that can help us to make these decisions.

With that said, we're not really

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always sure what measures are coming forward to us through the MUC because we've constituted these groups well in advance of when the MUC is released, but we do try to make sure there's a balance. We may not have all of the specialties represented on there because we have funding for a finite number of people on the workgroups, but we are trying to be sure there's bounds. So we're trying to make sure we can confirm that number.

MEMBER QASEEM: So, absolutely. And
I'm --

MS. MUNTHALI: And we want to change that, yes.

MEMBER QASEEM: -- not questioning that you need to have every single sub-specialty represented there, but you still need to have enough number of threshold. But you're passing vote is 60 percent. We've all dealt with public members, and we have public members actually sitting here at the table, right? These measures get quite technical and I think to be fair is

lots that goes into the measure. Now if you're going to be using these measures for essentially evaluating the care that's being provided, I think you need to get the process right. But these measures are as only as -- if the process is right. So a little bit concerning now for me and actually --

DR. BAGLEY: Chip, my hand's up.

MEMBER QASEEM: -- when based on that, because after looking at some of the measures that I think I'm going to be pulling, now I get a better feel for it why those measures even make it through for our discussion today. But I think that my measures probably have been discussed at that group level.

CO-CHAIR KAHN: Okay. Thanks, Amir. From the phone?

DR. BAGLEY: Yes, Chip, this is Bruce Bagley. I'm the co-chair of the Clinician
Workgroup. And let me remind the Coordinating
Committee that most of these measures, especially the ones that have been through the NQF process,

have had a tremendous amount of very specialized clinician input way back in the process. So it's not like there's nobody who knows all the details that's had a chance to look at these before.

As you move up the line, whether it's to the Clinician Workgroup, or for that matter it's the Coordinating Committee, you're going to get people that are supposed to be looking at the big picture a little more than the details of the measure, because that's really already been done.

CO-CHAIR KAHN: Let me ask a question though. There are measures that aren't endorsed though, right? So they would not necessarily -- am I wrong?

DR. BAGLEY: That's correct. And there would be less, but most of those have actually been created by clinician groups or specialty organizations, and we've looked at those in the past. So you're right about that, but there's still a fair amount of work by clinicians that goes into the making of the measures or the early evaluation of the measures.

CO-CHAIR KAHN: Yes, Sam?

MEMBER LIN: Thanks, Chip. Just a point of clarification. On the Clinician Workgroup there actually are -- there are 13 physicians, about 3 or 4 who actually represents single specialty societies, but it's a broad base. I don't think we could have a workgroup with all of the specialty societies. Talk about chaos.

So I think it's -- to me there's sort of this effort to say what's the general consideration of physicians? And then you want to get into some specifics, perhaps some specialty input on those. But I think you've got to look at it generally, otherwise I don't think it will move at all.

CO-CHAIR KAHN: And you also have the dilemma I assume on the Task Forces -- are the Task Forces under the rule that you have to have a majority non-provider clinician, or are the Task Forces mixed?

PARTICIPANT: No, they're mixed.

1	CO-CHAIR KAHN: Okay. Because that's
2	an issue I guess in other places in the
3	hierarchy.
4	CO-CHAIR PINCUS: If need be, I mean,
5	we could send it around to the different groups
6	that are represented on the Clinician Workgroup
7	who'd want to see it.
8	MS. O'ROURKE: Why don't we pull up
9	the clinician roster?
10	Sheila, I just sent you an email with
11	it.
12	It's also on NQF's web site if anyone
13	wants to pull it up for themselves, but we'll get
14	it up on the screen in the room.
15	CO-CHAIR KAHN: Okay. And are we
16	ready, Sam? John?
17	MR. BERNOT: I was just going to
18	comment while we're bringing this up. I think
19	Koryn said there were four specialties, not four
20	clinicians, and I am we're verifying the
21	number of that. There is just looking through
22	the roster of actually credentials, it looks like

1	there's at least a dozen M.D.s or advanced
2	practice providers, or advanced care providers,
3	nurse practitioners.
4	So we'll get the final numbers, but
5	it's not four. I want to reassure the group that
6	it is the vast majority of the group does have
7	clinician credentials. We'll get the specialty
8	breakdown for you.
9	CO-CHAIR KAHN: Any more comments from
10	the floor?
11	(No audible response.)
12	CO-CHAIR KAHN: Are there any comments
13	from the public on the phone?
14	OPERATOR: Okay. At this time if you
15	would like to make a comment, please press star
16	then the number one.
17	(Pause.)
18	OPERATOR: No, no public comments at
19	this time.
20	CO-CHAIR KAHN: Okay. So we will
21	proceed. And I guess I'll ask John to go through
22	what's on the table and then I've got a couple of

1 things to say. The Committee will have the 2 opportunity to both put more things on the table or discuss further what we've got on the table. 3 4 MR. BERNOT: Great. Thank you so 5 And thanks, everyone, for being here and much. 6 welcome this afternoon back from lunch. 7 don't know me, my name is John Bernot. I am the 8 Senior Director on the Clinician Workgroup for 9 We also have two of our co-chairs on the phone, which I know Bruce Bagley you heard. 10 line is open. Amy Moyer is the other chair. 11 12 Amy, would you be able to just say 13 hello so that we know that your line is open and 14 you're able to hear us? Hi, this is Amy. 15 MS. MOYER: Sure. 16 Can you hear me? 17 MR. BERNOT: Great. Thank you Yes. 18 so much. 19 MS. MOYER: Thank you. 20 MR. BERNOT: So we'll just go over a 21 very brief overview of the two programs that the 22 Clinician Workgroup oversees, which is the MIPS

Program and the Medicare Shared Savings Program,
a little bit about the selection criteria, and
then a quick overview of the measures and the
results from the workgroup.

Next slide, please? So as you can see here, I mentioned the two programs: the MIPS Program, which is a merit-based incentive payment system, as well as the Medicare Shared Savings The vast majority of these measures Program. The 22 measures were in the were MIPS measures. MIPS Program and 3 in the Medicare Shared Savings I will note those three are also in the Program. MIPS. So there's 22 unique measures. Three of those are duplicate, which happen to be the three in the Medicare Shared Savings Program.

Next slide? So there were a few themes that came up and I think we're already starting to hear them from the public comments that I will just to try to highlight what the group spent a lot of time speaking on last December.

One of them was the cost measurement.

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This was something that there were eight cost measures that came through to the Clinician Workgroup, and there was a lot of discussion around this topic. One of -- there was an understanding and a discussion around the importance in any value-based program of being able to capture the cost aspect in addition to the quality aspect. So that importance was raised.

On the flip side, the concerns that came up with the cost measures were -- I shouldn't even say concerns, but the cautions were to ensure that these measures are appropriately risk-adjusted, both medically and from a social risk factor so that we have -- we know we have different populations, heterogeneous populations in some of these measures and try to make sure that that can level the playing field.

The last thing is since this is a newer style measure for MIPS was that whatever the decision is or whenever they are incorporated into the MIPS Program to make sure that these are

reevaluated on a constant basis. And that came up in the discussion a few times.

Next slide, please? The other topic that I think we had a lot of robust discussion on -- and you will see that there are some composite measures that were on the MUC list this year.

And there was -- on the plus side, once again these are really well-suited to capture the more holistic view of the patient's care, a little more comprehensive view of the performance for the MIPS Program.

The challenges were raised that we see for composite measures. We just talked about the attribution. We had a discussion about that attribution was one thing. So if there's multiple components to a measure, is one clinician able to effect the change on all of those components of the composite measure? So that was one of the discussions. And then just the technical challenges that the composite measures are more difficult measures from the measure development perspective to create.

1	Next slide? So before I go on I
2	should ask, Bruce or Amy, any other comments on
3	those two major themes or anything that I did not
4	mention?
5	DR. BAGLEY: No, I and I think that
6	covers it. I'd be glad to answer questions
7	especially about the composite measures and the
8	confusion that sometimes occurs between a
9	composite measure and a composite all or nothing
10	measure. They're quite different.
11	MR. BERNOT: Great. Okay. I'll
12	just
13	MS. MOYER: And this is
14	MR. BERNOT: Go ahead. I'm sorry.
15	MS. MOYER: Oh, sorry. This is Amy.
16	I have nothing additional to add.
17	MR. BERNOT: Okay. We'll continue.
18	We will have time for discussion, as was
19	mentioned.
20	This I will not read this slide to
21	you in its entirely. This was each year CMS
22	does put out the priorities and needs for the

programs, so this is the language that the -that guided our workgroup. And I mean, it's the
-- a lot of the topics that you've heard us talk
about before: outcome measures and the relevance
to the different specialties. There are some
high-priority domains which you can see, the
peer-reviewed aspect. So we're really seeing the
literature. And then not duplicative of -- in
other sets and really looking for the improvement
opportunities and keeping in mind the topped-out
measures. So that's what was provided.

The next slide just shows really the topics on there that really I thought were discussed at length within the workgroup. The outcome measure was definitely something that just -- that kept coming up. It was often brought up in the counter-discussion of the attribution. The composite measures we've already mentioned has been a topic that was one of the key themes of the meetings. Mentioned the importance of the efficiency and cost reduction measures. So these are falling in line quite

well with CMS' needs and priorities. And then the other thing with the appropriate use measures, but also considering the flip side of the inappropriate use on those topics.

Next? So very similar not surprisingly to MIPS, CMS puts out a list of the priorities and needs for the Medicare Shared Savings Program. A lot of these are very similar. Again, I will not read each one of these to you, but really are supportive of what we saw in the MIPS, but do have their own priorities and needs for each individual program, including the last one, measuring did align with the recommendations from the Core Quality Measures Collaborative.

On this I will -- since the three measures were duplicative, it's no surprise that these are identical findings: the outcome measures, the composites. And then that alignment with MIPS was brought up in our discussions about the Medicare Shared Savings Programs.

So that is just an overview of the two 1 2 And are we going to turn it over for programs. some discussion first and then present the 3 consent calendar? 4 5 CO-CHAIR KAHN: No. You mean about 6 the -- in terms of the general themes I guess the question is do we want to have discussion or do 7 8 we want to get into the calendar itself? It's up 9 to the -- up to you guys. Sam? 10 MEMBER LIN: General themes, if you 11 would, for a couple seconds anyway. 12 CO-CHAIR KAHN: Okay. Sure. We have 13 time. 14 MEMBER LIN: A couple things. Thank you, John. And I attended that workgroup in 15 16 December, and there was robust discussion 17 certainly and appreciate all their work. 18 A couple things. This paragraph at 19 the opening at the workgroup report about cost 20 measures, it's wonderful. And I think we ought 21 to have that same concern on the other 22 workgroups' reports as well. Why? Because one,

it's not just about cost. It's about accountability, transparency, equity, fraud, waste and abuse. And I think that concern ought to be laced by this group onto all the other ones.

No. 2, at the end you talk about the removal criteria. And then I look at the removal criteria on the other two workgroup reports. I don't know why they should be different. We're supposed to be about alignment and harmonization. And whether we remove it from PAC or hospitals or clinician, we remove it. What's the difference? I think -- and I think it was best written again in the clinician report, so I would encourage us to think about that.

The third part of my diatribe at this point is saying that here we are since 2011 into alignment and harmonization, and what's frustrating to me is that for example today we're doing 22 items on MIPS, a program that MedPAC has said is essentially dead in their books. So what are we going to do when they pronounce it and it

doesn't resurrect? Well, then we're going to either can that or we're going to have to wait until the next program comes out, go through the process again because that's what law requires, and do the same work again.

The count that we had on this particular meeting was 32 measures across the board, but when you actually sit and count the chart, it's 37, and the reason being there is duplicity across some of the programs. That's not alignment. That's not harmonization.

In my own situation -- so I've got somewhat controlled diabetes and my hemoglobin Alc is 6.5 last time and this morning my blood sugar, fasting blood sugar was 109. That's all that counts. I don't care whether it was Medicare Advantage or a model home or whatever you want to call it, or MIPS or MSSP, et cetera. We should be aligning, not trying to cater to separate different programs the same blessed measures. And I'll go through that diatribe over and over again. But our initial charge from 2011

is align and harmonize, not create more gaps and more distinct differences. End of story. Thank you.

CO-CHAIR KAHN: Okay. Other comments?

(No audible response.)

CO-CHAIR KAHN: Okay. So we've got certain measures that have been, in quotes, "pulled," which we'll go through. We then I guess create a list, which is the ones that were pulled and then any that the Committee here would like to add to that pull. Then we will I guess have a discussion on each, and when we discuss each of the ones that were pulled, a member of the Committee can choose to do a motion to change the status of whatever it is.

In terms of the measures that remain on the consent calendar that have not been pulled either by someone for this meeting or -- I mean, in an way, then when we finally vote on the total package, the presumption is that those will be the ones we're voting to approve, unless we've changed the status of any that I -- as I

described. I think I've gotten our process down. 1 2 And we have John Bott and Derek Robinson and Carl to be discussants. And then 3 4 obviously Bruce is on the phone, and staff is 5 here. So when we go through the list, I 6 7 guess you, John, will describe sort of the basic 8 circumstances about why it was pulled and if a 9 change was made why there was a change made. But to create a complete list, does 10 11 anyone around the room want to pull any beyond 12 the ones that we know are pulled, or do we need 13 to go through that so everybody -- everybody 14 needs to go through it? Okay. Let's go through the list that 15 16 have been pulled and if anyone wants to pull 17 others, they can do it. 18 MR. BERNOT: Sure, no problem. I'll give just some very quick stats on the --19 20 what the status of the measures are. I will list 21 the ones that have already been pulled. I will

start with the MIPS Program, but I do want to let

you know they are all on one consent calendar, so whatever is left there that is not pulled off would be for both programs. There is not a MIPS consent calendar and a separate MSSP consent calendar. What is left would be for both programs. So if there's any questions, we'll keep going on with the voting and help out.

But for the MIPS Program, as we mentioned, we had 22 measures. They had 3 that were support, 8 with -- 18 with conditional support and 1 that had refine and resubmit. Seven of those have been pulled, and I will read those seven right now.

The first one is the continuity of pharmacotherapy for opioid use disorder. We will have the chance to talk about each one of these, so I'll just list the ones that have been pulled so you know, if there's anybody who wants to pull other measures.

The next one is the optimal diabetes care, which is -- I should give you the number also -- which is MUC17-181, optimal diabetes

Next is MUC17-194, optimal vascular care.

Next is MUC17-215, which is diabetes Alc control

less than 8.0. Next, MUC17-234, ischemic

vascular disease, use of aspirin or antiplatelet

medication. MUC17-262, STE-elevation myocardial

infarction, STEMIs for percutaneous coronary

intervention. And lastly for the MIPS Program is

the MUC17-363, intracranial hemorrhage or

cerebral infarction. So those are the MIPS

measures that have been pulled.

I guess at this point perhaps we should see if anyone from the MIPS would like to pull, and then I will go over the MSSP.

CO-CHAIR KAHN: Yes, is -- any other -- anybody that would like to pull from MIPS?

Amir?

MEMBER QASEEM: I just want to make sure that I'm -- there might be some repeats, so I wanted to have 181, 215, 194, 210. I'm trying to figure out how many should I pull. Sorry.

Three-sixty-three was pulled. Three-sixty-seven.

Did I already say 310?

1	CO-CHAIR KAHN: Yes.
2	MEMBER QASEEM: Three-ten as well.
3	Three-one-zero.
4	MR. BERNOT: So for the ones that
5	you're proposing to pull that have not already
6	been pulled do you want to pull them for
7	discussion or do you want to pull them for a
8	different motion?
9	MEMBER QASEEM: Well, I'll actually
10	probably have a different motion.
11	MR. BERNOT: Okay.
12	MEMBER QASEEM: Because my problem is
13	that conditional support has always worried me
14	actually. Conditional support ends up getting
15	implemented. That's the bottom line, right? So
16	any of the conditional supports many of them
17	actually are conditional support. Conditional
18	support and maybe you guys can educate me
19	again. Conditional support measures end up
20	getting implemented. And that's why I'm being a
21	little bit more stricter on that one.
22	So actually most of them are

1	conditional support and that's why I actually
2	have a different motion, unless I can get some
3	sort of another point of view that conditional
4	support does not mean that they're going to get
5	implemented.
6	CO-CHAIR KAHN: Okay. So to deal with
7	this in an ordered way, and since I don't know
8	if everyone has all the paper. Can how many
9	do we have total?
LO	MR. BERNOT: That would be two more.
L1	I don't I think did you mean 310 when you
L2	said 210? I don't believe there is a 210.
L3	CO-CHAIR KAHN: I think
L <b>4</b>	(Simultaneous speaking.)
L5	MR. BERNOT: Okay. Right. Yes, I had
L6	210 and 310. Did I miss it?
L7	MS. BUCHANAN: That's in the hospital.
L8	MR. BERNOT: So there would be two
L9	additional measures. That would be Measure 310
20	and 367, both which would need a motion as to
21	what the new motion would be.
22	CO-CHAIR KAHN: Okay. And how many

were pulled total?

MR. BERNOT: Prior to that we had seven pulled, so this puts it to nine.

CO-CHAIR PINCUS: Okay. So I suppose if the Committee will allow us then, why don't -- I guess we should go one by one from -- I mean, by the numbers, and discuss each one and then just make sure with each one whether there's just interest in discussion or -- but let's just -- let's sort of -- so what's the first one?

Oh, I'm sorry. Did I skip anybody who had any to add? I'm sorry. Bruce?

MEMBER HALL: Well, I have more of a question than wanting to push forward one measure, but procedurally then if we don't pull -- like of these 22 or whatever we've pulled 9. There's 10 or 11 others that are also recommended for conditional. If we don't pull them, we are saying we agree with the conditional comments that the prior committee entered.

CO-CHAIR KAHN: Yes, anything that's on the consent calendar, and the consent calendar

is now made up of all the recommendations of the Task Force, that are not pulled, they will be voted on in the end as they are in the calendar. So if you want to have discussion and possible action on any of them, any of the measures that are not already on the list for discussion, then we'd have to pull it to change it. Otherwise, it will be considered in the final vote.

Now the final vote is a vote, so -but presumably by the time we get there, we'll
have sufficient votes. We'll have to wait and
see.

MEMBER HALL: Okay. So some of the conditional comments though are nebulous.

CO-CHAIR KAHN: Well, let me add this, that as we said in the introduction that if you want to pull a measure, that doesn't mean that we have to take action on it. We could just have a discussion on it so that if there's a measure that you think you don't want to fight city hall on the conditional, but you think it requires a little more discussion and at least some comment

which would then go along with it as we pass things off to CMS, then that would be a reason to pull it. You don't have to have a motion on it.

So I leave it to you to decide whether there are any --

MEMBER HALL: I'm just trying to get the procedure straight. So like I'm just reading one example where it looks like the comments have already been put into our perspective, the comments saying MAP recognized, MAP did this, MAP did that. And that is meant to reflect that we did --

CO-CHAIR KAHN: Right the Coordinating
Committee, both to simplify our process and allow
us to come to a conclusion -- and actually the
Task Forces have a base to work from, the consent
calendar. The consent calendar was created
initially by the staff taking the measures
through the algorithms and coming up with a draft
recommendation based on the algorithms. The Task
Force then can change that recommendation, and in
some cases it does.

1	And then it comes to us. And then we
2	have the prerogative well first others can
3	pull prior to here that we have the prerogative,
4	the Committee members, to pull others, but we
5	only consider either for discussion and/or for
6	action further discussion and/or for action
7	the item is pulled. That's the procedure.
8	MEMBER HALL: All right. I'm sorry.
9	I misunderstood. I had sort of assumed that
LO	anything that was conditional we would at least
L1	review the language around that, but that's not
L2	the case.
L3	CO-CHAIR KAHN: Well, no, no. We can
L <b>4</b>	review the language, but you got to pull it to
L5	MEMBER HALL: I understand.
L6	CO-CHAIR KAHN: have a discussion.
L7	That's what I'm saying.
L8	MEMBER HALL: Okay.
L9	MEMBER ROBINSON: Just for final
20	clarification on the procedures, so if the
21	consent calendar says that we're going
22	recommendation is for resubmission and we vote to

approve it, those that are voted for approval would be approved. Those that were voted for or recommended for resubmission, would be for resubmission. Is that correct?

CO-CHAIR KAHN: Correct.

MEMBER ROBINSON: Okay.

CO-CHAIR KAHN: So that the -whatever is on the consent calendar when we vote at the end, we'll be voting on that unless it's If it's pulled, there can be two things pulled. One, we can just have a discussion which would go into the record on it, and then whatever the recommendation of the Task Force is would go in the -- back into the consent calendar. Or if people want to change the recommendation, then we can have a motion, have discussion and have a vote and see if the recommendation would be something different than was made previously. we're putting together a package basically that's been already prepared for us.

John?

MR. BERNOT: Sure. So unless there

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was anything else, any other we had two more
measures from MIPS pulled. I'll just very
quickly go over the MSSP measures, which is a
little bit easier. Again, three measures on
here. They all received conditional support. So
far two of them have been pulled of the three
measures. The remaining measure that has not
been pulled from MSSP is the MUC17-234. Yes,
that was pulled by MIPS, but it has not been
pulled for discussion for MSSP yet. And they
would need a separate vote for each program. So
if
PARTICIPANT: I'm sorry. Those two
measures are it's the
MR. BERNOT: It's the same measure for
two different programs, correct. The one pulled
for MIPS was pulled by pre-pulled by Bill
Kramer, but the one from but it only lists the
MIPS. It does not list the MSSP as having been
pre-pulled.
CO-CHAIR KAHN: You mean it is
identical?

1	MR. BERNOT: It is identical. I don't
2	know if there was a reason that he only wanted it
3	in one program or not.
4	PARTICIPANT: Yes, there is a reason.
5	I have to know the reason. He does not
6	CO-CHAIR KAHN: Then I'll pull it.
7	MR. BERNOT: Sure. All right. So
8	PARTICIPANT: We do have a reason.
9	CO-CHAIR KAHN: Then when we consider
10	it, we consider it for both programs at the same
11	time.
12	MR. BERNOT: We would talk about
12 13	MR. BERNOT: We would talk about the way we had done it the clinician program, we
13	the way we had done it the clinician program, we
13 14	the way we had done it the clinician program, we talked about it in one discussion and then took
13 14 15	the way we had done it the clinician program, we talked about it in one discussion and then took two votes. After that discussion, is there
13 14 15 16	the way we had done it the clinician program, we talked about it in one discussion and then took two votes. After that discussion, is there anything that would change your mind on that
13 14 15 16 17	the way we had done it the clinician program, we talked about it in one discussion and then took two votes. After that discussion, is there anything that would change your mind on that it belongs in one program versus the other?
13 14 15 16 17	the way we had done it the clinician program, we talked about it in one discussion and then took two votes. After that discussion, is there anything that would change your mind on that it belongs in one program versus the other?  CO-CHAIR KAHN: Okay. Well, I'll
13 14 15 16 17 18	the way we had done it the clinician program, we talked about it in one discussion and then took two votes. After that discussion, is there anything that would change your mind on that it belongs in one program versus the other?  CO-CHAIR KAHN: Okay. Well, I'll it's pulled, so

1 CO-CHAIR KAHN: Now let's proceed in 2 an orderly way and -- hopefully, and we'll go to the first measure that was pulled. 3 MS. MOYER: This is Amy. Can I --4 sorry, it's hard to raise my hand virtually. 5 6 CO-CHAIR KAHN: Oh, sure. Just speak 7 up, Amy. Thanks. 8 MS. MOYER: As I hear the measures 9 being pulled and some of the discussion around 10 that, I know something that we struggled with in 11 our workgroup and I've heard that other 12 workgroups struggled with was what exactly 13 conditional support meant versus revise and 14 resubmit. And so I'm wondering; and I apologize because I came in at the clinician portion, has 15 there been a discussion of the one status versus 16 17 the other with this group to make sure that the 18 questions are stemming from disagreement and not 19 from misunderstanding or a lack of clarity? 20 CO-CHAIR KAHN: Let me do this: 21 First, we'll have discussion tomorrow about --22 not that it helps us right now, but about how we

go forward in the future on these items. In order for consistency with this, I'm going to look at the staff and ask them to provide us with their perception at the Task Force level of what the meaning of those two categories were because they were at each of the levels and I wasn't. So I'd prefer them to explain.

And then let me just say we could have discussion on it right now, but my view would be and my hope would be that the Committee will just go with whatever the staff definition is, because otherwise we're just going to go around in circles.

So would the staff provide their working perception for the purposes this afternoon of conditional versus -- of the categories?

MS. O'ROURKE: Sure. So in the algorithm that staff used to make our preliminary recommendations we really use if a measure has been fully tested as the line in the sand for if the algorithm gives it a conditional support or

it a refine and resubmit, so the preliminary analysis the workgroup received based off of that. If it was not tested, it went into this refine category.

The workgroup's operationalized the refine and resubmit as a category to reflect when a measure needed a significant change. They wanted to see testing completed. They had concerns that it was tested at one level, but not at the level of the program that was being under -- being considered for. So they wanted to reflect that. Any other sort of significant change they would like to see the measure have made before it goes into the program.

Conditional support was more of a minor thing, if you will, usually around gaining NQF endorsement. Refine and resubmit they operationalized as something more significant that would perhaps require the measure to go back to the developer and have something re-specified.

CO-CHAIR KAHN: Let me ask a question.

How many times was conditional support given for

reasons other than it lacking endorsement, and 1 2 what were those reasons? MS. O'ROURKE: So I don't have a hard 3 4 number off the top of my head. CO-CHAIR KAHN: Oh, just roughly. 5 Maybe there were 5 out 6 MS. O'ROURKE: 7 of the 35 total where they attached a different 8 condition other than NQF endorsement. Some that 9 come to mind were for some of the hospital measures that you'll review this afternoon, a 10 11 recommendation that it be considered by the NQF 12 Methods Panel even though it may be closer to a 13 process measure than an outcome measure, some 14 concerns about examining the need for say SES adjustment some -- during the endorsement review. 15

CO-CHAIR KAHN: Pierre?

So any kind of special consideration about the

DR. YONG: Well, I was wondering whether -- and I don't mean to put Harold on the spot, but -- because this discussion did happen at the webinar discussion about how to use refine

measure.

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and resubmit in terms of guidance for the
workgroup committees, which I think, Harold, as I
recall you were the one that sort of verbalized
that. But Harold also had the benefit of
actually sitting in on the Clinician Workgroup,
and so how they sort of operationlized and
applied that sort of guidance. I was wondering
-- it's up to you, but I thought it may be
helpful for folks to hear from you.

CO-CHAIR PINCUS: Well, I mean, the way I conceptualize it; and it may not be the way it's intended to, is conditional support, if there's a very specific condition that needs to be met and if that condition is met, they would support it. So the modal version of that is NQF endorsement. There may be some other potential little things that are very specific to a particular measure that might -- are in the potential in the short term to be fixed in some way, but that was it.

Anything that needs more work basically that has sort of multiple issues or

needs more work in general would be refine and resubmit. That's the way I've been interpreting it.

it back to Amy in a second, but -- and the problem actually with both of them I think is for some people to say; well, and for me, too, is the feedback loop, because in both cases there's a -- there's something -- the action needs to be, at least from the view of the Task Force and/or the Coordinating Committee, action needs to be taken on that. And I guess the dilemma for some around the table is that we obviously can eliminate recommendations, but CMS will choose to go forward with them sometimes regardless. But that's where we are.

Amy, do you have more to add?

MS. MOYER: I was just going to add that an additional reason that for a couple of these we used conditional support was that there are existing measures in the program that are competing measures. And our understanding from

CMS was that those other measures will be being 1 2 removed from the program, so there won't actually be duplications, but we wanted to kind of 3 4 formally capture that in our recommendation. 5 other than NOF endorsement that was also a theme in our conditional support. 6 7 CO-CHAIR KAHN: Okay. That's actually -- how many times do you think that happened? 8 9 MR. BERNOT: So in terms of this, looking -- the condition being that we wanted to 10 11 ensure that there was no duplication. And that 12 was considered. It happened on I believe three 13 measures. It would be the optimal -- both the 14 optimal vascular and optimal diabetes, the Alc and the antiplatelet. So I think that's four 15 16 measures of which some of those were duplicate in 17 MSSP and MIPS. 18 CO-CHAIR PINCUS: To fit the criteria 19 I'd put forward that this is a very specific thing. 20 And if that were done, then it would be 21 supported.

Okay.

At least to

CO-CHAIR KAHN:

proceed, is there then a general understanding of what these different categories mean? We can deal tomorrow with the results I think in terms of conditional and refine and resubmit as to what we would like CMS to interpret them to mean. We can talk about that tomorrow.

I'm sorry, Bruce. I didn't mean -
MEMBER HALL: So just final

clarification then. Several of these definitely
say conditional pending NQF approval. And then
they go on to say when it's in the endorsement
process MAP would request the following things be
examined. But regardless, if the NQF endorsement
did not happen, the conditional is taken away,
this is communicated again, or how is it handled?

CO-CHAIR KAHN: I'll look at the staff
and ask them what they do.

MS. O'ROURKE: So that's an excellent point. So the way the MAP process is currently set up it's really only looked at once. I think if the measure were to have that happen where MAP conditionally supported it and then it failed the

endorsement process, that feedback would go to CMS and they would need to address it when proposing the measure, that it was not ultimately endorsed. But I think we don't really have a process for adjudicating that kind of disparity between CDP and MAP, but I think that's perhaps something we could put on the feedback loop conversation for tomorrow as how we check that and what that means.

CO-CHAIR KAHN: Yes, I think that we really ought to think about that. We will have an opportunity to talk about that tomorrow. And if -- and obviously our recommendations go over with the conditional language and -- but then if things are put in place in April, or I guess whichever month you do the other -- the regulations for the clinicians, and it's there and it still hasn't been endorsed, I think we do need to figure out at what point the Coordinating Committee or somebody sends a letter to CMS of either concern or question as to why something hasn't happened.

I mean, maybe we need to do that.

Because in some ways you could say it's maybe partly incumbent upon us when we're doing something conditionally to at least have a notion of some length of time and then to go back, because CMS could argue, well, it's in the process. We have to -- we have our regulation.

It's -- we got to meet our deadlines. And we're in the endorsement line and we can't wait. I mean, and there's no argument because the HHS Secretary has the power to do these, whatever we

So I think we need -- I think it's incumbent upon NQF to come up with some process to at least come back and query or set something in motion so we could bring something back to the Coordinating Committee. And I mean, obviously CMS can do what they decided to do. But we -- I think this is a -- in some ways I guess it's our fault. I think that we don't have the feedback loop that we ought to.

MEMBER ROBINSON: Mr. Chairman, I

recommend.

agree with your comments.

And to the staff, is it possible to let the Task Force know what the approximate date of the NQF endorsement process -- when that would actually occur and be considered for the measure relative to the regulatory process just so that the recommendation can be offered with a little bit more certainty. So if we anticipate that we won't know what the endorsement outcome is prior to the regulatory decision in summary we recommend that you go forward with it versus not?

MS. O'ROURKE: So I think we don't necessarily know what you're -- CMS is considering a measure for or when they would be wanting to propose a measure, so there's a little bit of uncertainty. But I think we could look at what you're saying especially with our redesigned CDP process where we have consistent calls in the spring and the fall of each year and perhaps highlight when there may be opportunities to submit a measure.

I know the workgroups were questioning

that a bit more as they realized the changes to the CDP process, and I think we're asking the developers some more pointed questions over when exactly will you be bringing this in now that the endorsement is consistent, when there would be the opportunity to submit rather than having to wait for a call.

So I think that's a great point,

Derek. We can try to work that into how our

calendars map with the rulemaking calendars and

what information could potentially be available

and try to deal with some of that nebulousness

that -- about the timing.

measure that gets conditional is ready for prime time other than it may not have endorsement or it may be duplicative, but it's a measure that's ready to go. And we don't know that CMS would put it in the next cycle, but if it's ready to go, presumably they probably would in most cases, whereas the other category is quite different if it's not fully tested or whatever.

Okay. Any other discussion? 1 2 PARTICIPANT: Elisa has housekeeping, some disclosures. 3 4 MS. MUNTHALI: Yes. Yes. 5 CO-CHAIR KAHN: Okay. MS. MUNTHALI: So we do have two new 6 7 members that have joined us, and before we 8 continue discussion and vote we wanted them to 9 orally disclose. So Mary Barton from NCQA and Mary Beth from the American Nurses Association. 10 11 We'll start with Mary Barton. 12 MEMBER BARTON: Thanks. This is Mary I'm the Vice President for Performance 13 Barton. 14 Measurement at the National Committee for Quality 15 Assurance. NCQA is a measure developer and we 16 design measures that principally are used in 17 health plan evaluation, which we do because we 18 accredit health plans, both Medicare, commercial 19 and Medicaid. Thanks. 20 MEMBER BRESCH WHITE: Mary Beth Bresch 21 White, Director of Health Policy at ANA.

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nothing to disclose.

CO-CHAIR KAHN: Okay. Are we ready?

(No audible response.)

CO-CHAIR KAHN: Okay. So now we're going to go down and with each one John will lay it out. Comment. We have commenters who can comment. Bruce can comment. And I guess then it's incumbent upon the people that had -- whoever had a problem with it to either discuss it further or make a motion.

MR. BERNOT: Great. Thank you, Chip.

And I know that was a little bit confusing. We

-- I do believe we have everything that we need.

We'll be able to walk through this in an

organized manner now that we know which measures

we're looking at. I'll do my best to let you

know what the workgroup had -- was thinking in

their thoughts, what the motion is if we have a

motion. Or if we need a motion, we'll lay that

out. And then we can have the discussion.

So the first measure to discuss is MUC17-139, top of the list here. This is the continuation of the pharmacotherapy for opioid

use disorder. The workgroup discussed this and gave it a recommendation of conditional support for rulemaking with the condition that it's tested and endorsed at the clinician and clinician group level.

They encourage the Relevance Standing Committee to specifically evaluate the attribution method, reliability and validity of this measure at the individual clinician and practice level. The reason for that was that this measure was not tested previously at the clinician level.

So the motion -- this was -- this measure was pulled by Carl and it -- the motion that was on the table was to change the conditional support to a refine and resubmit category. Rationale was that it's only been specified tested and endorsed at the health plan level. Because it's not yet developed and tested at the clinician or practice level, we do not believe it should receive conditional support.

CO-CHAIR KAHN:

Neal R. Gross and Co., Inc.

Washington DC

Okay.

Discussion?

MEMBER SIRIO: Well, I mean, you in part articulated the argument that I gave you guys in an email, which is you have the same discussion. This is consistent I think with Amir's conversation, which is if it's not quite ready for prime time, it's not conditionally approved. I mean, it should be -- we should take a step back. So I'm making the motion that we refine and resubmit rather than conditional support.

CO-CHAIR KAHN: John?

MEMBER BOTT: Yes, what Carl is suggesting makes sense to me. And it gets into this -- again, just to articulate what my understanding of this definition of "conditional" and "refine" means, it's not simply resubmitting to -- submitting it to NQF, getting it tested and getting it endorsed. It needs to be refined so that attribution can be appropriately conducted at the physician or group level.

I've been involved in measurement at the medical group level and at the health plan

level, and it's a whole other kettle of fish to come up with attribution for the physician or group level than the health plan level. So it's not as simple as send it to NQF, get it endorsed with testing because it has to be reconfigured to make it work at that level. So that -- I'm just expressing that's my understanding of the definition of particularly the word "refine," because it does need to be refined, not just simply sent to NQF with testing done. So it's -- so right now I support Carl with that understanding on some of those words.

CO-CHAIR KAHN: Okay. Are there any other comments?

MR. BERNOT: Just to your point, I just wanted to -- one thing I did not mention was even though this did not happen frequently, this is one where the staff preliminary assessment was refine and resubmit. Due to the importance of the epidemic is the reason -- was part of the discussion for the Committee to change it to a conditional support. Just wanted to state that.

1	CO-CHAIR KAHN: Good. Okay. So
2	CO-CHAIR PINCUS: Just I'm sorry.
3	Just one clarification. And this is an endorsed
4	measure, correct?
5	MR. BERNOT: At the health plan level.
6	PARTICIPANT: Not at the health plan
7	level.
8	CO-CHAIR PINCUS: No, no, but it is an
9	endorsed it's an endorsed measure so that it
10	wouldn't make sense to re-endorse it in a sense,
11	but you'd want to but, no, it just needs to be
12	refined. It just it needs to be refined to
13	apply it at a not at the health plan level.
14	DR. AMIN: Right, I just want to be
15	clear that the endorsement is for the measure
16	specifications at the health plan level. So we
17	wouldn't I mean, to state that it's endorsed
18	is accurate, but it's endorsed at the health plan
19	level. It's
20	CO-CHAIR KAHN: Right, but I think
21	that's an important point, that one can have a
22	conditional to for it to be endorsed even

1	though it has been endorsed for something else,
2	but endorse it for this particular purpose.
3	MR. BERNOT: I mean, it fits for
4	purpose here.
5	CO-CHAIR PINCUS: Right.
6	CO-CHAIR KAHN: Derek, do you have a
7	comment?
8	MEMBER ROBINSON: I concur with the
9	comments that were made as a discussant on this
10	particular measure.
11	CO-CHAIR KAHN: Okay. So, Carl had
12	made a motion. I need a second. Do I hear a
13	second?
14	PARTICIPANT: Second.
15	CO-CHAIR KAHN: Is there any more
16	1, ,
	discussion?
17	(No audible response.)
17 18	
	(No audible response.)
18	(No audible response.)  CO-CHAIR KAHN: The motion is to move
18 19	(No audible response.)  CO-CHAIR KAHN: The motion is to move this to the resubmit category. I guess does our

1	CO-CHAIR KAHN: Well, let's just
2	MS. OGUNGBEMI: It works for me, so
3	hopefully
4	CO-CHAIR KAHN: Let's take this
5	vote will be a test. So I'm going to pass it off
6	to the staff to give us instructions on the vote.
7	And those on the phone, they should they know
8	how to submit by email.
9	Are we ready for a vote?
10	(No audible response.)
11	MS. OGUNGBEMI: And we're just going
12	to have it displayed also. It's on the back.
13	And people participating via phone, please email
14	your vote in.
15	Mira, I just received yours. Thank
16	you.
17	CO-CHAIR KAHN: Okay. You ready?
18	MS. OGUNGBEMI: It should come up in
19	a moment. Hold on one second.
20	CO-CHAIR KAHN: Okay. We're going to
21	give it 10 seconds and then okay.
22	(Pause.)

1	CO-CHAIR KAHN: Okay. Does it look
2	good?
3	MS. OGUNGBEMI: Hold on. This is the
4	test. This is a test.
5	CO-CHAIR KAHN: Okay.
6	MS. OGUNGBEMI: Okay.
7	CO-CHAIR KAHN: Everybody, you ready?
8	MS. OGUNGBEMI: Hold on. Hold on.
9	Let this clear out.
10	Okay. Here we go. Polling is open.
11	CO-CHAIR KAHN: Okay. One is yes; two
12	is no.
13	MS. O'ROURKE: And, Maureen, we also
14	received yours. Thank you.
15	MS. KAHN: Good to know.
16	CO-CHAIR KAHN: Did it work?
17	MS. O'ROURKE: So it looks like we're
18	waiting for a couple more people to come in. So
19	it would be we're looking for 25.
20	CO-CHAIR KAHN: We're waiting for
01	II
21	people on the Internet or people here?

1	for patient population here. I have Mira and
2	Maureen's from on the phone. We also have
3	Stephanie Glier substituting for Bill Kramer.
4	She sent in a no vote.
5	CO-CHAIR KAHN: Okay. So that
6	wouldn't be here, though?
7	MS. OGUNGBEMI: So she needs to
8	Stephanie, could you please email
9	mapcoordinatingcommittee@qualityforum.org?
10	MS. GLIER: Yes, thanks.
11	CO-CHAIR KAHN: Okay. How are we
12	doing?
13	MS. OGUNGBEMI: We're doing great. We
14	have 24, waiting for one more.
15	CO-CHAIR PINCUS: One more here or one
16	more
17	NG OCIDICADANT No. de al
	MS. OGUNGBEMI: No, just
18	PARTICIPANT: Remotely.
18 19	
	PARTICIPANT: Remotely.
19	PARTICIPANT: Remotely.  CO-CHAIR PINCUS: Remotely?

1	MS. O'ROURKE: We got it.
2	CO-CHAIR KAHN: Okay.
3	MS. O'ROURKE: So we're working now.
4	CO-CHAIR KAHN: And so what's the
5	vote?
6	MS. O'ROURKE: So this was just a
7	test.
8	CO-CHAIR KAHN: I thought we were
9	testing by the vote. Okay. Well, let's I
10	understand. Let's hurry this up. We got a lot
11	to do this afternoon. If this is going to work,
12	it's got to work now or we're going to vote by
13	hand.
14	MEMBER QASEEM: So it's we are
15	voting refine and resubmit, to support refine and
16	resubmit?
17	CO-CHAIR KAHN: Right. Right.
18	MEMBER QASEEM: So I vote yes, that
19	means
20	(Simultaneous speaking.)
21	CO-CHAIR KAHN: A yes vote is in favor
22	of the motion to change the categorization for

1	this to the refine and revise and resubmit
2	rather than conditional. That's what the vote is
3	about.
4	MS. OGUNGBEMI: Okay. We are voting
5	on MUC17-139. The motion is refine and resubmit.
6	Yes to refine and resubmit; not to not support
7	that motion.
8	CO-CHAIR KAHN: Okay. Great.
9	MS. OGUNGBEMI: Voting is open.
10	(Voting.)
11	CO-CHAIR KAHN: Let's see what the
12	result is.
13	PARTICIPANT: We'll vote until we get
14	it right.
15	CO-CHAIR KAHN: Yes.
16	(Laughter.)
17	CO-CHAIR KAHN: We got the 25. Oh,
18	okay. And we got
19	MS. O'ROURKE: So our N is 25.
20	CO-CHAIR KAHN: What happened to it?
21	Well, I saw 80 percent, and 80 percent is more 60
22	percent.

1 MS. O'ROURKE: Okay. Eighty percent 2 yes to support refine and resubmit. CO-CHAIR KAHN: So we have a consensus 3 4 by Bill Kramer's definition, or anybody's 5 definition. So let's go to the next one. MS. BUCHANAN: And this is Kate 6 Just a quick housekeeping 7 Buchanan. 8 People on the teleconference are announcement. 9 having a very challenging time hearing people, so 10 please make sure to speak very loudly into your 11 microphones so that everyone can participate. 12 Thank you. 13 CO-CHAIR KAHN: Yes, make sure you're 14 always putting your red button on. Maybe that's 15 a problem, too. And then make sure you turn it 16 off when you finish. Probably only allows so 17 many at a given time. 18 Okay. John, let's go. 19 MR. BERNOT: All right. One down. 20 So we are on -- moving to MUC17-181. 21 This is the optimal diabetes care measure. 22 has been pulled for both MIPS and MSSP. I will

give you the quick read of the Committee's recommendations, and they are very similar between the two programs.

So the recommendation for this is conditional support for rulemaking. And I'm going over the MIPS recommendation. The condition is that there are no competing measures in the program and that the measure is updated to the most current clinical guidelines. And that is in reference to the fact that there's a blood pressure reading in the composite measure.

That's the MIPS.

For the MSSP it is the exact same recommendation, conditional support with the same condition.

So as a matter of process we'll have one discussion, but we will need to have two votes on this. And I will tell you why this was pulled. This was also pulled by Carl for further discussion. And since Carl's here, I'll just in the interest of time allow you to state your --

(Simultaneous speaking.)

MEMBER SIRIO: Yes, so real briefly.

I'm not suggesting we change the motion on this

for either of the programs. The concern

basically is -- and I think Sam alluded to it in

his personal commentary with respect to his

capacity to manage his diabetes.

I mean, there are going to be sociodemographic factors and other health factors that are going to play into the capacity of an individual to in fact hit targets, so what we're advocating for is that when CMS does this, it takes a hard look at actually appropriately restratifying the folks that in fact are being measured.

CO-CHAIR KAHN: I'm sorry, stratifying --

MEMBER SIRIO: The re-stratification issue, which is not clearly addressed. It's a comment basically that it probably warrants a discussion if CMS is going to move forward. I'm not going to change the recommendation. It's just a comment on the specification of the

measure.

increased 33 percent.

other commenters have any other comment? Amir?

MEMBER QASEEM: So I actually -- I

agree what you said, but I'm more leaning towards

actually changing the vote on this as well,

because I just pulled a slide, a CMS slide,

Medicare beneficiaries, a record of under
treatment versus over-treatment for diabetic

patients. Hyperglycemia admissions decreased by

39 percent for the period of past -- since 1999

until 2014 and hypoglycemic events admissions

CO-CHAIR KAHN: John, or any of the

So you can see there's an issue with this. The bottom line is that eight is not an absolute number. It might be okay for certain younger patient population. Actually you probably want to go lower. But for certain other aging populations you would go higher. And this, the CMS Medicare data is showing that there is a difference that under -- actually hyperglycemic events decrease, but hypoglycemic events

1	increase, and hypoglycemia is more dangerous if
2	you look at any of the new research.
3	So I'm more leaning for that this
4	measure is again I think it's going to cause
5	more harm than benefit over here. So I'm
6	actually have a motion that we need to change
7	this as well to revise and resubmit, or
8	something
9	(Simultaneous speaking.)
10	CO-CHAIR KAHN: Okay. So let's
11	well, actually just to make it clean, so Amir's
12	made a motion. Is there a second to the motion?
13	MEMBER MULLINS: Second.
14	PARTICIPANT: I would agree. I
15	support it.
16	CO-CHAIR KAHN: Okay. So now we're in
17	discussion around the motion, which would be to
18	actually go beyond simply a comment from us to
19	actually changing its status.
20	So, Amy, will you
21	MS. MOYER: Yes, I agree with Amir.
22	When you look at this measure, you have to

realize this is not just Type 2 diabetes. This is also Type 1 diabetes. And they are treating to an Alc of less than eight with -- in patients up to age 74. And I think that that is dangerous and I think that the evidence is out on that. I don't think it should be incorporated into a composite measure.

administrative burden associated with this sort of composite measure, and that goes against everything we're trying to accomplish in reduced administrative burden on clinicians. And so the measurement process around this measure is extremely complicated. So if it's in the MIPS Program, that's one thing. You don't have to choose it. You wouldn't, because why would you? But if it's in MSSP, you might be stuck with that, and that is unacceptable.

CO-CHAIR KAHN: Carl?

MEMBER SIRIO: One other comment --

CO-CHAIR KAHN: Sure.

MEMBER SIRIO: -- which is I'm not

sure how you want to handle this procedurally,
but the comments I was going to make on 194,
which was the change it to -- change that one to
conditional support. Conceptually they're
linked. So it would seem to me just as a prelude
to the comments I'll make in a few moments,
whatever we do for this one intellectually makes
sense to make -- do it for the second one.

CO-CHAIR KAHN: Okay. Chris?

MEMBER QUERAM: Thank you. I want to speak in favor of this measure and actually am inclined to opposed the motion that's been made, and depending on the outcome of the vote would propose that we change the recommendation to support for rulemaking.

We've been reporting this measure, and for that matter we've been reporting the optimal vascular care measure for seven, almost eight years now. It's been in use. We have not encountered the type of problems that we've heard from the previous speakers in terms of the burden or how cumbersome it is to collect the data. We

adopted it for precisely the reasons that are 1 2 cited elsewhere in various narratives about the complexity of identifying an accountable 3 4 physician. 5 The reason that we adopted this was 6 because it forces a higher standard of care in 7 care coordination and care communication across 8 multiple disciplines that are involved in caring 9 for these patients. 10 I would even go so far as to say I 11 think for purposes of MIPS we should remove the 12 duplicate component measures and put all of our 13 emphasis on the composite. 14 CO-CHAIR KAHN: Okay. The --15 DR. BAGLEY: Chip, my hand's up. 16 CO-CHAIR KAHN: Okay. Is that Bruce? 17 Would you speak? 18 DR. BAGLEY: It is, yes. I would concur with those comments. And I have a couple 19 20 of other things that might help. 21 With -- especially with MSSP there is 22 an individual clinician component or

responsibility, but there's also an organizational component, a responsibility to make sure that there are systems in place that all of these things happen. So I'd be far more in favor of this as an MSSP measure. And we could talk more about whether it should be MIPS as well.

The second thing that I would say is that with any measure that we have ever considered the outcome is a bell-shaped curve and a physician's brain immediately goes to the most -- the sickest patient they've ever seen. Oh, my God, how am I going to get that person under eight? And that's just not the way it works. Biology is not that way.

So we don't know what the optimal outcome for a particular individual clinician might be, but it isn't 100 percent. It's, you know, 90 percent. But unless we measure that, we'll never know what optimal outcome is.

The final comment I'll make is that this clearly is a composite measure which is all

or nothing. And if there are four components to it and you average the four components; in other words, you take individual measures and average them, then you'll get -- like let's say for each of the four components you've got a 75 percent. If you average them, of course the average is 75 percent, but in an all or nothing measure the answer is 0.75 times 0.75 times 0.75 times 0.75, which actually is about 32 percent.

So what happens is in order to do well on a composite all or nothing measure it forces the organization and the clinician and the practice to have a systematic way to get all four of these things done every time. It really is something that drives quality improvement that's not true of an individual measure.

CO-CHAIR KAHN: Okay. Raj?

MEMBER DAVDA: Thanks. I know we speak a lot about MSSP and CMS here, but my job is really to represent the health plans and commercial side of the business as well. And we frequently pick up on what's done here and use

the same measures to reduce the burden for physicians.

So in this one, in addition to 215, I think the -- I agree with the comments that were in support of this. I would agree to support it, again because there are -- there's a larger population also at stake here that has diabetes, and it's not just the CMS population.

So from a health plans perspective Alc less than eight is clearly appropriate for our commercial population in general. And we were forced really to use an Alc greater than nine, and it really doesn't hold true for our population, so I would also support this.

CO-CHAIR KAHN: When you say "support," you mean support the initial one?

MEMBER DAVDA: Yes.

CO-CHAIR KAHN: Okay. Marissa?

MEMBER SCHLAIFER: So I'm not going to comment on the clinical argument speak, but just looking at a little more process. And going back to -- I guess I'm a little uncomfortable with us

questioning -- I'm very uncomfortable with us questioning the appropriateness of the measure that's been NQF endorsed.

And there is an opportunity to click on -- because staff's done such a great job of putting this together, to go back and look at the NQF endorsement review from 2015. A lot of the things being said right now are in the discussion and in the comments that were taken in that the NQF Endorsement Committee took into consideration and then endorsed the measure.

So I think as a MAP our responsibility is to say whether it's appropriate to be used in this program, not to question -- not to revisit the endorsement process. So I just want to make sure that we're kind of staying within our charge of work. And it sounds like we're trying to rethink the endorsement process.

CO-CHAIR KAHN: Any other comments?
Oh, Amy?

MEMBER MULLINS: So thinking about which program it should go into; and I want to

kind of dovetail off of what you said, if you put a measure like these diabetes composites into MIPS and you take out all the other diabetes measures, what's going to happen is no one's going to measure diabetes anymore, because you're not going to pick this measure because it's too hard, it's too many components in one measure. You get credit for one measure, but you're doing four or five at one time. So what you're going to see in effect is no one's going to be measuring diabetes, because why would you?

I mean, you just have to think realistically about what six measures are you going to pick? Are you going to pick one measure that's four different things plus five others? Are you going -- you're just not going to do that. You're an overworked clinician. This is administratively burdensome. And so you're going to quit measuring diabetes. That's not good for anyone.

And this Alc less than eight, this is new and it really hasn't -- the evidence right

now is debatable on this six to nine range versus 1 2 over nine. So to hold physicians accountable for this less than eight, especially in older 3 populations and Type 1 diabetics I think is 4 5 questionable. And then also this -- you have to make 6 7 sure all of your diabetic patients are now non-8 tobacco users. If you visited the states where I 9 used to practice, that is sometimes completely outside of your control. And that's part of this 10 11 composite measure as well. 12 So I mean, I seconded the refine and 13 resubmit. My initial thing was don't support at 14 all, but I would be fine with refine and 15 resubmit. 16 CO-CHAIR KAHN: Okay. Any other 17 comments from the phone? 18 MS. GLIER: Yes, this is Stephanie 19 Glier from PBGH. Can you guys hear me okay? 20 CO-CHAIR KAHN: Sure. Yes. 21 MS. GLIER: Thanks. I want to follow 22 up on a couple of the comments that were just

made. I totally agree that MAP's job is not reendorse a measure, or to re-adjudicate the endorsement of a measure. And I think this is an endorsed measure. The population has been clearly specified. If the guidelines change for Alc levels, then I think that's something that we would want to consider or that CMS would consider, but until that's true I think it's appropriate to move forward with an endorsed measure.

The second piece to Amy's point that she just made, I think this is actually a program design issue, so a comment to CMS. If you're going to continue with a measure that allows clinicians the opportunity to pick and choose the measures that they're most likely to succeed on, I think that's a flaw in the program. I don't think MAP should be keeping good high-value measures out of the program because we're concerned that people will game the program or choose the measures that are easier to report.

Instead, back to what Chris had said

earlier; maybe it was Bruce that said it earlier,

I think MAP's job is to keep the high-value

measures in the program and to remove the lower
value measures so we're pushing people to report

the things that matter more to patients.

MEMBER QASEEM: Okay. So just a couple of things. We're not going to go dive into the evidence in this one, but I mean, at any point in time I can give you a quick summary of this.

The newer evidence is actually pointing towards individualized decision making. If you look at any of the new guideline recommendations that have come out, they do say that you target, talk to the patient and it's a joint decision making. No one is saying anywhere seven to eight at this point because of the newer evidence. And if you look at even the VA guidelines that just came out last year, right, they also said individualized decision making, just one thing.

The second thing is you can already

1	see there's enough we're not agreeing on a
2	measure. Generally speaking, and again looking
3	at you, Kate, I mean, if there is there's
4	plenty of low-hanging fruit. First we need to
5	improve quality of care versus start targeting
6	where we are not agreeing on something. I mean,
7	that's to begin with should not be used for
8	accountability as a fundamental principle.
9	That's just a personal opinion.
10	CO-CHAIR KAHN: Okay.
11	MEMBER QASEEM: There is a motion.
12	CO-CHAIR KAHN: Oh, wait. No, is
13	there any other comments?
14	(No audible response.)
15	CO-CHAIR KAHN: Okay. Let's have a
16	vote on this.
17	PARTICIPANT: Which one.
18	CO-CHAIR KAHN: I know.
19	PARTICIPANT: There's only one.
20	CO-CHAIR KAHN: On this vote, just
21	remind people, we have two separate measures I
22	mean, two separate measures or well, one

1 measure that is used in two separate programs. 2 And if you allow me, I will package this so that -- because I think if we're going to -- if you're 3 4 going to vote one way, you're going to vote one 5 way on both. So if it's agreeable, I'm going to 6 7 package this into one motion, but the motion 8 would affect both programs. And the issue is 9 moving this diabetic -- diabetes measure from conditional to refine and resubmit. 10 That's what 11 the motion is. I'm -- Carl? 12 MEMBER SIRIO: No, I was going to let 13 you finish. Go ahead. 14 CO-CHAIR KAHN: And that's what we would be voting on, if it's acceptable to the 15 16 group, because I'm repacking it a bit just for 17 simplicity. 18 Yes? 19 PARTICIPANT: For process we need 20 individual votes, so --21 CO-CHAIR KAHN: Okay. 22 PARTICIPANT: -- we have to do it

twice.

CO-CHAIR KAHN: Then we will vote twice.

MEMBER SIRIO: Actually, look, I don't want to get real crazy, but this is getting a little much. We don't need two votes. If in fact the Committee wants to split it -- but if we want to split it, basically what you're saying is the motion includes both. Someone around this table could say split the thing, otherwise you get one vote. I mean, we're spending 20 minutes worrying about a clicker.

Let me get to my point. So the point that Chris made earlier, which is what happens if this motion fails.

Chris, the motion as I understood it would revert to where we are, which is conditional support for rulemaking, which would -- I don't think you need a motion for. And I would just like to make sure that the comments that were made with respect to some of the stratification issues go forward. That's all.

CO-CHAIR KAHN: I was going to -- what I was going to do -- well, let me -- I'm going to make an alternative suggestion then. Let's vote -- one, let's have the vote and then I think we can look around the table and revert back to the comments going over if the vote fails.

Let me -- MIPS -- why don't we do MIPS first. Where is Amir?

MEMBER SIRIO: Why don't we do this by hand then? We're wasting way too much time.

CO-CHAIR KAHN: No, no, I understand, but let me offer -- let me make clear people need to know what they're voting on. We'll vote on MIPS, and frankly my argument would be if the vote is clear one way or the other, then I'm happy to have a vote on the second item, but I don't -- but it's only going to be clear if the motion carries that it's worth having a second vote.

So we'll have one vote and then we'll see whether Amir who offered the motion wants to have a second vote, because he's got to win the

first one to have the second one.

Chris?

MEMBER QUERAM: Chip, just to be clear, then, Carl, on your motion you're not recommending -- or the motion doesn't propose changing the workgroup's recommendation, but adding the requirement that the measure be risk-adjusted or stratified?

MEMBER SIRIO: I'm just suggesting
that -- I'm asking that the commentary go along.
I'm not even going so far as to propose that the
measure change. So I think what it would revert
to is the -- is Bruce Bagley's committee's
report, which was conditional support with a memo
going over to CMS with some additional
commentary.

CO-CHAIR KAHN: Yes, actually -- well, let's get through the motion and then we'll --

MEMBER SCHLAIFER: I was just going to mention that as was brought up earlier, it might not influence the vote, but a reason for two votes is MIPS is -- it's -- you get to use it as

1	a voluntary measure, where for MSSP it's a
2	mandatory measure. So there is there could be
3	a difference in voting. I don't think there
4	would be, but I mean as far as just wanted
5	CO-CHAIR KAHN: Okay. MIPS. Are we
6	ready to use this thing?
7	MS. OGUNGBEMI: Yes. No, we're voting
8	with clicker.
9	CO-CHAIR KAHN: I'm willing to use it
10	if we do it fast.
11	MS. OGUNGBEMI: So we're voting for
12	the recommendation for MIPS, optimal diabetes
13	care, MUC17-181, refine and resubmit. This is
14	for MIPS. Yes or no voting is open.
15	CO-CHAIR KAHN: Okay.
16	(Voting.)
17	MS. BUCHANAN: And I have Mira's.
18	Steve, I don't know if you are able to vote, but
19	if you are, please email. Stephanie, Bill is
20	back, so he is voting. And so but I just have
21	Mira's right now.
22	Okay. Maureen. I have vours.

Okay.
CO-CHAIR KAHN: Okay. And the vote is
overwhelming no by a consensus amount, so that
the motion fails.
So, Amir, the question do you want to
bring it up for a vote on the other program?
(No audible response.)
CO-CHAIR KAHN: Okay. Now, but to be
clear in terms of Carl's comments, is let me
just ask. We're not going to vote, but is it
sort of generally around the table a feeling that
it's fine that as one of the conditions to ask
that CMS consider some stratification? I don't
think you defined exactly what it was.
We're not voting on the second one
because we're going to revert back to the consent
calendar. Yes.
MS. OGUNGBEMI: And for the record we
have 32 percent yes and 68 percent no.
CO-CHAIR KAHN: Right. Okay. We're

And the vote's the vote. So now let's go

going to go forward with the comments that Carl

made.

21

1	to the next item.
2	MS. O'ROURKE: Procedurally John
3	pointed out that now that we took a motion, that
4	is off of the consent calendar. So we would need
5	to now take a vote for the conditional support.
6	CO-CHAIR KAHN: Well, wait a minute.
7	Wait a minute. It reverts back to the consent
8	calendar. Why do we need to have another vote?
9	MS. O'ROURKE: This came up in
10	clinician; and I would ask Bruce to correct me if
11	I'm wrong, but per parliamentary procedure once
12	you take it off a consent calendar, it can't go
13	back on. Is that correct, or can
14	DR. BAGLEY: That's correct. You have
15	to vote on it.
16	PARTICIPANT: So moved.
17	CO-CHAIR KAHN: Okay. Let's vote.
18	We're voting on we're now voting on whether or
19	not we make a conditional recommendation
20	including Carl's comments, on the measure on MIPS
21	regarding diabetes.
22	So one is wes and two is no.

Actually, can we have a show of hands 1 2 in terms of it being conditional? (Show of hands.) 3 CO-CHAIR KAHN: This is the MIPS. 4 5 have to vote on MIPS again because we -- since it is now off the consent calendar because we had a 6 7 motion. So it's clear that we have consent. 8 MS. BUCHANAN: And so I have two 9 people on the phone who said yes. Four, five, 10 six, seven, eight, nine, ten, eleven, twelve, 11 thirteen, fourteen, fifteen, sixteen, seventeen, 12 eighteen, nineteen, twenty, twenty-one, twenty-13 two, twenty-three. Twenty-four, because Maureen 14 just said yes. So we have 24 for yes. 15 CO-CHAIR KAHN: Okay. And I assume we 16 have one no? Right. 17 Okay. Now -- and we didn't -- and 18 since on the other we didn't vote on the motion, it remains on the consent calendar, on the second 19 20 You made a motion but we -- it was 21 withdrawn, so --22 MS. O'ROURKE: Okay.

1	CO-CHAIR KAHN: it remains on the
2	consent calendar.
3	CO-CHAIR KAHN: So to clarify, optimal
4	diabetes care remains on the consent calendar for
5	MSSP.
6	CO-CHAIR KAHN: Right, because there
7	was no motion voted on by the body.
8	Okay. Let's move to the next one,
9	please.
10	MR. BERNOT: Sure. Hurry, hurry,
11	hurry.
12	So we will go to the next one, which
13	is MUC17-194. And this particular measure was
14	pulled for both or sorry, for the MIPS
15	Program. The preliminary recommendation from the
16	workgroup was support for rulemaking, and the
17	preliminary assessment from the staff was
18	conditional support with the condition that the
19	that there's no duplicate measures.
20	At the meeting CMS had said that they
21	would just be swapping this measure with a
22	preexisting measure. The motion was subsequently

changed to support, so the workgroup 1 2 recommendation is support for rulemaking. This particular measure was pulled again by Carl. 3 4 CO-CHAIR KAHN: Carl? No, this is what we're supposed to be doing. 5 6 MEMBER SIRIO: No, no, no. I mean, in 7 terms of the voting and all that good stuff. 8 CO-CHAIR KAHN: No, no. Move forward. 9 MEMBER SIRIO: So on this one actually 10 I'm encouraged on the one hand to hear that the staff recommendation is the same thing that I'm 11 12 actually asking us to revert to. It's interesting; and I don't know if 13 14 it's worth a conversation, this will be the second time in 20 minutes where in fact the staff 15 16 recommendation and the Committee recommendations 17 were different in terms of their conversation and 18 we're reverting back to the staff. So I think 19 there's some signal in there that we should talk 20 about maybe tomorrow. 21 But that having been said, my comment If the one 22 really is to align the two measures.

that we just discussed is conditional and these are in essence linked in terms of the care of the diabetic patient, that this should be conditional also. So all I'm now suggesting is the motion would be to revert to the staff recommendation, which is to support conditional.

CO-CHAIR KAHN: Okay, So themotion is

CO-CHAIR KAHN: Okay. So themotion is to make this conditional. Do I have a second?

(No audible response.)

CO-CHAIR KAHN: I need a second.

MEMBER QASEEM: Second.

CO-CHAIR KAHN: Okay. Amir. Well, it's for the person offering the motion to explain what the condition is.

MEMBER SIRIO: Sure. So there's a couple of points: The first is that there -- it's consistency in terms of the care of the diabetic. And secondly, there is some -- there are some blood pressure guidelines that need to be accounted for. So one would at least need to look at the newer guidelines to make sure that the measure is consistent with current science.

know if the measure developer is on the line, but they are  (Simultaneous speaking.)  MEMBER SIRIO: To assess whether the measure is consistent with best practice, and if it's not, to align with best practice. So it would be to tweak it in a way that actually would be making the measure current.  CO-CHAIR KAHN: I'm sorry. Is this a endorsed measure?  MR. BERNOT: Yes, it is.	
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endorsed measure?	
	n
MR. BERNOT: Yes, it is.	
MS. MOYER: And this is Amy. It's	
currently undergoing revision to address	
compliance with that guideline.	
CO-CHAIR KAHN: Oh, okay.	
MEMBER QASEEM: So but then it	
shouldn't be conditional. So that measure is	
going through revision. It's not even ready yet,	
so how can we shouldn't it be refine and	
resubmit and you look at the new measure?	
CO-CHAIR KAHN: Well, Marissa	
CO CIMILIA ACITA MOLLA M	

MEMBER SIRIO: Well, Amir, I'm going to suggest for political purposes that is likely to fail again in terms of where we're going.

CO-CHAIR KAHN: Well, Marissa, what were you going to --

(Simultaneous speaking.)

MEMBER SIRIO: So my only point is that if it's already endorsed and it's a tweak on one part of it, it probably doesn't need to be tested all over again.

MEMBER SCHLAIFER: I'm fine with the recommendation, but I think I disagree with the fact of saying that it's not really out there because it's being -- I mean, updated. It's still -- there is a measure out there that is NQF-endorsed until the revision comes forward. So I'm not opposed to the discussion. I just want to make sure that we get that right.

MS. MUNTHALI: I just wanted to clarify. It's going through a maintenance review, so it isn't currently endorsed, but it is being reassessed after -- per our process after

_	three years, yes.
2	CO-CHAIR KAHN: Well, let me ask a
3	but technically well, it is endorsed, but
4	there is a likelihood that it would be revised in
5	this technical I mean, in this technical
6	review or
7	MS. MUNTHALI: I think this is
8	included in the revisions, so this would be a
9	significant revision to the measure. But is
LO	under review right now. It is currently an
L1	endorsed measure. It's going through its
L2	maintenance process.
L3	CO-CHAIR KAHN: So when something's
L <b>4</b>	going through a maintenance process, it will come
L5	back for re-endorsement?
L6	MS. MUNTHALI: If it's re-endorsed, it
L7	will be endorsed again for three years.
L8	CO-CHAIR KAHN: Well, it is different
L9	than the other measure that was just endorsed.
20	Marissa?
21	MEMBER SCHLAIFER: This is just a
22	process question. When you when a measure

gets revised through the normal -- and goes 1 2 through the normal NQF process, does CMS automatically revise that, or how does that work? 3 4 MEMBER GOODRICH: So we always want to 5 use the most updated form of the measure. that relates to the MAP though, the answer is it 6 7 depends as to whether it comes back here. 8 comes back here to the MAP if the revision is 9 determined to be a significant revision, and there are parameters around what "significant" 10 11 If it's a minor insignificant revision, means. 12 like a few codes are added for new drugs that 13 came on the market or something like that, we 14 don't bring it back. 15 You would update MEMBER SCHLAIFER: 16 it? 17 MEMBER GOODRICH: We would update it, 18 We always in our regular ongoing process of 19 updating measures would use the updated measure, 20 yes. 21 CO-CHAIR KAHN: But it does say that 22 doing conditional would be recognized.

1 going to recognize it in a sense by what they're 2 going to do. I guess does anyone -- Stuart's on the 3 Does he or she -- who is it? 4 phone. 5 PARTICIPANT: Colleen. Colleen, 6 CO-CHAIR KAHN: do you have 7 any comment? 8 MS. PITZEN: Hi, this is Collette 9 Pitzen --Oh, Collette. 10 CO-CHAIR KAHN: 11 sorry. 12 MS. PITZEN: -- from the Minnesota 13 Community Measurement. As these measures were submitted for the call for measures in June the 14 15 new guidelines had not yet come out. They were 16 fairly recent the end of 2017. And we are taking 17 action to review those to connect with the Measure Development Workgroup and determine what 18 19 we are going to do with the guidelines and some of the controversy around them. And we will have 20 an answer before this comes back to MAP review 21 22 again.

1	CO-CHAIR KAHN: Okay. That was oh,
2	Chris?
3	MEMBER QUERAM: Thank you, Chip. my
4	comments on this measure are similar to the ones
5	that I made on the optimal diabetes care. We've
6	been reporting this as well. It's an accepted
7	measure. It has been well embraced by the
8	clinical community in our state. I come from
9	Wisconsin. Run a quality collaborative in our
10	state. I know the same is true in Minnesota.
11	I oppose the motion that would change
12	this to conditional support and prefer that we
13	maintain the recommendation from the workgroup
14	that it be supported for rulemaking.
15	CO-CHAIR KAHN: Okay. Carl, anything
16	else?
17	MEMBER SIRIO: No.
18	CO-CHAIR KAHN: Okay. Any other
19	comments? I'm sorry. Leah?
20	MEMBER BINDER: Yes, I would support
21	Chris' recommendation. I just also want to make
22	a process point that I don't think we have any of

us months to sit on this committee and relitigate all of the endorsement protocols that have already happened. I'm just concerned that we won't even get through this agenda if we keep re-litigating the endorsement process, which I think we all know. So we put so much time into that process. It's a thorough process and I think we should have some confidence in it.

CO-CHAIR KAHN: Okay. I think we've heard the points. Anybody else? Any other -- oh, I'm sorry.

MEMBER KRAMER: I also think that we ought to be careful about how we -- what kind of conditions we put on. If something is lacking NQF endorsement and needs to go through the endorsement process, that seems to be an appropriate condition. I think a simple updating for updated clinical guidelines is more like a maintenance kind of function which would be done automatically. This is already an endorsed measure. I don't think it warrants applying a condition. So I would not support -- I recommend

1	that we not support this motion.
2	CO-CHAIR KAHN: Okay. So let's then
3	on this measure I guess the committee approved
4	it, or recommended that it be sent forward with
5	approval. And the motion is to change that to
6	conditional. And so a yes would be for
7	conditional; a no would be for the Task Force's
8	recommendation that it be recommended to CMS.
9	So with that, is everybody ready? You
10	ready?
11	MS. OGUNGBEMI: Yes.
12	CO-CHAIR KAHN: Okay. Let's one is
13	yes and two is no.
14	MS. OGUNGBEMI: Yes, we're voting on
15	a recommendation for MIPS, optimal vascular care,
16	MUC17-194, conditional support. One is yes; two
17	is no. Voting is open.
18	(Voting.)
19	MS. BUCHANAN: I'm still waiting to
20	hear from people participating through the phone.
21	MEMBER SIRIO: general comment?
22	CO-CHAIR KAHN: Yes.

MEMBER SIRIO: It does strike me as worthy of a discussion at some point that we've had now two episodes of stuff that gets pulled where we've had discord between the staff and the committee.

And for this group to understand that, in terms of being the final adjudicator of the recommendations that go forward, we need some mechanism I think to understand when that disconnect or that discord happens between the staff assessment and the workgroup, you know, what, a little bit more background in terms of the whys and the wherefores.

It's in here, but I think an explicit discussion might be warranted insofar as it would maybe inform this conversation a bit better.

CO-CHAIR KAHN: Well, I think, so if we, if this happens, we're going to discuss more of them, then, one, we know from the discussion this morning that the staff follows this algorithm to come to its conclusion.

So, if you could, when you describe,

1 explain why you think the task force made a 2 different recommendation than was in your draft based on the algorithm, they can explain it. 3 4 Well, you explain it. 5 Okay. So do we have the vote? 6 We have 24. So, if that's sufficient, at 25? let's go ahead and close it. 7 8 I apologize. MS. BUCHANAN: I think 9 we'll have to scrap the voting as far as again. So I think moving forward we won't be doing that 10 11 anymore, so many apologies. I think we're going to have to handle it. It did not capture. 12 13 CO-CHAIR KAHN: Okay. All of those in 14 favor of the motion to move this measure from recommend to conditional recommendation, please 15 16 vote aye and raise your hand if you're for it, 17 the motion. 18 (Show of hands.) 19 CO-CHAIR KAHN: Okay. And now, 20 everyone who is opposed to the motion, please 21 raise their hands. 22 (Show of hands.)

1	CO-CHAIR KAHN: Okay. So the motion
2	fails, and we voted. So let's go to the next
3	measure, please.
4	MR. BERNOT: Sure thing. So the next
5	measure is MUC17-215. This is the diabetes A1C
6	control less than 8.0. It has been pulled for
7	both MIPS and MSSP. It was identified for MIPS
8	by Bill Kramer. I will let him go over his
9	rationale. And it was pulled by Carl for both
LO	MIPS and MSSP.
L1	CO-CHAIR KAHN: Okay. I'm sorry. So
L2	we have two people pulling it for different
L3	reasons presumably. Okay. So who pulled it
L <b>4</b>	first?
L5	(Off-microphone comments.)
L6	CO-CHAIR KAHN: No, no, no, seriously,
L7	because we have to, I mean, we got to stack this
L8	linearly, or I'll just choose.
L9	MR. BERNOT: Bill's was first on the
20	list.
21	CO-CHAIR KAHN: Who?
22	MR. BERNOT: Bill Kramer's was first

on the list.

CO-CHAIR KAHN: Okay. So Bill is first. And I hope we can maybe even get a motion, if there is one from Bill, unless it's just a comment. So we can then get going through it.

MEMBER KRAMER: Yes, okay. The recommendation applies from -- the reason I asked for this to be pulled and reconsidered and would ask for a revote is with regard to the use of this measure in MIPS, not MSSP.

CO-CHAIR KAHN: Okay.

MEMBER KRAMER: So it's just one piece of this. The rationale is that this is a voluntary -- two parts, it's a part of a larger composite measure. It's one component of a composite measure.

Second, it exists in a program, MIPS, being proposed here for a program, MIPS, which there is voluntary reporting. In other words, clinicians can choose to report on this or something else.

So this, while the measure itself seems like a fine measure, the danger is that in the use of this measure in MIPS, it might be chosen by clinicians instead of the composite measure, in which case we, the stakeholders in healthcare, consumers, purchasers, and clinicians, would lose information about what is happening with the optimal care.

So, because of the risk of how it would be used in MIPS, displacing a more valuable composite measure, I recommend that it not be approved for use in MIPS so that the information that will be gathered will be on the composite measure.

CO-CHAIR KAHN: Okay. So, I'm sorry, so your recommendation is that it be removed from MIPS. Is that --

MEMBER KRAMER: That it not be, that we would recommend, or that the -- I would move that MAP recommend that it not -- to CMS -- that it not be used, this measure not be used in MIPS.

CO-CHAIR KAHN: I'm sorry, in --

That this measure 1 MEMBER KRAMER: 2 would not be used in MIPS. MIPS, okay. 3 CO-CHAIR KAHN: So it 4 would be removal. He's recommending to remove 5 So you've made a motion. Can the staff sort of describe anything else about this, 6 7 who our expert is? 8 MR. BERNOT: Sure, sure. So the 9 discussions around this -- and, Bruce and Amy, feel free to jump in if I missed anything. 10 11 they said that, even though the composite was, 12 that this was discussed with the workgroup, that 13 there was a composite and a subcomponent of the 14 composite. And they did talk about the tension 15 16 between, that there was some folks who felt there 17 was still value in this particular measure for 18 several reasons. It was part of some programs, part of things that people were tracking. 19 Also the attribution of the composite 20 21 was a concern to the workgroup that they may not

be, fully be able to affect all of the

1	components, but this would be a way for them to
2	track diabetes. That's my very quick summary
3	unless, Bruce or Amy, you have other
4	recollection.
5	DR. BAGLEY: No other comments.
6	CO-CHAIR KAHN: Okay. Are there
7	comments around the table? Thoughts? Chris?
8	MEMBER QUERAM: Do you need a second
9	for the motion?
10	CO-CHAIR KAHN: Oh, yes, actually.
11	Thanks.
12	MEMBER QUERAM: Second.
13	CO-CHAIR KAHN: Yes. Thank you. I
14	apologize. Any other comments?
15	MEMBER SIRIO: So my only comment
16	would be I don't think I would get as draconian
17	as withdrawing this. I think the ask that I was
18	going to make for both of these was the same as
19	earlier. It's just that we look at the issue as
20	a recommendation to Kate and her team with
21	respect to appropriate risk stratification. It's
22	not even risk adjustment; it's risk

stratification. 1 2 So I won't support it, not because I don't think that the argument's got merit. 3 But I think that there's value in the communication. 4 CO-CHAIR KAHN: Yes, I think --5 MS. PITZEN: This is Collette from 6 7 Minnesota. May I make an additional comment --8 CO-CHAIR KAHN: Oh, sure, yes, please. 9 MS. PITZEN: Yes, we had proposed 10 putting forward this component. Believe me, we 11 are big believers in the composite. The purpose of this was to replace the poor control A1C 12 greater than 9 measure that's currently existing 13 14 in programs, because when you're looking at poor control, there's the unintended consequences that 15 16 anything less than 9 is good control. And that's 17 simply not true and not evidence-based for 18 diabetes care. Thank you. 19 CO-CHAIR KAHN: Continue other 20 discussion? Bill, do you have anything else to 21 say?

MEMBER KRAMER: I'll say this.

not opposed to the measure, per se. Just to clarify, this isn't a complicated argument I'm making, that the problem is how the measure is used in MIPS. So I fully support it being used in MSSP, where it is a required measure. The problem is how it's used in MIPS has the -- would have the unintended effect of crowding out a more valuable measure, the diabetes composite measure.

CO-CHAIR KAHN: It's really a structural argument. Chris?

MEMBER QUERAM: And I just build on Bill's comments a little bit and some of my earlier observations. What animates my thinking about this motion is that we need to begin to take steps, however measured or however big they may be, to up the bar a bit in terms of what it is we expect of clinicians in these programs.

A gentleman made the comment earlier that there's lots of low hanging fruit out there, and we should be focused on that. Many of us have been around these tables for a lot of years, and we keep talking about low hanging fruit.

When are we going to move to higher hanging fruit? And I think this is a good example of taking a measured step in that direction.

CO-CHAIR KAHN: Good. Okay. Any other comments?

MEMBER O'BRIEN: I think the way that CMS would describe the whole point of the choice within MIPS is that, so you can, a clinician can find the measures that are the best fit for their practice.

When talking about a component of a composite measure, I think unless there's a pretty clearly articulated and valid reason why the composite measure itself isn't a -- wouldn't generally be a good fit, I think there is a -- to me, there only seems to be downside to including a component measure. And the downside is the risk of someone choosing the component measure, not so much because it's the best fit for their practice, but it's going to -- they know they're going to do well on it. It's going to put their practice in the best light.

So I don't see any upside to including 1 2 this, only downside, when it comes to MIPS itself. 3 4 CO-CHAIR KAHN: Okay. Enough 5 Let's just -- we're going to vote by Yes, please. Okay. The motion --6 hands? 7 MS. MUNTHALI: Sorry, we have one more 8 voter who's joined us. Rich Antonelli's on the 9 phone. Rich, can you introduce yourself? let us know if you have anything to disclose. 10 11 MEMBER ANTONELLI: Dr. Richard Antonelli, Medical Director of Integrated Care, 12 13 Boston Children's Hospital. And I'm representing 14 the hospital. And I don't have any disclosures. CO-CHAIR KAHN: Rich, I'm happy to 15 have you. Okay. So the motion is to remove this 16 17 measure. And we've had discussion. 18 (Off-microphone comments.) 19 CO-CHAIR KAHN: Oh, I'm sorry. Ι 20 thought remove and not support was the same 21 thing. Or not support this measure or recommend 22 not to support. So a yes vote is supporting the

motion not to support. A no vote is to maintain 1 2 the current recommendation. 3 MS. BUCHANAN: Not to support for 4 MIPS. 5 CO-CHAIR KAHN: Not to support for MIPS, I'm sorry. Yes, it's for MIPS. 6 It doesn't affect the other; only voting on MIPS. 7 Okay. 8 All those in favor of the motion please raise 9 their hands, and on the phone, you can vote yes or no now by email. Okay, here. 10 11 (Show of hands.) 12 MS. BUCHANAN: One, two, three, four, 13 five, six, seven, eight, nine, and then I have, 14 on the phone, I have Maureen, Steve, and Mira's I don't -- Rich, I don't know if you're 15 16 connected to the, your email right now. 17 you're not, if you just wouldn't mind verbally 18 saying yes or no. 19 MEMBER ANTONELLI: Actually, I just 20 clicked send. Could you just confirm that you 21 receive it in the next five seconds? MS. BUCHANAN: 22 So --

1	MEMBER ANTONELLI: The vote is in the
2	subject line.
3	MS. BUCHANAN: So I didn't did you
4	send it to mapcoordinatingcommittee@qualityforum
5	.org?
6	MEMBER ANTONELLI: Yes.
7	MS. BUCHANAN: I did not receive it.
8	MEMBER ANTONELLI: I'll send it again.
9	MS. BUCHANAN: Okay. Rich, I still
10	haven't received anything. So maybe just
11	verbally
12	MEMBER ANTONELLI: Okay.
13	MS. BUCHANAN: Oh, I got it. Thank
14	you.
15	MEMBER ANTONELLI: Yes.
16	CO-CHAIR KAHN: Okay.
17	MS. BUCHANAN: And then yes, you need
18	yeses.
19	CO-CHAIR KAHN: And then we vote no.
20	MS. BUCHANAN: So we voted no. We
21	have 11 noes. And now for yeses.
22	CO-CHAIR KAHN: How many?

1	MS. BUCHANAN: 11 noes. So for yeses
2	
3	CO-CHAIR KAHN: But we didn't vote no.
4	We didn't do the votes for no.
5	MS. BUCHANAN: Sorry, we have 11 for
6	yes. We have not done the noes.
7	CO-CHAIR KAHN: That's what I thought.
8	Okay. So can we have the votes for noes?
9	MS. BUCHANAN: One, two, three, four,
10	five, six, seven, eight, nine, nine, and then we
11	have three. So that's 12. Okay.
12	CO-CHAIR KAHN: So what do you have
13	a
14	MS. BUCHANAN: I do. I do not have 25
15	people, though.
16	CO-CHAIR KAHN: Yes, Mira's not
17	actually, do you have enough to have to meet the
18	quorum test?
19	MS. BUCHANAN: We do have enough for
20	quorum.
21	CO-CHAIR KAHN: Then, if you're not at
22	the table, you don't get a vote. If we have a

1	quorum, that's it.
2	MS. BUCHANAN: Okay. So we had 10 for
3	yes, 12 for no, which gives us
4	(Off-microphone comments.)
5	MS. BUCHANAN: No, we have 10. I was
6	wrong the first time.
7	CO-CHAIR KAHN: We have to get to 60-
8	plus
9	MS. BUCHANAN: And so we're at 45
10	percent. So the motion does not pass.
11	CO-CHAIR KAHN: Okay. So the
12	recommendation goes forward as made by the task
13	force. Let's go to the next one.
14	Oh, do we have to vote again?
15	MS. BUCHANAN: Yes.
16	MR. BERNOT: For MIPS.
17	MS. BUCHANAN: For MIPS. So this is
18	a vote for MIPS to conditional support.
19	CO-CHAIR KAHN: But what if it doesn't
20	get the 60-plus? Okay. So everybody vote now.
21	MS. BUCHANAN: So, to clarify for the
22	record, this is the diabetes A1C control measure.

1	And you are voting yes to conditional support.
2	No is a disagree with conditional support.
3	CO-CHAIR KAHN: Any noes? I think we
4	made it.
5	MS. BUCHANAN: We did. I just want to
6	make sure, because I just wanted to capture our
7	online people who are voting in as well.
8	CO-CHAIR KAHN: While you're capturing
9	those
LO	MR. BERNOT: Sure thing. Next one is
L1	MUC17-234. This is the ischemic vascular disease
L2	use of aspirin or anti-platelet medication. The
L3	workgroup came up with a recommendation of
L <b>4</b>	conditional support for rulemaking. And the
L5	conditional was that there's no competing
L6	measures within the program or that is rectified.
L <b>7</b>	That's for MIPS.
L8	They also had a similar recommendation
L9	for MSSP conditional support, with the condition
20	that there are no competing measures in the
21	program, exact same thing.
22	So this was originally pulled by Bill

Kramer for discussion for MIPS only I believe, 1 2 Bill. And then, Chip, you had pulled it for MSSP to have a discussion. 3 4 MEMBER KRAMER: Which measure are we Sorry, I thought we were still on the A1C. 5 on? 6 MR. BERNOT: 234. MEMBER KRAMER: 7 Okay. I thought --8 didn't Carl also pull -- okay, okay. So we're 9 not reconsidering the A1C for MSSP and MIPS. 10 understand. Okay. Sorry. I was --11 CO-CHAIR KAHN: Can you repeat --12 Absolutely, yes. MR. BERNOT: measure is MUC17-234, ischemic vascular disease 13 14 use of aspirin or anti-platelet medication. And this was pulled for MIPS by Bill Kramer. 15 16 MEMBER KRAMER: The rationale for this 17 recommendation is similar to the one we just went 18 through, which is that the aspirin use is a 19 component of the optimal vascular care composite. 20 Again, if it is used in the MIPS 21 program, it has the risk that it will crowd out a 22 more valuable composite measure. So it's

basically the same rationale as on the previous 1 2 measure. So my recommendation is that we do not 3 support for use in MIPS. 4 MR. BERNOT: Correct. This was a 5 conditional support that there was no -- that it was rectified with competing measures within the 6 7 program. 8 CO-CHAIR KAHN: That was, that's what 9 the staff --That was both staff as 10 MR. BERNOT: 11 well as the workgroup. 12 MEMBER MACKAY: I support Bill's recommendation to not include it in the MIPS 13 14 I'm seconding his motion. program. MEMBER ROBINSON: Yes, I just wanted 15 16 to offer a comment. I think if, I think we have 17 to be careful about the assumption that if we 18 take out the component measures that are in the 19 composite measures that it will sufficiently 20 shrink the pool of other measures such that

composite measure. I don't know that the math

providers have no choice but to pick the

21

necessarily equates with that.

I mean, like in the end you could end up with no individual component measures selected because we pull them out. The composite measure isn't selected because the provider chooses not to select that one. And they select a universe of other measures.

I think if the rationale is that there are ten measures and we have to choose six and are going to pull out three or four, and so that means you by default have to choose the composite, then I think, you know, it maybe seems more rational.

MEMBER KRAMER: You're right there.

That might be the case in some cases. That might be the case. And we're talking about the, you know, problem we all know exists, the structure of MIPS on the choice of measures.

I think the net effect will be, if this component measure stays in, there will be less reporting of the composite measure. And if we take it out, there will be more reporting of the composite measure.

But you're right. It won't be -there will be some cases in which they choose not
to report the composite measure. But I think the
net effect will be, there will be -- we'll get
better information for patients, for consumers,
purchasers, and clinicians, and taxpayers about
how the, like the quality of care is just being
delivered.

MEMBER ROBINSON: This is a quick follow up. Are there any other examples where this has happened that, you know, can serve as a model for us? I mean, it's sort of an assumption versus having seen this happen in two or three other cases, and it's substantial enough to inform our decision.

MEMBER KRAMER: I'm not aware of any evidence relying on kind of our, it's, how human nature would work. But the -- I will say, too, as just an additional point, we've been talking about trying to remove the clutter in our measurement system. There are too many measures.

This kind of goes along the point that Chris was making earlier that we should be moving toward better measurements and removing measures that are of lower value. This is a lower value measure because it's a component. We have a better measure, the composite. Let's support that. Let's get rid of the clutter, the low value measures.

CO-CHAIR KAHN: And there is some discussion. I mean, I've heard discussion about that, I mean, that if you don't like MIPS, you can play the minimums. So I don't, I think it is possible. But, Leah?

MEMBER BINDER: Actually, I agree with everything Bill said.

CO-CHAIR KAHN: Shaun?

MEMBER O'BRIEN: Yes, I just wanted to amplify Bill's response to Derek. One of the things we do know is that consultants advise clinicians, you know, to, I'll use the pejorative term, game the system, but, you know, to select those measures that they already do, they know

they already do well on. And that's the risk here when you have a component of a composite, and they're both in the same set.

And so, you know, look, in my mind, having, you know, a component and a composite undermines the credibility of the system. And, yes, there's a structural problem here, but I just don't think CMS should be feeding that structure. I think that's what this does.

MEMBER MULLINS: Raise your hand if you've ever had to actually report a measure.

Okay. It is hard to do. And by doing composite measures, you're not reducing the number of measures. You're packaging measures together.

But the same number of measures still exist.

You can't just start removing component measures and assume people are going to report composites. As an overworked clinician, it's not going to happen. And until the systems we have support them and make it easier to report measures, people are not going to report composite measures. They just don't have the

time and the energy and the resources to do it.

As a rural clinician with one, you know, support staff in your office that's typically maybe your MA or your, you know, spouse that's running your office to report all these measures, I mean, just isn't -- it's not feasible to have a set of MIPS measures that are a bunch of composite measures that now six measures are now 24.

You just can't take away all the component measures. You just -- the system is not set up to support that at this time. Maybe at some point; not right now.

MEMBER DAVDA: One other point I would bring up to keeping individual measures is really the data collection issue. A lot of the composite components tend to be hybrid data. And there is truly a data collection and burden issue that at least our health plan and many of the health plans face because of the lack of interoperability. So, and it is a very manual process.

So I don't know where the motion is, 1 2 but I would support keeping the individual 3 measures. 4 MEMBER ROBINSON: I'd echo Raj's 5 And the other question is whether this comments. is the right mechanism for addressing the 6 7 opportunity to better employ the use of composite 8 measures. 9 I mean, should another tactic be looking at increased weighting of composite 10 11 measures in the program versus the individual 12 measures as a mechanism for incentivizing 13 providers to select the composite measure? Just 14 trying to think about, you know, different ways 15 we can use this forum versus other forums to get 16 to that end goal. 17 MS. BUCHANAN: Microphone. 18 CO-CHAIR KAHN: What? 19 MS. BUCHANAN: Microphone, if you wouldn't mind. 20 21 CO-CHAIR KAHN: -- kind of vote as the 22 previous one, which is to recommend against a

measure, if I've worded it properly, going
forward. And so let's we're going to vote by
hand.
So those voting in favor of the motion
would be to recommend that this measure be
removed, this measure not be recommended. Okay.
So all those voting in favor of the motion,
please raise their hands.
(Show of hands.)
CO-CHAIR KAHN: And on the phone,
please vote by email. Those voting against the
motion here in the NQF offices, please raise your
hand.
(Show of hands.)
CO-CHAIR KAHN: Okay. I can just tell
we didn't get to 60-plus percent. So did we make
it?
MS. BUCHANAN: We did not. So we had
10 for yes, do not recommend, and we had 12 for
no, to oppose the measure.
CO-CHAIR KAHN: And we made 22 votes.
MS. BUCHANAN: We made 22 so far.

1	CO-CHAIR KAHN: So we had enough
2	votes. Okay. Let's go to the next one, please.
3	MR. BERNOT: So we just have to go
4	I'm sorry. But we just go back and
5	CO-CHAIR KAHN: Oh, I'm sorry.
6	MR. BERNOT: the staff conditional
7	support
8	CO-CHAIR KAHN: So now we have to vote
9	in support of, in a motion. Do we have to offer
LO	a motion for that, or can the Chair just
L1	MS. BUCHANAN: If there's no other
L2	motions, you can go forward. Then we vote on the
L3	staff.
L <b>4</b>	CO-CHAIR KAHN: Okay. So, now we're
L5	going to re-vote on the recommendation of the
L6	task force actually. So all those in favor of
L7	the recommendation of the task force, please
L8	raise their hands.
L9	(Show of hands.)
20	CO-CHAIR KAHN: And on the phone,
21	please vote by email. All those opposed to the
22	recommendation of the task force, please raise

1	their hands.
2	(Show of hands.)
3	CO-CHAIR KAHN: And as long as we get
4	to 22, I sense we've gotten the 60-plus percent.
5	MS. BUCHANAN: That is correct.
6	CO-CHAIR KAHN: Okay.
7	MS. BUCHANAN: We have yes.
8	CO-CHAIR KAHN: So now let's move to
9	the next
LO	CO-CHAIR PINCUS: Actually, can I call
L1	a point of order
L2	CO-CHAIR KAHN: Sure, yes.
L3	CO-CHAIR PINCUS: to get
L <b>4</b>	parliamentary?
L5	CO-CHAIR KAHN: Sure.
L6	CO-CHAIR PINCUS: Okay. Could we make
L7	a general procedural motion that in the case
L8	where a motion has failed to pass to override the
L9	workgroup recommendation and that there's no
20	other motion on the floor, that automatically the
21	workgroup motion goes ahead, so we don't have to
22	keep doing this over and over again?

1	CO-CHAIR KAHN: So Harold has
2	suggested that we vote on a well, he would
3	like to make a motion
4	CO-CHAIR PINCUS: Right. I would like
5	to make a motion.
6	CO-CHAIR KAHN: that in cases where
7	the motion to change the task force
8	recommendation fails that
9	CO-CHAIR PINCUS: And there's no other
10	motion on the table.
11	CO-CHAIR KAHN: and there was no
12	other motion on the table, that we would
13	automatically support the task force position,
14	and thus we wouldn't have to vote twice. Is that
15	acceptable to the legal opinion of the staff?
16	CO-CHAIR PINCUS: I believe according
17	to parliamentary, Robert's Rules of Order, you
18	can do that.
19	CO-CHAIR KAHN: Okay.
20	DR. AMIN: So the voting procedure
21	that we've agree to is not that. However,
22	recognizing that the committee is trying to move

1 things along, and we are trying not to just have 2 arbitrary sort of rules of procedure, we can agree to move forward with that. 3 4 CO-CHAIR KAHN: Okay. So all in 5 favor, then, of a blanket reversion -- do you have a --6 7 MEMBER HALL: Well, then is it 8 portrayed as a unanimous decision? 9 That's how -- because, yes, DR. AMIN: 10 that's the only thing that we have to work with. 11 CO-CHAIR KAHN: Well, that would be 12 Well, actually, let's do this. the case. made a motion. I second the motion. Now let's 13 14 have a discussion. No, we have to have a discussion of it because -- and the discussion 15 16 just began. And the point is, the question is 17 whether it would be considered unanimous, I 18 guess. 19 MS. O'ROURKE: So we don't actually 20 have unanimous decisions at MAP. The vote is 21 really just to get to a decision category.

don't pass along that 24 people voted, 18 voted

conditional support, the others disagreed with 1 2 that motion. So we just say that MAP conditionally supported the measure. MAP members 3 4 said X, Y, and Z. However, others raised 5 concerns about A, B, C. So there's not --CO-CHAIR KAHN: That's true. 6 7 people have been -- we haven't gotten a unanimous 8 vote on any of this really. So, Bill, do you 9 have a comment? I'll just make the 10 MEMBER KRAMER: 11 observation that this will inform our discussion 12 tomorrow about voting rules and voting process 13 and consensus. 14 I remember earlier that there's just a comment made that sometimes consensus takes a 15 16 long time. And it's difficult to achieve and 17 measure. And voting can be more efficient. I 18 think we've seen a situation which voting 19 actually makes it less efficient right now. 20 I think we could actually say, 21 modifying Harold's suggestion, that we could say 22 that we have a consensus that when a proposal has failed to move forward, in other words there's not consensus, that the measure will revert, that if there's consensus, it will revert. We don't have to have a vote. We can say, we can agree by consensus that that's what we will do.

So I just, I don't want to get us overly tangled up, but just an observation and a suggestion about we could make a better use of consensus --

CO-CHAIR KAHN: Well, I would like to accept that as an amendment. But I'm not sure that will pass muster with the staff because, in terms of their advice to us.

MS. MUNTHALI: Yes, I think we should take a five-minute break here because we want to make sure that we're not making this painful for you. But we also want to make sure we're being consistent and not changing rules midstream. We understand tomorrow we're going to discuss this further. So we need to --

CO-CHAIR KAHN: Well, before we do that --

MS. MUNTHALI: Yes.

CO-CHAIR KAHN: -- I just want to go back to Bruce and ask, do you -- or let me ask the broad question. The staff will make a determination whether we can or can't do it. But assuming we can, is there an objection to this by anybody? I mean, I just -- do you object to this approach?

MEMBER HALL: Well, I would just highlight the second part of Harold's statement. And that is that if this group confirms that there's no alternate motion, then I would be willing to accept that everyone is therefore saying. But I think there does have to be a confirmation that after one motion is voted and rejected, that there needs to be confirmation at that point that no one else wants to raise a motion.

CO-CHAIR KAHN: Well, yes, well, first, let me say two things. One, I think that is accepted. We have only gone to the reaffirmation when there were no more motions on

1	the floor. So I don't even think we I mean,
2	that, procedurally, if anybody else had made a
3	motion, we would have accepted it.
4	CO-CHAIR PINCUS: Yes, that's explicit
5	in my motion.
6	CO-CHAIR KAHN: Yes, that's explicit.
7	MEMBER SCHLAIFER: I just have one
8	comment and one
9	MEMBER HALL: Yes
10	MEMBER SCHLAIFER: possible
11	alternate suggestion.
12	CO-CHAIR KAHN: Marissa was going to
13	go. Then
14	MEMBER SCHLAIFER: I'm sorry.
15	CO-CHAIR KAHN: I'll recognize.
16	MEMBER SCHLAIFER: Well, one, just as
17	a reminder, as Bill said, we've got a thorough
18	discussion about this tomorrow. So I think if we
19	do anything today, it should be kind of a
20	temporary until we have time to
21	CO-CHAIR KAHN: Right.
22	MEMBER SCHLAIFER: thoroughly

discuss this tomorrow.

And the other thing is I think what we said, I think what you were saying was that if we vote down the modified recommendation but then we accept that the committee recommendation passes, I'm wondering, just for staff discussion in the five-minute break, if it makes sense just to say it goes back on the consent calendar because then we still --

CO-CHAIR KAHN: No, no, they said we couldn't do that. That's the reason we --

MEMBER SCHLAIFER: Well, they
originally said that. But I think that would at
least allow, there would be a vote. Otherwise,
there's no vote. And so I just, that's just, was
going to throw that out there just for
consideration.

CO-CHAIR KAHN: Okay. I'm sorry. On the phone, was it -- I don't know who it was.

DR. BAGLEY: Yes, this is Bruce
Bagley. Let me make this suggestion that if a
motion is made and fails, procedurally you revert

1	to the motion on the table. And what you could
2	do is at that point just say is there any
3	objection to accepting the motion on the table.
4	And then you only get, basically you're getting
5	only the no votes.
6	CO-CHAIR KAHN: Okay.
7	DR. BAGLEY: So
8	CO-CHAIR KAHN: Well, actually, that's
9	the best way to go.
10	DR. BAGLEY: that might speed
11	things along.
12	CO-CHAIR KAHN: Okay.
13	DR. BAGLEY: So, in essence, you're
14	voting on the motion
15	CO-CHAIR KAHN: Okay. So
16	DR. BAGLEY: but you're only asking
17	for objections.
18	CO-CHAIR KAHN: Well, procedurally, we
19	can do that.
20	MS. BUCHANAN: We can do that, yes.
21	DR. BAGLEY: Yes.
22	CO-CHAIR KAHN: Okay. Well, I think

you just solved the problem. And we just spent

15 minutes figuring it out. I hate to take a

break because -- let's just keep rolling because

-- what's the next one?

MR. BERNOT: The next measure is MUC17-262. That's the ST-elevation myocardial infarction with percutaneous coronary intervention. This is for the MIPS program.

The workgroup recommendation was conditional support for rulemaking. The condition was that this was pending NQF endorsement, with even some further guidance saying during the NQF endorsement review, the MAP encouraged Cost and Resource Use Standing Committee to specifically consider the appropriateness of risk adjustment model to ensure clinical and social risk factors are reviewed and included when appropriate.

They also expressed concern over the precision of the cohort definition, whether it was sufficiently large cost performance distribution within the measure.

This was pulled by -- identified by Carl with a new motion to refine and resubmit.

Koryn, are you stepping -- okay.

MS. RUBIN: I'm handling it in his absence. So the AMA put forward a motion to change the decision category to refine and resubmit based on some of the comments that were made by the American Heart Association and American Stroke Association in terms of the emerging evidence that's expected to change the way that patients with STEMI will receive care.

And it doesn't appear it's captured in this cost measure. And so it will have implications moving forward and does not appear this measure is ready to move forward.

I would just want to highlight that

CMS did engage in a pretty collaborative process

as they were developing these cost measures. And

we thank them for that. But with this cost

measure, we don't feel comfortable moving

forward.

CO-CHAIR KAHN: Okay. Did staff and

1	the task force agree?
2	MR. BERNOT: The staff recommendation
3	on this was a conditional support. That is also
4	what the workgroup came up with.
5	CO-CHAIR KAHN: Okay. Other comments?
6	Leah?
7	MEMBER BINDER: Just a question, if
8	the developer can
9	CO-CHAIR KAHN: Is the
10	MEMBER BINDER: insight on this?
11	CO-CHAIR KAHN: Oh
12	DR. NAGAVARAPU: Yes, we're online.
13	This is Sri Nagavarapu from Acumen. And I'm
14	joined by Dr. Do from Acumen
15	CO-CHAIR KAHN: Great, thanks. Leah
16	has a question.
17	MEMBER BINDER: If they could comment
18	on the objection raised by AMA.
19	CO-CHAIR KAHN: Yes, would you all
20	comment on the objection raised by AMA in terms
21	of the status of the measure for use?
22	DR. NAGAVARAPU: Sure. We'd be glad

to. So we think this is related to a concern expressed in the public comments regarding emerging recent evidence for multi-vessel PCI in selected STEMI patients.

We do have an approach. So multivessel PCI was considered during the measure
development process. As was mentioned, the
measure development process included a clinical
subcommittee with over 30 members from an array
of specialty societies. And Dr. Do can walk
through those.

So we did think about multi-vessel PCIs there in order to address it as carefully as possible. We came up with the specific approach that Dr. Do can talk about to both address issues with staged procedures, as well as in cases where there's multi-vessel PCI performed during the triggering event, the inpatient stay.

The cost measure does reflect

potential savings from reduced adverse events

from that performance. So that is included in

the cost measure.

Going forward, as evidence continues to emerge from ongoing clinical trials, I think the clinical subcommittee has expertise that makes it really well-positioned to continue monitoring that evidence and make adjustments going forward as needed.

But I'll turn it over to Dr. Do. He's a cardiologist on our team that's worked with this. So --

DR. DO: Yes, thank you, Sri. This is

Rose Do. I'm a cardiologist. And I had the

opportunity to work with the 40-plus members

represented, ACC/AHA, Society for Cardiovascular

Angiography and Intervention.

I do know that during the conversations we did talk about multi-vessel PCI. We understand the data that is out there with PRAMI, culprit, the Danish study with AMI and, therefore, did talk about staged PCI.

I also want to note that we are aware that there is new evidence emerging. The guidelines so far for 2015 in the ESC and AHA/ACC

guidelines state that a multi-vessel PCI for STEMI is still Grade IIb, classification raised from Grade III, Class III a few years prior.

So I think there's still some development to be occurring in the clinical community in terms of consensus for stating whether PCI, multi-vessel PCI should happen on the trigger date.

That said, I do want to reiterate what Dr. Nagavarapu said, which is that we do account for staged PCI in some of the service assignment rules that we have. So once we have the trigger, we are assigning a revascularization only if there are some diagnoses that are present such as a repeat STEMI or new angina, a new chronic disease that's found within the claims. So we tried very hard to be restrictive when that did occur.

And then I would say at this time that the measure does capture, you know, potential cost savings. So, if a clinician does decide to do a multi-vessel PCI, and they do have some

savings in the sense that there's fewer admissions for angina or revascularization needs, then that will be seen at this time.

We also have ways of measuring, just kind of following people, because I think at this time at least in the clinical community, we want to gather more data. And we do have that capability with our measure now.

We have some upcoming meetings with the clinical subcommittee. And we're going to be addressing these comments and then talking about service assignments and risk adjustments. So those are things that we can tweak.

I think also our process allows for us to consider new data as it's coming. I would say there are some meta-analyses in the last two months or so that have come about to bring discussion into mind.

And so I think our process does allow for us to kind of think how strong the data is, has it been widely accepted by the community, is it in the guidelines, is there something that we

should be accounting for. 1 2 Well, I'll pause right there to see if anybody has any other comments or questions. 3 4 CO-CHAIR KAHN: Comments? MS. RUBIN: Yes, just to speak also 5 6 just a little bit further is this is one of the 7 measures that the public hasn't seen the testing 8 data. And I'm not sure if the specs have been 9 updated. I do -- you know, and to what regard, because there was an additional clinician 10 11 subcommittee meeting after the clinical workgroup 12 met. 13 And so we're talking about something 14 that's, you know, we're talking about stroke, 15 pretty complex disease. And the public have not 16 had access to the most updated specs or the 17 testing data to really evaluate and then confirm 18 what's being stated. 19 CO-CHAIR KAHN: Are there any other --20 I mean, anything else from the developers? 21 DR. NAGAVARAPU: Yes, this is Sri

Nagavarapu from Acumen. Just quickly on that, so

we did go through a field testing process in which all attributed clinicians with at least ten cases received field testing reports on these measures.

Shortly after that time, we created a national summary data report on these measures that were published online so that people could get a sense of what the cost distributions were like and so on and compare them to their own field testing reports.

As the commenter noted, in December there was a round of measure refinement activities to discuss substantive aspects of the measure and ensure that the clinical subcommittees felt comfortable with them.

Those refinements were submitted to our team here. And we worked with them and incorporated them. We'll be sharing an updated national summary data report with the clinical subcommittee members before we talk to them in the upcoming refinement webinars that will hopefully share the sort of information that was

being requested there.

I will say that we did compute updated reliability metrics based on the changes that were recommended in December that have now been fully incorporated into the measures. And the reliability at the team level for the STEMI with PCI measure is at least .75, at a case minimum of 30. And so that surpasses the usual standard for high reliability.

And we'll be sharing these sorts of numbers with the clinical subcommittees soon and can talk with you on that about sharing updated numbers when we've got them.

CO-CHAIR KAHN: Okay. Let's go to Bill and then over there. And then let's call it a day. Bill.

MEMBER KRAMER: I have just a process question. I'm not a clinical expert to be able to judge or comment on these things, but a question for staff and maybe other experts.

It sounds like these are the kind of issues that will be discussed and resolved in the

endorsement process. And my understanding is 1 2 that we're not trying to second guess endorsement or other -- or judge the measures in that way. 3 4 There are clinical experts and others who will be 5 doing that. So our recommendation here is, my 6 understanding it came from the workgroup, was 7 8 conditional support based on the condition of NQF 9 endorsement and which -- and if it passes through endorsement and it addresses these concerns, then 10 11 it seems to me that's, that works from the MAP's 12 perspective. 13 I don't think this is the place to re-14 litigate whether something should be endorsed or 15 not. We don't have the expertise. We're not 16 charged with that. And so I'm comfortable with 17 the -- but I may be misunderstanding. So I look 18 for anyone --19 CO-CHAIR KAHN: Let me just --20 MEMBER KRAMER: -- to clarify that for 21 me.

CO-CHAIR KAHN: Where is this in the

endorsement process?

MS. O'ROURKE: So this has not been submitted for NQF endorsement, still in development from what we just understood.

But I think to your point, Bill, we can certainly pass all of these, MAP's input and concerns, along to the endorsement subcommittee. We can also make sure that the reports from that reflect the concerns raised by the AMA. But it's obviously to the committee if you feel this is a conditional support or a refine and resubmit.

Well, it already is conditional.

MS. O'ROURKE: Like if agree with the AMA's motion to refine rather than the workgroup's standing --

CO-CHAIR KAHN: So this is not an issue of endorsement. It hasn't gone through the endorsement process yet.

MEMBER KRAMER: Right. The condition, though, the recommendation from the workgroup was that it be approved, conditional upon NQF endorsement. Is that correct? So it hasn't gone

1	through NQF endorsement. Is that right? But if
2	it is, does go through endorsement and addresses
3	these concerns, then the question is are we okay
4	with that. That's
5	CO-CHAIR KAHN: Well, okay.
6	MEMBER KRAMER: Because it just seems
7	to me these are the questions that will be
8	addressed in the endorsement process, and we're
9	not trying to second guess that.
LO	CO-CHAIR KAHN: Okay. I don't know.
L1	Did I get a second on the motion? I'm not sure I
L2	did. So where are we? Do you want to do a
L3	motion? We need to do the
L <b>4</b>	MS. RUBIN: Yes, I would like to make
L5	a motion to move to refine and resubmit.
L6	CO-CHAIR KAHN: Okay. Is there a
L7	second?
L8	MEMBER QASEEM: Guys, this is Amir on
L9	the phone now. I second it.
20	CO-CHAIR KAHN: Okay. Amir seconds.
21	Is there any oh, I'm sorry. Bruce, I didn't
22	see I was going to ask

1	MEMBER HALL: Yes, this is the same
2	question I asked earlier. So it looks to me like
3	13 of these 22 measures are not NQF-endorsed.
4	This is one. And for all 13 of the 22, we're
5	saying we approve it if it gets endorsed. But
6	we've been told earlier in the day there's really
7	no feedback loop to take that information back to
8	CMS. So that makes me really
9	CO-CHAIR KAHN: But wait, wait.
LO	No, I don't think it's not a question of
L1	feedback loop. CMS will get a recommendation
L <b>2</b>	from us that we are only recommending, we're
L3	conditionally approving it based on the notion
L <b>4</b>	that it's got to be endorsed.
L5	The feedback loop is then to us and us
L6	taking action as to whether or not CMS ignored
L7	with our recommendation and went on and made it,
L8	you know, required it anyway.
L9	MEMBER HALL: Right.
20	CO-CHAIR KAHN: And then the question
21	is: what do we do about it?
22	MEMBER HALL: Which is their

1	prerogative, right?
2	CO-CHAIR KAHN: Yes.
3	MEMBER HALL: So they would be acting
4	on the assumption that we've conditionally
5	endorsed, when in fact this measure's not even
6	submitted for endorsement. And they
7	CO-CHAIR KAHN: Well, no, no, we are
8	
9	MEMBER HALL: They will move forward
10	with the work.
11	CO-CHAIR KAHN: No, in defense of us,
12	we're only recommending conditionally that it go
13	forward based on it becoming endorsed. If they
14	choose to go forward with it, then that's their
15	prerogative because they have authority to do
16	that.
17	What I think we'll discuss tomorrow
18	is, when they do that and let's say it doesn't
19	get endorsed, but when they do that, what do we
20	do. I mean, do we stand on our hands, or do we
21	go back to them and start some process and

complain and do whatever we can? Obviously, we

1	can only jawbone. But we haven't done that in
2	the past.
3	MEMBER HALL: Yes, this feels to me
4	like we're
5	(Simultaneous speaking.)
6	CO-CHAIR PINCUS: CMS doing it in
7	the absence of it being endorsed, or if it goes
8	ahead and gets endorsed and gets implemented, you
9	know, are we comfortable with that?
10	MEMBER HALL: Yes, I don't have a
11	problem with these ultimately getting endorsed.
12	I wish that were going to be true of all of them.
13	I think what we're doing is we're
14	actually taking a stand on something that's not
15	specified. That's how I see it, because, in
16	fact, the conditional statements on 10 of these
17	13 ask for very specific issues to be addressed.
18	And all we're doing is saying we hope that
19	happens someday. In the meantime, we're going to
20	let it move forward.
21	CO-CHAIR KAHN: Well, we're
22	MEMBER QASEEM: And another problem

that I raised when I was at the meeting is, well,
I think that that's important. And in this case,
I'm really strongly pushing for that we need to
just -- the measure is we don't even have the
full information if the measure is ready for
prime time. And whether the measure developer
decides to submit it to NQF for an endorsement,
that's their decision to make.

But at this point, the information that we have in front of us, what I'm hearing from the commission workgroup is that they were unable to make that decision, because they didn't even probably have the expertise. And they want NQF to look into this measure, decide whether it's a good measure or not.

And in that case, the measure developer needs to make a judgment call whether they go through the NQF process or not. But our decision needs to be is whether this measure is ready to be used for accountability.

At this point, what is presented in front of us and that's what I think Horan and AMA

has been saying is there is not enough information for us to be able to make that judgment call.

I don't think we need to get into this issue of what the measure developer does, what process they follow. We need to make the decision based on the information that's in front of us.

Can we make, do we have enough information that we can say this measure should be used for physician accountability and we firmly believe it's going to improve the patient outcome?

If you feel like that, we should give it a thumbs-up. If you don't feel like that and go with what the clinician subgroup is saying that we don't know and NQF needs to review this measure in detail, then it's a revise and resubmit.

CO-CHAIR KAHN: Okay. Let me -- and I know I need to go to Leah. But let me go to Kate, and then I'll recognize Leah, to sort of

have CMS response so we have some sense of -
MEMBER GOODRICH: Yes, I do feel like
we have this conversation every year just for the

But, anyway, glad to have it again.

So what I do want people to just, as a point of information for the committee, this measure, while, yes, we're sort of in the final stages of testing the measure as the developer -- actually, it's done, right? We just have to get all the information out there.

Just I do want to say the measure is specified. This is a specified measure. It is not unspecified. This is not just a concept. So I want to be really clear with everybody how far along this measure is.

The other thing just for the committee to know, and many of you know this but some of you who are newer may not, at CMS every single measure that we develop -- and we are the stewards of this measure. Acumen is our contractor. Every single measure we develop we put through NQF endorsement pretty much. I mean,

record.

there's been one or two here or there that are not used in accountability programs that we haven't.

But that's our practice, because we are required by law for our programs to use NQF-endorsed measures except where there isn't an NQF-endorsed measure for a particular quality area.

And so I just want folks to know that it is not -- I think Amir was saying we can decide, the developer can decide whether or not they want to. That is, of course, technically true. But we always put ours through endorsement. And this one will be put through endorsement.

CO-CHAIR KAHN: Leah.

MEMBER QASEEM: And, Kate, if you're saying that it's a revise and resubmit, it will be perfectly fine to bring it back the next year. That's right. I mean, for this year's deadline the measure is not there yet. I absolutely agree. It might be in the final stages, but the

deadline was missed.

CO-CHAIR KAHN: Well, Amir, I'm going to let Leah comment. And then I'll come back, and I'll ask a question of Kate. Leah.

MEMBER BINDER: Well, from my understanding and I'm not a clinician either, but my understanding is that the objection was that there is some new science that has not been considered in the measure.

I was actually reassured by the comments of the measure developers that they have considered it. And they are continuing to evolve the measure. And then I am certain that the process of endorsement would also go over those same issues once again, and I think in a thorough way with all of the stakeholders.

So I would think from that point of view this would be -- that the recommendation of the workgroup would make the most sense.

And I will add that the measure is important, and that I would not want to see it delayed. I think this is a critical issue of

cost, resource use, and quality. This is a very high-stakes issue for purchasers and consumers. So I would very -- unless there's some major, major issue, I would not like to see it delayed.

CO-CHAIR KAHN: So let me ask a question of CMS then. Okay. So, if we go forward with the recommendation, it says conditional based on endorsement. Knowing, I mean, endorsement takes a while, and the measure is not quite ready to go to the endorsement process it sounds like.

So which cycle -- maybe you can't answer the question. But in an ideal world, which cycle is this measure in, going back to Amir's issue of is this going to go, you know, forward in the current cycle, or is it a year away because of the development process? I mean, what's the -- how much can you tell us?

MEMBER GOODRICH: So what I can say is, again, this issue does come up every year, which I think is one that NQF is trying very hard to resolve in partnership with a lot of us here

around the difficulties of the timelines between
endorsement and MAP process.

In an ideal world, we would be able to
have only endorsed measures come through the MAP

recommendations of the MAP very seriously.

process. We are not in that ideal world,
although I am very optimistic about the progress
that is being made. So we do take the

It is true. We could decide because this is a gap area, because it's really important, whatever we decide, that we want to propose it in rulemaking this year.

We also could decide that, based upon input from the MAP, that we want to wait another year. That is at the Secretary's discretion for us to do.

You are right. I cannot tell you exactly what we're going to do. I'm not allowed to do that.

CO-CHAIR KAHN: Well, let me say that at least as a matter of course, a recommendation that's conditioned on endorsement is making a

statement from us that it's not ready for this year, I mean, basically, I mean, in a sense.

Now, that doesn't mean you can't go ahead and do it. As you say, you have authority, and it depends on -- you're making decisions for a lot of reasons.

endorsed, you know, in time for this year's regulatory process even if endorsement was, you know, only a three or four-month process, because there's just not time. So, and as you say, you can't tell us what you're going to do. And you may not even really know yet what you're going to do.

But, you know, we are recommending in a sense against going forward with it without endorsement. Bill.

MEMBER KRAMER: Yes, and just to put it another way, I think by recommending to move forward conditioned upon endorsement sends a sense of the MAP Coordinating Committee that this is an important measure, fills an important gap.

It needs to be endorsed soon and put in place as 1 2 soon as feasible and appropriate. Saying that we should revise and 3 4 replace and send it back -- it's not repeal and 5 replace, is it? (Simultaneous speaking.) 6 That would send a 7 MEMBER KRAMER: 8 different message that we think a lot more work 9 needs to be done on this. It's not, the concept 10 isn't right. It's not, or maybe it's not 11 important, doesn't meet our selection criteria. And it would probably delay it for at least 12 13 another year. 14 And so I think we -- there is a difference between support with conditions versus 15 16 -- what is the other category? Revise and 17 resubmit. 18 CO-CHAIR KAHN: Okay. 19 MEMBER KRAMER: So I would recommend 20 that we support. I'll also say a lot of this was 21 discussed and I understand happened in the 22 clinician workgroup already.

1	CO-CHAIR KAHN: Right.
2	MEMBER KRAMER: And they've hammered
3	it out and came out with this conditional
4	support. Unless there's a compelling reason to
5	overturn the workgroup's recommendation, then I
6	think we ought to stick with the workgroup's
7	recommendation.
8	CO-CHAIR KAHN: Okay. So we have on
9	the table a motion to revise and resubmit this
10	rather than the recommendation of the task force,
11	which was conditional on endorsement. We're
12	going to have a vote now. All those in favor of
13	the motion raise your hands. You can vote for
14	the AMA.
15	(Off-microphone comments.)
16	CO-CHAIR KAHN: The motion was refine
17	and resubmit, yes. Okay. So all those in favor.
18	Okay. All those opposed.
19	(Show of hands.)
20	CO-CHAIR KAHN: And please vote on the
21	phone. Okay. Clearly
22	MS. BUCHANAN: Yes, so, for the motion

1	of refine and resubmit for MUC262, we have 1 yes
2	and 24 no.
3	CO-CHAIR KAHN: Okay. And if there's
4	no objection, then we will consider reverting
5	back to the consent calendar. And seeing anybody
6	object?
7	MS. RUBIN: Yes, I'd then like to add
8	a condition, to make a motion to add a condition
9	that, as the evidence emerges in terms of
10	changing with how STEMI is treated, that the
11	measure is looked at for and incorporate, looked
12	at whether the latest procedures are
13	incorporated. I'm trying better
14	CO-CHAIR KAHN: Okay. Is there
15	MS. RUBIN: phrase that.
16	CO-CHAIR PINCUS: That's an
17	assumption. I mean, that's really an assumption.
18	That's what the endorsement process looks at.
19	CO-CHAIR KAHN: Yes, but she
20	CO-CHAIR PINCUS: Yes.
21	CO-CHAIR KAHN: offered a motion.
22	CO-CHAIR PINCUS: Okay.

1 CO-CHAIR KAHN: I mean, I --2 CO-CHAIR PINCUS: Okay. CO-CHAIR KAHN: Is there a second to 3 4 the motion? Okay. There's no -- any second on 5 There's no second to the the phone? Okay. Is there any objection to go forward? 6 motion. 7 Okay. We'll go forward. Let's go to the next 8 one. 9 Okay. So we are going to MR. BERNOT: MUC17-310. I'll give you a bit of history from 10 11 The workgroup recommendation was this one. 12 conditional support for rulemaking with the condition of NQF endorsement and that it's 13 14 updated to reflect the most current clinical 15 quidelines. 16 The discussion of the workgroup revolved around the fact that there is a current 17 18 vaccine with evolving guidelines that may propose 19 a different vaccine with a different age group. 20 So this was pulled not previously, but 21 just in the discussion by Amir. And you did not

have a motion yet. Or are we just going to

1	discuss this?
2	CO-CHAIR KAHN: I'm sorry
3	MR. BERNOT: It's to Amir, sorry.
4	CO-CHAIR KAHN: Amir, are you on the
5	phone?
6	MEMBER QASEEM: Yes, I am.
7	CO-CHAIR KAHN: And do you want to
8	I'm sorry. First, the recommendation was
9	conditional
10	MR. BERNOT: Conditional support for
11	NQF endorsement.
12	CO-CHAIR KAHN: And was the staff
13	algorithm the same?
14	MR. BERNOT: Yes.
15	CO-CHAIR KAHN: Okay. Amir, how do
16	you want to proceed? Do you want to make a
17	motion, or do you just want to
18	MEMBER QASEEM: Yes, just quickly, I
19	just wanted to hear the conditions from the
20	staff. What are they asking for with these
21	things?
22	CO-CHAIR KAHN: I'm sorry?

1	CO-CHAIR PINCUS: What were the
2	conditions?
3	MEMBER QASEEM: What are the
4	conditions you guys are asking to put in place
5	CO-CHAIR KAHN: What is the condition
6	
7	MEMBER QUERAM: from the staff
8	perspective?
9	CO-CHAIR KAHN: Yes. What is the
10	condition that's placed on this by the task
11	force?
	MR. BERNOT: The condition was NQF
12	
	endorsement and inherently at that point that
12 13 14	endorsement and inherently at that point that it's updated to reflect the most current clinical
13	
13 14 15	it's updated to reflect the most current clinical
13 14	it's updated to reflect the most current clinical guidelines. But that was specified in the
13 14 15 16	it's updated to reflect the most current clinical guidelines. But that was specified in the condition.
13 14 15 16	it's updated to reflect the most current clinical guidelines. But that was specified in the condition.  CO-CHAIR KAHN: Okay, so it's
13 14 15 16 17	it's updated to reflect the most current clinical guidelines. But that was specified in the condition.  CO-CHAIR KAHN: Okay, so it's conditional based on endorsement which presumes
13 14 15 16 17 18	it's updated to reflect the most current clinical guidelines. But that was specified in the condition.  CO-CHAIR KAHN: Okay, so it's conditional based on endorsement which presumes that it will be updated, I guess, to the latest

add some conditions there and just want to make 1 2 sure that, you know, the medical exclusions are just not even listed in this one. 3 4 There's so many examples and even in a compromise, you can't really give this vaccine. 5 It's a very expensive vaccine, guys. 6 I mean, you 7 can keep it in mind what you're asking for. And, this does not even follow the CDC 8 9 recommendations. It's not even consistent based 10 on what CDC says. 11 They even talk about allergies, being 12 a compromise and all that. 13 So, essentially, this measure needs to 14 be revised. I can live with it, keep it under conditional, but I just want to add the condition 15 16 that at least reflect what CDC is saying. 17 CO-CHAIR KAHN: Does -- do we know, 18 are the developers available for this? 19 know what kind of stratification they've done on 20 this measure? 21 MR. BERNOT: We did not, but more of the discussion was the fact that there is a 22

1	potentially evolving change, significant change
2	to the vaccination guidelines and that's why the
3	workgroup had recommended
4	CO-CHAIR KAHN: But the guidelines,
5	because of what the proper vaccination is or
6	because of the issues that Amir's raising?
7	MR. BERNOT: It was about really about
8	the proper vaccination was the discussion.
9	CO-CHAIR KAHN: Because he's really
10	he's raising the issue that, you know, there's
11	certain circumstances you wouldn't provide
12	that you wouldn't choose to give the shot. Which
13	means that's not
14	CO-CHAIR PINCUS: But, is the measure
15	fully specified at this point? Does that include
16	the exclusions?
17	CO-CHAIR KAHN: I don't let me look
18	at CMS and ask the question. Pierre, do you know
19	anything?
20	DR. YONG: Yes, I believe it is fully
21	specified, but it does need to be updated. But,
22	let me check on the phone, I know there's CMS

staff there. Do you know, is the developer on the line?

MS. ERDE: Hi, this is Susan Erde, I believe the developer was supposed to be on the line. But the measure itself is fully developed. The situation is one of where the Zostrix vaccine is coming out. And, I believe that the developer wants to make sure that's incorporated into the new measure guidelines or measure specifications.

CO-CHAIR KAHN: The issue that Amir's raising really --

(Simultaneous speaking.)

MEMBER QASEEM: Yes, and so essentially we need to have the CDC recommendation or the exclusion who shouldn't be given this vaccine.

And then, generally, just for the group to think about, you know, it's an expensive vaccine. A lot of hospitals are giving it to underserved population, no insurance. I am not really sure, but I won't really get into the operational issue.

1	At the least, we should be following
2	what the current guideline by the government
3	says.
4	DR. AMIN: It might be helpful just to
5	clarify, the specifications that we have doesn't
6	include any exclusions.
7	So, what we can do is we could just
8	add this into the narrative, all of Amir's sort
9	of points of just clarification as elements that
10	the endorsement committee should review when it
11	goes through endorsement.
12	So, we could just add that to the
13	discussion.
14	CO-CHAIR KAHN: Okay. Would that be
15	acceptable, Amir? We'll add in your concerns.
16	MEMBER QASEEM: Yes, that's fine.
17	CO-CHAIR KAHN: Okay, good.
18	MEMBER QASEEM: I'm nicer over the
19	phone than I am in person, I guess.
20	CO-CHAIR KAHN: We love you in any
21	form.
22	Okay, let's move on to the next one.

DR. BERNOT: Sure.

so, the next one is MUC17-363. This is the intracranial hemorrhage or cerebral infarction. This is for the MIPS program.

The workgroup recommendation for this was conditional support for rulemaking similar to the STEMI measure, this was for NQF endorsement with the same specific considered appropriateness of clinical cohorts defining the measures.

That meaning that there was two clinical cohorts. There was the ischemic stroke as well as the hemorrhagic stroke where the two clinical cohorts that were brought up in the discussion.

And, that the appropriateness of the risk adjustment model for both the clinical and social risk factors.

The other thing that -- the point that was raised was the need to ensure that the measure appropriately handles transfers from tertiary medical centers, that they may receive transfer patients without -- with a more severe

presentation that may not be reflected in the 1 2 administrative data. That was all part of the conditions, 3 4 a long condition. NQF endorsement considering 5 then with some guidance of what things they should be considering. 6 7 CO-CHAIR KAHN: Okay. And, then who 8 9 MS. RUBIN: The AMA pulled this. 10 CO-CHAIR KAHN: The AMA, okay. 11 Yes, we pulled this and MS. RUBIN: 12 made a motion for refine and resubmit. The conditions placed on this measure 13 14 are structural issues and potentially will change the intent. As highlighted in the comments the 15 16 MAP received and then in the Clinician Workgroup 17 discussion, there's many questions that need to 18 be answered and further examined, including the 19 appropriateness of including the two conditions 20 within one cost measure. 21 And, each requires treatment from 22 different specialties. And, you're including it

1	all in one measure.
2	And then, also, the lack of precision
3	of the coding to capture patient severity.
4	So, also, to, you know, ensure that
5	there is really the appropriate risk adjustment
6	in there.
7	CO-CHAIR KAHN: I'm sorry, this is up
8	for endorsement or this has been endorsed?
9	MR. BERNOT: This is the exact same
10	thing as the STEMI, so it will the CMS had
11	stated that this will go through the endorsement
12	process.
13	CO-CHAIR KAHN: Okay. Has anything
14	that's been said not, I mean, not included in the
15	comments that are going? I mean the comments
16	okay, do you want to have a motion?
17	MS. RUBIN: Yes, to me, the type of
18	the the comments that were made and the
19	conditions placed on it warrant refine and
20	resubmit. We're looking at potentially a whole
21	different measure in the end.

CO-CHAIR KAHN: Okay, but, so there's

1	a motion to refine and resubmit.
2	Is there a second for this?
3	Okay, sorry, no second and there is a
4	recognition of all your comments in the
5	conditional. So, let's go forward with the next
6	one.
7	MR. BERNOT: Okay. Do we need to make
8	sure there's no objections from any
9	CO-CHAIR KAHN: No, no, but there was
LO	no motion. I mean, the motion but the motion
L1	wasn't seconded. So, there's no real motion.
L2	MR. BERNOT: My error, my error.
L3	CO-CHAIR KAHN: Unless I'm sorry,
L <b>4</b>	Derek?
L5	MEMBER ROBINSON: Yes, I just had a
L6	question in terms of, I mean, I share some of the
L7	concerns around the, you know, the cohorts and
L8	how management of those cohorts can vary quite
L9	substantially.
20	When we look at the total cost, and
21	this may be a little bit more down in the weeds,
22	but does this include like air ambulance

transport and all the things associated with the 1 2 cost for these patients? DR. YONG: Sri, are you still on the 3 line? 4 5 DR. NAGAVARAPU: Yes, I'm here. 6 So, this is Sri Nagavarapu from 7 Acumen. 8 One thing we wanted to clarify is that 9 we think there is an important misunderstanding in the comments that were received by the MAP 10 11 about these two conditions. 12 This was actually an issue that the 13 clinical subcommittee, when they started 14 deliberating on this early last summer, was 15 sensitive to. 16 And, what they did was they subgrouped 17 the two conditions separately. So, what this 18 means is that observed spending for ICH episodes 19 are compared only to expected spending for ICH 20 episodes and observed spending for stroke is 21 compared only to expected spending for stroke

episodes.

So, the way that's accomplished in the 1 2 measure, the way subgrouping is implemented is that there's entirely separate risk adjustment 3 models estimated for each case. 4 5 So, ICH is only compared to ICH and stroke is only compared to stroke. 6 7 result of that is that no clinician or clinician group is penalized or rewarded simply if they're 8 9 focusing ICH cases or only on stroke cases. And, this is something the clinical 10 11 subcommittee chose to do specifically for the 12 types of concerns that were raised in the cost. 13 After the subgroup calculations are 14 performed, they're brought together in one The clinical subcommittee chose to do 15 16 this for several reasons. 17 One is that the subgrouping approach 18 that we kind of just ran through directed the 19 clinical condition concern. The second is that often the same sets 20

It may also be the case that several of

of clinicians may be involved in both types of

cases.

21

1 the included services that are assigned such as 2 adverse events and complications may be common across the two cases. 3 4 And, then ICH cases were perceived as 5 extremely important to address and not segment them off. 6 7 The final point is that Medicare does 8 consolidate these cases into a single set of 9 DRGs. In terms of the inclusion of certain 10 costs, the clinical subcommittee did decide to 11 12 include certain types of costs like ambulance and air transport and exclude other costs that they 13 felt were not in control of the attributed 14 15 clinician. There's also several other issues that 16 17 were brought up regarding risk adjustment. 18 is dealt with by --19 CO-CHAIR KAHN: I think we got the 20 picture. 21 DR. NAGAVARAPU: Okay. 22 CO-CHAIR KAHN: That was very helpful,

1 though. 2 So, where are we? Where are we? MS. O'ROURKE: So, I believe it was 3 4 one -- Amir had pulled the HIV screening measure, 5 MUC17-367. Everything else was still on the consent calendar. 6 7 MR. BERNOT: Correct. So, I can go 8 through the 17-367. This is the HIV screening. 9 The workgroup recommendation for this was conditional support for rulemaking. 10 11 They requested the standing committee 12 review the patient cohort definition how 13 community is handled in the endorsement process with the --14 So, this would be a recommendation for 15 16 NQF endorsement with a very particular requests. 17 And, this was pulled by Amir, again. 18 So, we do not have a motion. So, Amir, it would 19 be up to you whether this is for discussion or --20 CO-CHAIR KAHN: Amir, on the HIV, do

do you want to express concerns or have a motion?

you want to -- what are your concerns?

21

22

I mean,

MEMBER QASEEM: So, yes, and I'll just express concerns and then that makes me feel better. I'll leave it.

As you do know, all of you are in primary care, one-time screening in a lifetime, it just doesn't make sense. That has no evidence that shows that one time screening is going to have a better clinical outcome.

And this, I mean sure that we have actually we had discussed also, I don't actually -- so, they are in our population health committee, this keeps on coming up.

And, for whatever the reason, on the back for years, then we come in here and the measure is going to get revised and nothing get revised.

And, the reason I wanted to pull this measure and want to bring this issue up is, many of these conditions supports and all that and we had asked for changes from the measure developers.

The reality of things is, many of

these measures don't change. We hope they will, 1 2 and they don't. And, that's my little spiel based on the evidence. 3 4 So, it's a one time screening in a 5 lifetime doesn't make sense. CO-CHAIR KAHN: If I can ask CMS, so, 6 I mean, just logic, why was this included? 7 MEMBER GOODRICH: This is a measure 8 9 from the CDC, is that correct? And, obviously, HIV screening is important. 10 11 I think for the issues that Amir 12 raises, I'm not going to speak to them clinically 13 except to say that, those are exactly the kind of 14 issues that, through the development of the measure and the endorsement process would 15 16 definitely come up. 17 So, you know, I think sometimes things 18 that come up at the MAP may or may not get 19 addressed in the measure because of the MAP 20 process, but they may get addressed because of 21 the endorsement process. But, I don't know the answer, Amir, to 22

1	your question about why that particular choice
2	was made for this measure. I would have to defer
3	to the CDC.
4	CO-CHAIR KAHN: And, this measure
5	so, it hasn't gone through is it
6	MR. BERNOT: It is not NQF endorsed.
7	PARTICIPANT: And, this is the CDC
8	here. I can speak to some
9	CO-CHAIR KAHN: Oh, yes, thanks.
10	PARTICIPANT: of the concerns.
11	CO-CHAIR KAHN: Yes, please respond.
12	PARTICIPANT: So, whether you could
13	I assume that there are individual concerns
14	with the once ever screening, but it's not valid
15	to say that there's no evidence for that because
16	there is evidence for that, it is not just the
17	CDC recommendation, it is also a USPSTF
18	recommendation with a Grade A assigned to it.
19	And the USPSTF is reviewing that and
20	is expected to re-endorse it again with a Grade
21	A.
22	So, the evidence for this

recommendation is solid, that it does have meaningful individual and public health ramifications.

In terms of not making any changes to the measure, this measure did go before the Population Health Committee with NQF and, based on some concerns about how we had defined the denominator, the measure was reconfigured to expressly address those particular concerns around the denominator which is why it didn't move forward with that review.

So, I think it would be fair to say that we do take concerns when expressed very, very seriously and we have tried to make changes.

So, this measure has gone through changes in response to specific feedback we have received including from previous iteration when we visited it at NQF.

CO-CHAIR KAHN: Okay, so it's in the endorsement process? It's been --

MS. MUNTHALI: It came through the process but it was not successful. It sounds

like the developer took that feedback, is 1 2 revising the measure. It's not back through the process yet. But, we expect -- it sounds like 3 4 it's coming back through the process. CO-CHAIR KAHN: Okay, we have 5 6 questions. 7 Sam? Thank you, Chip. 8 MEMBER LIN: 9 I'm looking at the specs on this and it says review for scientific acceptability. 10 The measure failed to meet the 11 12 scientific acceptability criteria, et cetera, et 13 cetera, and it explains why. While the committee -- review 14 committee was generously supportive of the 15 16 measure, several concerns were raised about the numerator and denominator. 17 Ultimately, the 18 measure failed the reliability criterion. 19 So, for what that's worth. PARTICIPANT: Which is this from? 20 21 MEMBER LIN: This is from the spec 22 sheets for the coordinating committee here.

1	Let's see, as to that particular it's the
2	summary of the NQF endorsement review in 2016.
3	MEMBER QASEEM: So, it's for would
4	it be possible that we just put this measure
5	through NQF process first without getting into
6	the debate about the evidence and all that,
7	right?
8	I think this measure needs to be taken
9	a look at and especially if it's going to go
10	through this process. Unless they then are
11	we going to see it? It's probably going to
12	population health first, right?
13	CO-CHAIR KAHN: Yes, I think it's
14	already gone to population I think you said it
15	had gone to population health.
16	MS. MUNTHALI: It failed, yes.
17	CO-CHAIR KAHN: And that concerns were
18	raised and that they're having to adjust the
19	measure.
20	I think
21	MEMBER QASEEM: No, but I thought that
22	I heard that they had revised the measure based

on the feedback and they're going to resubmit. 1 2 CO-CHAIR KAHN: Yes, yes, well, yes. PARTICIPANT: 3 Yes. 4 CO-CHAIR KAHN: Yes. But, you're 5 raising a fundamental issue, I think, that -let's continue with the points here and we'll see 6 7 where we get. 8 Mary? 9 MEMBER HALL: Thank you. 10 I actually was on a technical panel 11 for this measure. And so, I am going to recuse 12 myself from any voting. 13 But, I wanted to just add that, one of 14 the very often problems as someone who has over 80 measure NQF endorsed, speaking from NCQA's 15 16 point of view, is the difficulty in mapping a 17 guideline with great precision to a measure. 18 And, while, of course, Amir, you would 19 expect that in practicing in primary care, you would assess somebody's risk factors routinely 20 21 and you would re-screen them every time they had

had a potential new exposure.

When you're creating a -- and that's what a guideline would support.

But, when you're creating a measure, you have to aim, unfortunately, for the, I don't want to say the floor, but, you know, the most common set of scenarios.

And, without the capacity using claims to assess risk level, you can't make a measure that's risk appropriate. You have to just make the measure that would apply to the whole population without regard to understanding each person's risk status.

And so, that's the argument for the validity of a measure that is of -- and, I believe that this is a first HIV screening measure to be used in federal programs that the point of starting here is to say, everybody should at least be screened once and that's the starting point.

MEMBER QASEEM: But, they have better measure that, again, are being reviewed in NQF that gets into the whole treatment because we

haven't had that just one time screening is not going to help. You have to follow it through. So, that's one thing.

And so to respond to your comment, I absolutely agree that this is the floor, but if that's the logic we're going to follow, we need to apply it consistently across all the measures we have reviewed today.

We can't say that this one is okay for this one, but the others we have gone through, which I am not too sure if they were.

CO-CHAIR KAHN: You know, Mary, I haven't studied much of this stuff, but I did study when I was in graduate school, we looked at the old guaiac stool study and it really, you know, it wasn't worth doing them.

And, this one, I don't know, it sort of raises -- because you're basically saying we need, you know, you can't see it all the time so we ought to at least do one time everybody, if I understood what you said.

Okay, Bruce?

MEMBER HALL: Well, I think some of my 1 2 questions may have been resolved. But, I'm just trying to clarify what we're looking at here, so 3 4 this was a measure that was previously submitted 5 for endorsement and failed. The developer says that, I think the 6 7 developer said that what we're looking at now has 8 been revised. But, I don't know whether that 9 means it's going to be in the endorsement process in a timely fashion because our comments as 10 11 recorded talk about the standing committee 12 reviewing the cohort definition and so on and so forth. 13 14 So, are we looking at something that is going to be re-submitted for endorsement and 15 16 is that more than a year away then? 17 CO-CHAIR KAHN: Well, my sense is it 18 either has or will be re-submitted for

endorsement, right? Do we know? Can CDC tell us?

PARTICIPANT: Yes, yes, it will be revised and re-submitted. In fact, I thought

19

20

21

1	that was part of the message that you get from
2	the conditional endorsement.
3	We condition it on further review
4	under the appropriate NQF subcommittee.
5	MS. O'ROURKE: So, just to clarify,
6	MAP does not grant NQF endorsement as to just be
7	precise with language.
8	But, I think just to clarify, the
9	conditional support is pending on re-submission
10	for NQF endorsement.
11	CO-CHAIR KAHN: Yes, we understand
12	that.
13	MS. O'ROURKE: Okay.
14	CO-CHAIR KAHN: The issue is
15	PARTICIPANT: Yes.
16	CO-CHAIR KAHN: that our
17	conditional support is based on endorsement. It
18	may be endorsed, it may if it doesn't get
19	endorsed then that answers the question.
20	If it gets endorsed, then it would
21	meet our conditions.
22	MS. MUNTHALI: And, I just wanted to

ask the CDC, so it sounds like you did make those 1 2 revisions to what the MAP is looking at now. And, if that's the case, would you be ready to 3 4 submit this year to NQF? 5 PARTICIPANT: Yes, so, the previous 6 time we went through NQF, we didn't actually get 7 to the validity and reliability data because 8 there was a conceptual concern about how we had 9 constructed the measure. 10 We redid the measure and have gone 11 through all of the testing process with 12 Mathematica, policy research according to CMS's 13 Blueprint version, what, 13 or whatever. 14 So, at this point, we are in the final stages of developing all of the materials that 15 16 would be needed to take it back to NOF. 17 MS. MUNTHALI: Okay, that's helpful, 18 thanks. 19 CO-CHAIR KAHN: Okay, John? 20 MEMBER BOTT: Yes, just because it 21 seems like there's a lot of gray area for when 22 something would fall the conditional support and

refine and re-submit, I just thought I'd go back 1 2 to, you know, these seven assessment criteria. And, Sam made a good point when he 3 4 read, and correct me if I'm wrong, that it didn't test out as being reliable and valid. 5 So, the way these, you know, somebody 6 put a lot thought into these criteria and the 7 8 So, if it fails that seven components. 9 assessment number six, then it falls to refine and re-submit. So, it seems like until it's been 10 tested and reliable and valid, it -- we can't go 11 12 -- it can't graduate to conditional support. So, it sounds like it falls in the 13 14 refine and re-submit bucket. 15 CO-CHAIR KAHN: I'm sorry, so you said 16 17 PARTICIPANT: This is Abigail, can NQF 18 speak to that? Because what we got from them was 19 that it did pass all of those. And, Mathematica 20 is also on the line to discuss the validity and 21 reliability.

I did hear the person that brought

1 that up reference 2016, which wouldn't be the 2 current version of the measure. 3 MR. BERNOT: Sure, yes. I can address 4 that. 5 So, the preliminary assessment that we had had information on the testing. 6 We had it 7 looked at by a number of different individuals, 8 including our Chief Methodologist and others. 9 And, there is, admittedly, a need to 10 go through the endorsement process. That was not 11 supposed to be there. They thought there was 12 enough information that it is quite conceivable 13 that it was going to get through the endorsement 14 process on the reliability and validity. So, we did not, obviously, have their 15 16 endorsement data, but it was looked at internally 17 to see if it appeared to be sufficient to get to 18 that point on a quick glance based on the 19 information and it did. 20 So, that's where -- that's why we were 21 able, in our preliminary assessment, to pass it as saying that it appeared to be valid and 22

reliable.

I will admit this is an unusual situation where the data was close like that.

And, that's why we went through the extra steps including at the actual Clinician Workgroup,

Helen Burstin was the lead discussant on there also had looked at the data herself.

And we had this discussion. But it was close, but that's why we wanted the endorsement process and that's what the workgroup set as a condition.

CO-CHAIR KAHN: Okay. So, let me ask this question. There really are two issues here. I mean, I think, one, that everyone's brought up now which is just questioning the measure.

There is the more fundamental issue,

I think, from Amir, which is, is this the right

approach and should this measure be in here?

So, let me first as Amir, do you want to offer a motion? You were the one that pulled it.

I'm not getting any --

1	DR. AMIN: Amir, are you on the line?
2	CO-CHAIR KAHN: Amir, are you on the
3	line?
4	Well, okay. So then
5	DR. AMIN: Operator? Let me just make
6	sure. Operator, can you just verify that Amir's
7	line is open and is he on the line?
8	OPERATOR: Amir has actually
9	disconnected.
10	CO-CHAIR KAHN: Okay. So, that solves
11	that.
12	So, let's go to the second issue that
13	Bruce and others have raised.
14	Is there any interest in any kind of
15	motion from the room to change the status from
16	conditional with the proviso about endorsement
17	and the other issues that have been raised to
18	revise and re-submit? Okay, going once, going
19	twice.
20	(No audible response.)
21	CO-CHAIR KAHN: Okay, so we had our
22	discussion and clearly it's not ready for prime

1	time, but hopefully it will become so in the
2	process.
3	DR. AMIN: We'll clarify, we'll add
4	all this commentary.
5	CO-CHAIR KAHN: Okay. And so, is
6	are there any other measures for consideration?
7	MR. BERNOT: There are no other
8	measures. So, everything else would have
9	would be on the consent calendar that would just
10	need to be accepted.
11	CO-CHAIR KAHN: Do I have motion to
12	accept the consent calendar?
13	PARTICIPANT: So moved.
14	CO-CHAIR KAHN: Do I have a second?
15	Okay, we have a second.
16	All those in favor of proceeding with
17	the consent calendar. And, please vote on the
18	phone by email.
19	MS. BUCHANAN: And, we're just getting
20	in some of our phone ones.
21	CO-CHAIR KAHN: Okay. While they're
22	doing the tabulation, let me say, I'd like to

take a ten minute break. The Chairs have to 1 2 confer about -- with the staff about how we proceed with the agenda. 3 4 So, we'll determine what we do for the 5 rest of the afternoon. Can I just clarify? 6 DR. AMIN: 7 CO-CHAIR KAHN: Yes. 8 DR. AMIN: Can I just clarify one 9 thing? There was a measure, that optimal vascular measure for the MIPS program was pulled. 10 11 And, the motion did not pass. We're assuming 12 that that went on to the consent calendar that 13 was just voted on. Just want to confirm that 14 there's --15 CO-CHAIR KAHN: That one -- was that 16 the one where we made the transition to asking if 17 there was an objection? 18 DR. AMIN: Yes. 19 MS. BUCHANAN: Yes. 20 DR. AMIN: Right, it was before that. 21 So, I just want to confirm that the consent calendar has been --22

1	CO-CHAIR KAHN: There wasn't any where
2	we didn't vote on them but until we changed the
3	policy.
4	DR. AMIN: Okay, just wanted to
5	confirm.
6	CO-CHAIR KAHN: Yes.
7	DR. AMIN: For the record, that's what
8	it is.
9	CO-CHAIR KAHN: For the record, there
10	was no objection. We proceeded.
11	DR. AMIN: Thank you.
12	CO-CHAIR KAHN: Okay. And the vote?
13	MS. BUCHANAN: Yes. So, we have 26
14	yeses to support the clinician consent calendar.
15	CO-CHAIR KAHN: Okay, it's passed.
16	Sorry this took so long, but this is
17	what our process is.
18	So, we're going to take a ten minute
19	break and then reconvene.
20	PARTICIPANT: Then we'll get to the
21	hard stuff.
22	CO-CHAIR KAHN: Right.

1	(Whereupon, the above-entitled matter
2	went off the record at 3:41 p.m. and resumed at
3	3:56 p.m.)
4	CO-CHAIR PINCUS: Okay, so, we're
5	about to approach the report from the Hospital
6	Workgroup. And so, if everybody can take their
7	seats?
8	MS. O'ROURKE: And, do we have Ron
9	Walters and Cristie Travis on the phone?
10	DR. WALTERS: I am.
11	MS. TRAVIS: Yes, I'm here.
12	MS. O'ROURKE: Okay, excellent, I just
13	wanted to be sure.
14	MS. TRAVIS: Yes, we're both here.
15	MS. O'ROURKE: Be sure we had you guys
16	and your lines were open.
17	CO-CHAIR PINCUS: So, before we get
18	the report from the Hospital Workgroup, is there
19	anybody in the room from the public who wants to
20	make a comment?
21	(No audible response.)
22	CO-CHAIR PINCUS: Seeing nobody, what

about on the phone? Anybody on the phone that 1 2 wishes to make a comment about hospital programs? Bridget or Kathy, could 3 MS. O'ROURKE: 4 you open the line up for public comments? 5 If you'd like to make a OPERATOR: public comment, please press star, one on your 6 7 telephone keypad. That's star, one to make a 8 public comment. 9 And there are no public comments. 10 CO-CHAIR PINCUS: Okay. 11 MS. MARINELARENA: Great, thank you. 12 Hi, my name is Melissa Marinelarena. I'm the 13 Senior Director. I'm working on the MAP Hospital 14 Workgroup. I want to thank you for hanging in there this afternoon. We've got an hour and 15 16 hopefully we can get through this. 17 And it's kind of refreshing to see 18 that we weren't the only ones that had voting 19 I think my colleagues will agree with 20 that. 21 I'm just going to turn it over to my 22 colleague, Kate, it's been a long day, Kate.

She's going to provide just an overview of the 1 2 themes that our workgroup discovered. And then, turn it over to our co-3 4 chairs to give an overview of the recommendations. 5 And then, we're here to answer any 6 7 specific questions that you have on the 8 workgroup's recommendations. 9 MS. MCQUESTON: Great, thank you. So, we promise to make the introduction very quick. 10 11 This year, there were nine measure for 12 five of the hospital and setting specific 13 programs. 14 Four of the programs in the Hospital Workgroup purview did not have measures under 15 16 consideration for this year. And those are the 17 hospital acquired condition reduction program, 18 the hospital readmissions reduction program, the 19 inpatient psychiatric facility quality reporting 20 program and the value-based purchasing program 21 had no measures.

This is just a quick overview of the

meeting themes that were discussed during the hospital in person meeting.

The first major theme was the need to promote alignment and harmonization to reduce provider burden and provide better information to patients.

The workgroup discussed the need to balance costs and quality issues with measure burden that's placed on clinicians and providers as well, also remaining aware of measurement resource issues.

So, this discussion fell into four main buckets.

The first is alignment across payers.

They noted that it's important to align measures

across CMS programs and public and private sector

payers.

The next area was harmonizing similar constructs. They noted there was a need for increased harmonization of measures that evaluate similar constructs across setting and programs.

This, in particular, was discussed in

relation to MUC17-176, the dialysis facility medication reconciliation measure where the workgroup discussed opportunities to improve alignment of med rec measures across programs.

Next is the MAP role in advising on harmonization. They noted the importance of considering parsimony alignment and measure harmonization at MAP.

And, noted that there should be an active MAP role in examining measures used in CMS programs more broadly. And, this would include assessing opportunities across current programs and measure sets.

And, finally, discussed patient and family engagement and the importance of engaging these groups to improve measure harmonization.

The next item was balancing the need to address quality concerns with the need to ensure fair measurement. And we won't go too much into this because this has been discussed in depth today.

But they noted a lot of the same

issues that have been discussed previously in terms of balancing, driving improvements in key outcome areas with the need to ensure that there's fair attribution for providers that use measures and that the use of measures is reliable and valid.

And, they also stressed the importance of NQF endorsement as a mechanism to ensure that the measure is evidence-based, reliable and valid.

And, in particular, there looked at, you know, some of the issues with timing challenges, that measures are brought to -- that some measures are currently under development and have been reviewed for NQF endorsement when they're brought to the MAP.

And, that there were some concerns with the status of measure development where MAP struggled with balancing quality issues and patient outcomes with ensuring that there was the right information to evaluate if a measure is reliable, valid or actionable for providers.

So now we'll quickly go through the 1 2 discussion for each of the specific programs. So, I'll pass it to Cristie to summarize the 3 4 discussion of the ESRD program. MS. TRAVIS: Thank you, Kate. This is 5 Cristie Travis and I'm one of the co-chairs for 6 7 the Hospital Workgroup. 8 And so, we had measures that we looked 9 under the end stage renal disease quality 10 incentive program. 11 One which the workgroup supported was 12 the medication reconciliation for patients receiving care at dialysis facilities, that 13 14 there's an NOF endorsed measure. There were two wait list measures and 15 the workgroup had a robust discussion around 16 The decision was to 17 these measures. 18 conditionally support both of them based upon NQF 19 endorsement, they have not yet gone through the 20 endorsement process. 21 From the commenter standpoint, it was

broad support among stakeholders for the

medication reconciliation measure. 1 2 There were quite a few comments that came in on both of the wait list measures. 3 And, 4 I would say that the committee itself, our workgroup, also worked through a lot of these 5 same issues. 6 Perhaps the one to single out is the 7 8 issue of attribution to dialysis facilities 9 because, clearly, the transplant centers are also 10 a key player in whether or not patients are able to successfully get wait listed. 11 12 So, that was one of the major concerns 13 both discussed at the workgroup and with the 14 commentators. So, I'll turn it over to Ron for the 15 16 next one. 17 DR. WALTERS: You want me to do the 18 PCHQR, right? 19 MS. MCQUESTON: Yes, that's right, 20 Ron. 21 DR. WALTERS: Yes. So, the advanced 22 cancer center program had one measure which was a

readmission measure proposed by the exempt cancer centers because the all-cause readmission measure did not cover cancer patients.

This is an endorsed measure and was supported by the exempt cancer group. The Hospital Workgroup accepted that NQF endorsement and recommended that it be approved. And we're not taking individual questions, right?

The AFC quality reporting program also had one measure which was supported conditionally. That's a theme you're going to hear recurrent through this and was discussed extensively in your previous concerns.

The condition was that of NQF endorsement, I would say that the workgroup, in general, was very supportive of the rigorous review that the committee, sustaining committees perform for the endorsement process.

And therefore, recommended the measure of the hospital visits following an AFC procedure for conditional support pending that review.

There was, obviously, some discussion

regarding attribution and it was felt that the 1 2 review process would take that into consideration. 3 4 MS. TRAVIS: Thank you. In the 5 hospital outpatient quality reporting committee there was one measure under consideration. 6 7 lumbar spine imagining for low back pain and the 8 committee -- the workgroup did not support moving 9 this forward. This measure was not recommended for 10 continued endorsement by the NQF musculoskeletal 11 12 standing committee in 2017. 13 I think you can see on the slide what 14 the major concerns were from the standing There was significant concern that it 15 committee. 16 did not pass the validity standard. 17 There was one comment that was 18 received and they supported the workgroup's 19 recommendation. 20 DR. WALTERS: Okay, and then came the 21 inpatient quality reporting. And, again, there 22 was two risk-adjusted mortality measures.

those were extensively discussed and also an 1 2 opioid-related adverse respiratory event was extensively discussed. 3 There was a lot of discussion and, 4 again, it was felt that for the risk standardized 5 mortality measure, the best recommendation was 6 7 conditional support. 8 There were many conditions placed on 9 that. 10 The mortality measure was conditional 11 support and the opioid measure was refine an re-12 submit. There was a lot of discussion about 13 14 whether or not that was adequately defined at the 15 present time to even take to the standing 16 committee. 17 MS. MARINELARENA: Okay, thank you, 18 Ron and Cristie. 19 So, that's an overview of the measures 20 that the Hospital Workgroup reviewed. And I 21 think that there was one measure that was pulled off of the consent calendar. 22

1	CO-CHAIR PINCUS: I thought there were
2	several.
3	MS. MARINELARENA: So, there was one.
4	John Bott had pulled two, but I understand that
5	there is a conflict so those go back on the
6	consent calendar unless somebody else wants to
7	pull them.
8	So, as of right now, we have MUC17-
9	241, percentage of prevalent patients wait listed
10	for the end stage renal disease quality incentive
11	program.
12	MS. O'ROURKE: And, John, just to
13	confirm, he had pulled the two mortality measures
14	but he participated in the development, so he
15	declined rescinded his pull, if you will,
16	because of that recusal.
17	CO-CHAIR PINCUS: So we only have one?
18	So, there's only one measure that has been
19	formally pulled out?
20	MS. O'ROURKE: Yes.
21	CO-CHAIR PINCUS: Well, showed me up.
22	(Laughter.)

CO-CHAIR PINCUS: Well, now, but you 1 2 know, that -- is there any -- you know, from -are any other measures sort of that are being 3 4 proposed by the committee to be pulled for 5 discussion? (No audible response.) 6 7 CO-CHAIR PINCUS: If not, then we'll 8 proceed with the discussion of measure 241, is 9 it? 10 So, Bill, you were the --11 I was scheduled to be MEMBER KRAMER: a lead discussant, so maybe I'll be both the lead 12 13 discussant and a measure puller at the same time. 14 (Laughter.) 15 So, I think in my MEMBER KRAMER: 16 general comments first. First of all, I thought 17 it was very encouraging to see that, on the MUC 18 list, there were a number of very important outcomes measures. And, that's the direction 19 20 we're trying to move. 21 The ASC measure, the mortality measures and the readmissions for cancer 22

patients, I think are all good, important outcome measures.

And the Hospital Workgroup recommended full support or conditional support pending NQF endorsement for most of these measures which is also encouraging given the complex nature, attribution issues and so on for all of these.

And, I think, in particular, the ASC and the readmission for cancer patients measures fill important gaps in our current measure set.

So, I think there's -- this is a good set of measures to advance.

I do think the ESRD measures warrant further discussion and I suggest that we pull the 17-241, the percentage of prevalent patients wait listed, PPPW, for discussion, but I'm not recommending a re-vote just because I think there's some important issues raised in the --what this measure is and how it's discussed in the workgroup.

And, may be some useful information that gets captured in the discussion that would

be useful for CMS and its decision on whether and 1 2 how to use this. 3 Before I go on, though, to that 4 specific thing, were there other discussants? 5 Lead discussants who want to give some general I don't want to jump ahead to the 6 comments? specific thing if there are some other folks who 7 8 want to make general comments. 9 MEMBER KAHN: Hi, Bill, this is 10 Maureen Kahn and I think I was your partner as 11 one of the discussants on the hospital measures. 12 And --13 MEMBER KRAMER: And, I think Rich 14 Antonelli also. Rich, are you on now? MEMBER ANTONELLI: I'm on and I'm 15 16 Why don't we let Maureen go and then I'll 17 bat third in this lineup. 18 MEMBER KRAMER: Okay, that's fine. MEMBER KAHN: You know, I would agree 19 20 in the whole general comment, Bill, that you made 21 on the measures, you know, that it is good to see 22 some strong outcome measures.

And, I certainly have some reservation 1 2 for the wait list thing, putting a lot of accountability in one place and how do we take it 3 4 across to the transplant centers who make those 5 final determinations of who gets onto the list? And, you know, and I share John Bott's 6 7 when I look at all of our concerns on hospital mortality measures. 8 9 You know, running a hospital, we track our mortality measures and, you know, as you all 10 do, looking at specific diseases. 11 12 But, broad categories, I want to make 13 sure that we get actionable data to make change 14 and not just collect a lot of data. Anyhow, those are my only comments on 15 16 the measures. You know, I thought that a lot of them were well -- the discussions and the 17 18 feedback that was given, I think was on target. 19 MEMBER ANTONELLI: Yes, this is Rich 20 Antonelli. 21 So, I'll just weigh in. Basically, they pretty much mirror what Maureen had said. 22

1 had some concerns about the approach to the 2 mortality measures. The ESRD, I agree to the extent that 3 4 attribution to the dialysis centers causes an 5 opportunity, in fact, to fragment the types of accountability. 6 7 I do want to open this up to 8 conversation because I think that it's one thing 9 to measure -- to put a measure in place at one point in the system, but it's another to 10 11 misinterpret that there is a malaligned 12 accountability in that place. So, I would like to hear some 13 discussion about the ESRD wait list. 14 15 CO-CHAIR PINCUS: So, in the 16 discussion at the Hospital Workgroup, were there specific strategies or alternatives that were 17 18 suggested as a way to fix some of those problems? 19 It sounds like there wasn't a general 20 discussion of sort of moving to a lower category 21 because nobody's proposing that.

But, are there some particular

recommendations that were made or that came up in the Hospital Workgroup with regard to particular issues that should be addressed during the endorsement process?

MS. MCQUESTON: This is Kate speaking.

Yes, so, I can read the list of

conditions.

So, it includes NQF review and endorsement, also the workgroup recommended that the measure be reviewed by the scientific methods panel and the renal standing committee.

In particular, they asked that the endorsement process look at the validity of the measure, particularly, the risk-adjustment model, ensuring that it appropriately accounts for social risk.

And then, finally, the workgroup noted the need for the attribution expert panel to provide further guidance on the attribution model as well as the disparity standing committee to provide guidance on potential health equity concerns.

1 CO-CHAIR PINCUS: Why don't we open it 2 up for discussion here and see who wants to discuss it. 3 So, I see Raj is raised his hand. 4 5 That's good, we'll start with Raj. So, thank you. 6 MEMBER DAVDA: As a former nephrologist and a 7 8 transplant physician, I've got to say, I am so 9 glad to see this on the MUC list. I can't tell you how much disparity 10 11 and variation in care there is for patients 12 getting on the transplant list and for patients maintaining on the transplant list and the number 13 14 of living related transplants we do. Dialysis, at least on the commercial 15 16 side, is probably one of the most expensive 17 therapies we offer. 18 Transplant, I've tried to push many 19 ways and, really, there weren't any ways of 20 making anybody take accountability. 21 I do think accountability, as a former practitioner in this area, does rest with the 22

nephrologist and does rest with the dialysis facility.

They are the primary care givers of these patients in that sense. Most primary care physicians and other subspecialists will give up control of these patients.

The other thing I would say is that, and as Nancy said, I don't believe there's a risk of fragmentation, but a risk of more coordination of care.

When I was running the transplant program at two different places, it was very important when we had questionable patients to put on the transplant list if the nephrologist participated in the committee meetings.

And the nephrologists rarely came into the committee meetings to participate. And, a lot of times, we made blind decisions because we didn't know.

By putting the percentage of prevalent patients on the wait list, at least it allows the accountability and promotes nephrologists who

don't meet that criteria to go to the transplant 1 2 committee meetings and support and advocate their patients and at least provide information. 3 4 They may not have the final decision 5 but they can certainly influence the process and they can certainly ask for a vote. 6 7 All dialysis centers, by network rules, are aligned to a transplant center. 8 9 the nephrologist has to sign off on their thing every year. It has been just a paper push for 10 11 many years. 12 And, patients are disadvantaged by 13 that so I'm wholly in support of this and I'm 14 very glad this was on the MUC list. 15 Thanks. CO-CHAIR PINCUS: 16 Bill? 17 MEMBER KRAMER: Raj, I really 18 appreciate your sharing that because I had pulled 19 this list for a lot of those same reasons, but 20 not being a clinician, I'm glad you went first 21 and could describe it, what the situation is.

But, that's what we understand from

our perspective. There's -- there are issues around all the care provided by dialysis centers and the coordination, or lack thereof, fragmentation of care for people who need transplants.

And, the scary thing is, from what I can tell, we are lacking good measures of quality for ESRD patients and that this is -- moves in the right direction.

Is this -- are there questions about is this the right measure, the only measure? I don't think it is. I think it would be important for the MAP coordinating committee to send a send of the group to CMS and other measure developers that this is a gap that needs to be filled and needs to be -- an issue that needs to be addressed.

So, the question, I think, for us is, is this a measure that we think is good enough?

Is indicative of measure of quality? That it warrants moving forward conditionally on the endorsement, of course?

But, I think it is, but I also know it's a tough issue. And, since it's not a slam dunk from my perspective, but I was very encouraged to hear what you said, Raj, about the need for this from a clinician's perspective and it mirrors the perspectives we have.

CO-CHAIR PINCUS: I'm confused. So, why was there a vote like there was at the workgroup on this? This was not what you heard.

I mean, the feedback I got was this is not what you heard at the workgroup. What you heard at the workgroup was that a question as to whether this was the right place to be doing this, that the nephrologist is the keystone and that they don't even put them on the list.

So, they're going to have to, I guess you could say, aspirationally, it'll push the centers to push the nephrologists, I suppose.

But, you had a 60 percent vote. I

mean, if it had been -- if one more person had

switched, it would have been -- it would have

passed to remove. So, it was one vote away from

removal and we have kumbaya here. So, what's 1 2 going on? (Laughter.) 3 4 CO-CHAIR PINCUS: No, I want an 5 explanation. No, I mean, I think 6 MS. O'ROURKE: that's a great point. I think there were a lot 7 8 of very strong feelings about this measure at the 9 Hospital Workgroup. And some very difficult issues that 10 11 came up about this. I think it was a -- people 12 were trying to balance attribution concerns and 13 who actually makes the decision about putting a 14 patient on the wait list. 15 We did not have anyone, I think, to 16 represent the perspective Raj just brought up of 17 the transplant center. Rather, we had 18 representatives from dialysis facilities and I 19 think a nephrologist, but not at a transplant 20 center. I just want to represent people's 21 22 credentials correctly.

There were, I think, a lot of concerns about the potential disparities on the other side of who is -- to your concerns about who actually gets put on the wait list and that kidney transplants may not be equitably allocated and that there's issues there.

I think we also heard some strongly held concerns that dialysis facilities may be putting patients -- or profits over what might be best for their patient. And, I think that caused some alarm from some of the provider community that they were offended by that perspective.

And, I think it was a very challenging conversation working through some of these issues around disparities and potential social issues and conversation that I think people felt maybe insulted their professionalism and that whether they put their patients' best interests at heart.

So, I think there were quite a lot of things that came out on the table and the long list of conditions that Kate read reflects trying to broker a compromise of how we look at the

methodological challenges to this measure and ensure that it's valid and fairly attributed with some of the concerns that Raj and Bill just raised, that we need to drive progress in this area and make sure that we're addressing equity issues.

But, I think Melissa, Kate, Jesse, anything that I missed from the conversation?

DR. ROACH: No, I think -- I'm Jesse Roach, I'm a nephrologist at CMS.

And, I think you captured the contentiousness of the discussion.

I think one of the things you mentioned about whether it was fair to attribute this to the dialysis facilities, and I think that was -- I think that was one of the main issues that people had.

We brought up the point that, one, that this is sort of a shared attribution model. So, while the transplant facilities do have some responsibility, there is some responsibility on the part of the dialysis facilities to get

patients wait listed and to keep them healthy enough and get the education enough to keep them maintained on the wait list.

I think it's the same model that we have the readmission measure, for instance, where there's a shared accountability between the hospital and the dialysis facilities and encourages coordination of care.

The other point that I wanted to bring up is that the TEP, we had a TEP of experts that agreed that this was -- the dialysis facilities did have some accountability to keep these patients on the transplant list and getting them transplanted.

And, the -- and, a lot of this is education of the patient to have their options available to them so that they push for it and then maintain the whole status.

And then, lastly, from a patient perspective, this is very popular with patients.

Patients want to be encouraged to transplant -- get transplants. Patients want transplants once

they found out about it.

And, there are lots of disparities between who gets transplanted and who doesn't get transplanted.

DR. AMIN: If you don't mind, can I just add a few additional comments?

There were extensive public comments on this measure that I just wanted to make sure were accurately represented. And, we essentially have them in the room today.

But, Casey P. has provided extensive comments on the measure expressing concern primarily related, and I'd point everybody to these comments in the discussion guide, but a lot of it was related to the attribution model, specifically pointing out that there were social factors that significantly affect patients length and influence waiting list placement and wanted to make sure that that was included in the model.

And then, also, specifically pointed out concerns related to the reliability of the measure with an overall C statistic of .72.

And then, HA also sort of raised some 1 2 significant questions related to attribution around whether or not the dialysis facilities can 3 meaningfully influence patient include on wait 4 5 lists enough to provide measure -- provide responsibility for this measure. 6 7 I just don't want to -- I just wanted 8 to make sure we don't downplay the fact that this 9 took up a significant amount of the Hospital Workgroup's deliberations and there were 10 11 significant representation and sort of concerns 12 that were raised along these lines. This is Ron. 13 DR. WALTERS: 14 If I could say, I think you've heard 15 exactly the answer to your question. 16 This was a very difficult one. think it took us about an hour and a half or an 17 18 hour and 45 minutes to get to conditional 19 support. There were views all across the 20 21 spectrum on this one. 22 MS. MCQUESTON: And, just to recap,

because you mentioned the voting earlier. 1 2 So, there was a motion for do not support that came in very close, almost 50/50. 3 There was a second motion for refine and re-4 submit also along those lines. 5 But, ultimately, the final vote for 6 7 conditional endorsement was 23 yes, yes for 8 conditional support -- oh, sorry apologies --9 conditional support was 23 yes, three for no. 10 CO-CHAIR PINCUS: Jesse, you had 11 mentioned the fact that there was a kind of 12 shared accountability modeling this. 13 Can you explain? Because, what I'm hearing is that there was a concern that too much 14 of the accountability was on the dialysis 15 16 centers, not enough on the transplant centers. 17 And, just say a little bit more about how the 18 accountability model actually worked? 19 So, I'll use the example DR. ROACH: 20 of the readmission measure where, if you get 21 admitted to the hospital, it's partially the

responsibility of the hospital that discharges

you to make sure that the patient doesn't come back.

But, it's also the responsibility of the dialysis facility to follow up with that patient and to make sure that they are taken care of so that they don't end up readmitted to the hospital.

And, it's the same sort of idea with this measure where, yes, the transplant facility makes the ultimate decision as to whether to list someone, but a lot of things have to happen before they make that decision, before they even see that patient.

They have to have a patient that's educated about the transplant process. They have to have a patient that's healthy enough to do that. And, they have to have a patient that is sort of stabilized from a health perspective.

And, most of that is on the -- is under the purview of the dialysis facility.

So, both of these things -- so wait listing someone and getting the transplant labs

and doing -- that's the responsibility of the transplant center. But, getting the patient to a health status where they're able to be transplanted is the responsibility of the dialysis facility.

Furthermore, in the conditions for coverage, there are requirements that the dialysis facilities encourage referrals to transplant centers that they facilitate education of the patient so that they can be prepared to do the things for -- to ready themselves for transplant.

So, I think that it's pretty clear that some of the responsibility lies with the dialysis facility being the, for the most part, the primary provider of health for these patients.

CO-CHAIR PINCUS: So, what I am hearing, then, is this is not really a wait list.

I mean, wait list is not a functional thing you just sign somebody up.

They have to meet certain standards

with the patients that are set by the centers and there's not a cap on age or anything.

And so, and there's still going to be a predominance of real Medicare patients. So, and so, is this really sensitively --

I mean, so here, the stratification really makes a big difference. I mean, you could be a center that has a lot of 75-year-olds because of where you're located and, you know, you could be doing -- you could stand on your head and you're not going to get them to a condition that could really justify a bypass.

Is this going to be --

DR. ROACH: So, they're adjusted for age on the measures. And, the SWR, so, the initial wait list one is adjusted for certain comorbidities that would -- that we would expect to be increased mortality and make it less likely for you to listed on a transplant list. So that is taken into account.

And, also, one of the things that people brought up were that some facilities were

more served -- some transplant centers were more 1 2 stringent when -- than others. And so, at some facilities would be 3 4 penalized for having -- for living -- or for 5 being near a transplant center that was -- that didn't list people as well. 6 7 And so, they actually looked at that and I don't want to give you the exact number. 8 9 When they tried -- when they 10 controlled for regional transplant center rates, the reliability just went from 0.82 to 0.79 when 11 12 they adjusted for the regional differences in the 13 transplant centers. 14 So, it wasn't a significant factor. So, I have Derek and 15 CO-CHAIR PINCUS: 16 then Raj. I just wanted to 17 MEMBER ROBINSON: 18 rise in support of the measure. I think that 19 it's important to have that accountability as has 20 already been described. 21 And then, also just acknowledge that 22 there are substantial disparities in access to

transplants for kidneys.

And, I think while the -- some of the concerns expressed by the facilities regarding the attribution were noted, I think, also as something that's important.

And, I think any of you who drive through any impoverished communities today, one of the thriving enterprises and new buildings that, at least I see in the metropolitan area where I live, are dialysis centers everywhere.

So, we need to ensure that there's good visibility and to provide the right support system within the healthcare delivery system for an individual to be prepared for transplants.

And, I think everybody needs to play their role.

MEMBER DAVDA: Well, thanks.

You know, the other way to sort of think about this is that we worked for so many years, and CMS in particular, and I give them a lot of credit for setting up the ESRD QIP.

And, starting to make people

understand that this is a continuum of care for those patients.

That -- and so, the transplant centers, we tend to think of them as separate, but really, it's not. The patients don't think of them as separate, they think of them as continuum of care.

The second thing I would say is that for patients, transparency is very important.

Right now, they have no way of knowing which centers are doing a lot of transplants for the wait list and which centers are encouraging them.

And, you have to be aware that, even in today's age, there are still nephrologists out there that will tell a patient that they need to be on dialysis for a year before they can get transplanted or be referred for transplantation, which is just not true. It's countered and we have evidence based medicine, we know.

And so, although I agree that attribution is an issue, but I'm not really certain that attribution itself is an issue, it's

who should take responsibility for those patients and who should advocate for that patient in the healthcare system.

And, whether the dialysis centers want it or not, they are the people who advocate and should be advocating as CO's and CEC's the valuebased care is along that same line, they are the ones responsible for advocating.

So, I think this measure, although -and, I understand the resistance of attribution
from the facilities and not wanting any more
responsibility, but I think it's fully beneficial
to patients and I think it is a move forward
towards having a full continuum and continuity of
care for those patients.

CO-CHAIR PINCUS: Yes?

MEMBER GIFFORD: It's a little bit of an example of our earlier stretch over attribution dialogue.

I might suggest, and I would endorse the previous two comments on this measure, the importance.

It might be helpful in these 1 2 attribution things to add a recommendation or think about recommendations to CMS on how are you 3 4 measuring the impact these measures unintended 5 effects on access to care or shifts in care or stinting of care or, you know, denying care, you 6 7 know? 8 Because, you can start to game some of 9 these measures in ways that we never intended even though it was helpful out there. 10 11 And so, I don't think it underscores 12 the importance of using the measure and going 13 forward, but it's the advice to CMS in 14 rulemaking, how they might think about using the 15 measure, other consequences around it and 16 measuring it. 17 CO-CHAIR PINCUS: I think Rich --18 PARTICIPANT: Oh, sorry. 19 CO-CHAIR PINCUS: I think --20 DR. ROACH: That's just part -- I was 21 just going to say that that would be part of our regular maintenance of our measures to look at 22

unintended consequences. 1 2 For instance, with the transfusion measure, we noticed that there were regional 3 4 differences in coding and that was affecting how the measure was being implemented. 5 And so, they made changes and last 6 7 year, it was brought in. So, we regularly monitor those things. 8 MEMBER GIFFORD: 9 No, I appreciate 10 I think I'm just taking another step 11 forward. As you write rules for the program, it's in -- it's not just on when you come back, 12 13 but thinking about how that program design is. 14 This is not, as we've talked about here, this is not about endorsing the measure or 15 16 this measure spec, this is about how you're going 17 to use it in rules. 18 CO-CHAIR PINCUS: I think Rich 19 Antonelli on the phone had a comment, question. 20 MEMBER ANTONELLI: Yes, thank you. 21 Harold, it's sort of is related to the

point that you raised and I'm hoping that Raj is

still on the line.

In my brain, I'm trying to think about how this measure, you know, where it would fit in say to a logic model and that we could actually say that this measure would move us away from fragmentation to integration.

So, I'm obviously asking that question because that's what I'd love to see this measure do.

My concern is that, you know, if it's interpreted about where the air quotes now accountability lies, what would be the drivers, you know, what's in it for the nephrologists, for example, to reduce fragmentation and move toward integration?

Could you speak a little bit about how this measure would strategically support integration?

MEMBER DAVDA: Yes, thanks for asking.

So, I think it particularly helps in
the C models and the actual models. And, I -- it
clearly depends on how those value-based cares

are arranged and how the incentives are arranged within those value-based care.

But, I think this gives -- by including it in the QIP, it gives CMS some flexibility to be able to do that.

So, for example, if a dialysis facility was not meeting whatever the benchmark is set at, this would be an opportunity for them to engage their nephrologists, their -- the medical director of the facility and engage with the transplant center to see what are the particular issues.

And, we may -- and it's not, you know,

I know we kind of tend to think about it as it's

a -- the patient was too sick and couldn't be

wait listed. But, a lot of it is, and a majority

of it is really coordination of care.

You know, the patient didn't come to that one CT that they needed or they didn't come to the one appointment they needed. These are areas where really the facility who coordinates their care, which is the dialysis center can help

out and work with the transplant center to see 1 2 what the issues are to try to resolve. In terms of non-value-based care, I 3 4 think it proves a fair amount of transparency but we're moving very quickly in the dialysis world 5 toward value-based care. 6 7 And, I see this as, again, a micro 8 step forward, although I would love to see one 9 thing, is the transplant center part of the dialysis facility but it moves us closer to that 10 continuous care, you know, for that patient. 11 12 So, that's where I see the benefits of 13 this coming forward. 14 CO-CHAIR PINCUS: Any other comments on this measure? 15 16 (No audible response.) 17 CO-CHAIR PINCUS: Okay, and nobody's 18 interested in putting it forward another motion? 19 (No audible response.) 20 CO-CHAIR PINCUS: Good. 21 So, we have a little bit of time and, you know, a number of concerns have been raised 22

about the mortality measures. And, I just want to give people an opportunity that, if they do have any comments on the mortality measures to be able to discuss them now, if they -- the hospital mortality measures, if they do have any comments just to be able to open that up since there were some concerns and issues that were raised in the workgroup.

(No audible response.)

CO-CHAIR PINCUS: I guess everybody's tired out.

Melissa, did you want to make any other comments about the mortality measures in terms of just the concerns or issues that you think need to be addressed?

DR. AMIN: Harold, I think -- sorry to just jump in there.

Just, I think there is the question around the lack of variability that was raised in the comments before. I think we can just -- we can at least incorporate some of that into the current dialogue that we have for the measure.

1	So, I want to make sure that those
2	comments that were submitted before are
3	addressed.
4	CO-CHAIR PINCUS: Okay. So, any other
5	comments on any other issues pertaining to the
6	Hospital Workgroup?
7	(No audible response.)
8	CO-CHAIR PINCUS: Okay. So, I think
9	we're ready for public comment at the end of the
10	day.
11	MEMBER GIFFORD: Don't we have to vote
12	on the
13	CO-CHAIR PINCUS: Oh, we've got to
14	you're right.
15	Everything went so easy.
16	MEMBER GIFFORD: I think this will be
17	an easy vote.
18	CO-CHAIR PINCUS: Yes. I couldn't
19	hear you. Turn on your
20	MEMBER GIFFORD: I may have missed
21	what the public comment was on. Shouldn't we get

1	CO-CHAIR PINCUS: No, we've got public
2	comment before we voted.
3	MEMBER GIFFORD: Oh, sorry.
4	CO-CHAIR PINCUS: The very beginning
5	was public comment. Yes, at the end of the day,
6	we have public comments on the whole day.
7	So, okay, we have a motion to accept
8	the consent calendar? Second? Okay.
9	And vote by hand? Everybody in favor
10	of the consent calendar? And people on the
11	phone, phone people, send in yours.
12	MS. BUCHANAN: We have 22 in the room
13	and they are coming in on the phone.
14	Okay, great, so, we have for the
15	hospital consent calendar, we have 26 yeses.
16	CO-CHAIR PINCUS: Great.
17	Anybody opposed?
18	MS. BUCHANAN: And no oppositions.
19	CO-CHAIR PINCUS: Okay. Now, any
20	public comment from room about anything that was
21	discussed today?
22	(No audible response.)

1 CO-CHAIR PINCUS: Seeing none, any 2 public comment on the phone? If you would like to make 3 OPERATOR: 4 a public comment, please press star, one. That's 5 star, one on your telephone keypad to make a public comment. 6 7 And there are no public comments at 8 this time. 9 CO-CHAIR PINCUS: Okay. Well, I guess we're adjourned for the day unless you guys have 10 some final things that you want to --11 12 MS. O'ROURKE: Kate has one final 13 announcement, just to clarify about the PAC/LTC 14 vote. But before that, I just want to thank 15 16 everyone for sticking with us for today. 17 it was a lot of votes and a lot of good 18 discussion. 19 Thank you for bearing with us as we 20 worked through some of the process concerns and 21 hopefully it was some good case studies for tomorrow when we can put everything out on the 22

table and get your input about what we could do better next year.

You know, I think we want your feedback on everything, the voting process, the algorithm, the decision categories. So, if you have some time tonight to think it through, we're hoping to get all of your input on how we can do this better next year.

And, thank you for bearing with us through some of those challenging votes and revotes and a special thank you to Chip and Harold for expertly guiding us through all of that.

CO-CHAIR PINCUS: Chip did the heavy lifting.

## Bill?

MEMBER KRAMER: Just a question about since we did address or discussed the attribution model, or issues earlier today, we're going to come back to that again tomorrow or remodify the agenda?

CO-CHAIR PINCUS: No, modify the agenda so that we both have to do that.

Okay, so how is it 1 MEMBER KRAMER: 2 being modified? Maybe I missed that earlier. Well, part of it. 3 CO-CHAIR PINCUS: Do you have more that you want to talk about? 4 5 MEMBER KRAMER: No. 6 CO-CHAIR PINCUS: I mean, it would 7 shorten the day. 8 Yes, I think we're MS. O'ROURKE: 9 going to see if our colleagues from the Rural Health Workgroup could come earlier and give 10 their presentation in the morning and then 11 12 transition to the conversation about process 13 concerns and then if that wraps up early, just 14 let everyone head out. Okay, so it's moving 15 CO-CHAIR PINCUS: 16 rural health up earlier in the morning where the 17 attribution slot was and then continuing on with 18 the other ones around the voting process and the 19 decision algorithms as originally scheduled? 20 MS. O'ROURKE: And giving, yes, giving more time to that and then if that doesn't take 21 the whole day, letting people go early and --22

1	MS. BUCHANAN: And thank you, Erin.
2	And I just wanted to clarify, for the record's
3	sake, for the PAC/LTC consent calendar, CoreQ
4	short start discharge measure, MUC17-258, we had
5	23 voting yes, zero voting no and one abstaining.
6	I just wanted to get those numbers on the record.
7	And I wanted to ask my colleague to do
8	a quick announcement regarding the dinner.
9	(Whereupon, the above-entitled matter
10	went off the record at 4:45 p.m.)
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## <u>C E R T I F I C A T E</u>

This is to certify that the foregoing transcript

In the matter of: Measure Applications Partnership

Coordinating Committee Meeting

Before: NQF

Date: 01-25-18

Place: Washington, DC

was duly recorded and accurately transcribed under my direction; further, that said transcript is a true and accurate record of the proceedings.

Court Reporter

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### NATIONAL QUALITY FORUM

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# MEASURE APPLICATIONS PARTNERSHIP COORDINATING COMMITTEE

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# FRIDAY JANUARY 26, 2018

The Committee met at the National Quality Forum, 9th Floor Conference Room, 1030 15th Street, N.W., Washington, D.C., at 9:00 a.m., Charles Kahn III and Harold Pincus, Co-Chairs, presiding.

#### PRESENT:

CHARLES KAHN III, MPH, Co-Chair

HAROLD PINCUS, MD, Co-Chair

RICHARD ANTONELLI, MD, MS, Individual Subject
Matter Expert\*

JOE BAKER, Medicare Rights Center

LEAH BINDER, MA, MGA, The Leapfrog Group

JOHN BOTT, MSSW, MBA, Consumers Union

MARY BETH BRESCH WHITE, American Nurses
Association

RAJESH DAVDA, MD, America's Health Insurance Plans

TRICIA ELLIOTT, MBA, CPHQ, The Joint Commission DAVID GIFFORD, MD, MPH, American HealthCare Association

BRUCE HALL, MD, PhD, MBA, FACS, American College of Surgeons

GAIL HUNT, National Alliance for Caregiving MAUREEN KAHN, RN, MHA, MSN, Blessing Health System\*

MIRA IRONS, MD, American Board of Medical Specialties\*

WILLIAM KRAMER, MBA, Pacific Business Group on Health\*

- RACHEL LA CROIX, PhD, PMP, National Association of Medicaid Directors
- SAMUEL LIN, MD, PhD, MBA, MPA, MS, American Medical Group Association
- ERIN MACKAY, MPH, National Partnership for Women & Families
- AMY MULLINS, MD, FAAFP, American Academy of Family Physicians
- SHAUN O'BRIEN, JD, AFL-CIO
- AMIR QASEEM, MD, PhD, MHA, FACP, American College of Physicians\*
- CHRIS QUERAM, MS, Network for Regional Healthcare Improvement
- DEREK ROBINSON, MD, MBA, FACEP, CHCQM, Health Care Service Corporation
- MARISSA SCHLAIFER, MS, RPh, Academy of Managed Care Pharmacy
- CARL SIRIO, MD, American Medical Association
- STEVE WOJCIK, MA, National Business Group on Health

FEDERAL GOVERNMENT LIAISONS (NON-VOTING):

- MARY BARTON, MD, National Committee for Quality
  Assurance
- KATE GOODRICH, MD, MHS, Centers for Medicare and
  Medicaid Services (CMS)

## NQF STAFF:

TAROON AMIN, PhD, MPH, Consultant

JOHN BERNOT, Senior Director

KATHRYN BUCHANAN, Project Manager

KAREN JOHNSON, MS, Senior Director, Performance
Measures

ELISA MUNTHALI, MPH, Acting Senior Vice

President, Quality Measurement

YETUNDE OGUNGBEMI, Project Analyst

ERIN O'ROURKE, Senior Director

### ALSO PRESENT:

PIERRE YONG, MD, MPH, MS, Centers for Medicare and Medicaid Services (CMS)

\* present by teleconference

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1	P-R-O-C-E-E-D-I-N-G-S
2	(9:09 a.m.)
3	CO-CHAIR KAHN: Okay, let's bring the
4	group to order. And we're going to start as soon
5	as Pierre gets his food. We're going to start
6	with him on the input on measure removal
7	criteria.
8	So, for those of you who are looking
9	at your agenda, we're going to start with the
10	12:45 item, which is Pierre, because I think he
11	might have to leave this morning. So, we want to
12	cover first.
13	And then I guess will we do rural
14	next?
15	MS. O'ROURKE: We'll do the process
16	improvement and then rural.
17	CO-CHAIR KAHN: Okay.
18	MS. O'ROURKE: Process improvement.
19	CO-CHAIR KAHN: Okay. So we'll then
20	go to process improvement and then rural. So,
21	we've got people on the phone. And whatever
22	points you think appropriate on the phone, please

1 don't be shy to ask questions or make comments, 2 if you have them. And also, we will have points during 3 4 the day, I guess where we have opportunity for 5 public comment. Will they come at the same time, 6 is there only one time today or is there a few 7 times? 8 MS. O'ROURKE: We'll have one before 9 lunch and then one before we conclude for the 10 day. 11 CO-CHAIR KAHN: Okay. Okay, so 12 without further ado, Pierre, you have the floor. Thanks. 13 DR. YONG: Thank you very 14 much, Chip, I appreciate it. And so, good morning to everybody. 15 16 Very happy that today is our last official day of 17 the MAP in-person meeting since I've been here a 18 number of days, but it has been great this year. 19 I think this year, as you've noticed, we have taken a little bit of a different 20 21 approach to the MUC list as you've noticed. 22 MUC list is quite smaller than it has been in the past years.

As Kate mentioned yesterday, we've launched a new initiative called meaningful measures. Hopefully you have heard that presentation.

We've done a number of them across the MAP workgroups on the webinars as well as the inperson December meetings. And thought you probably have had a chance to hear it so I didn't want to repeat it here.

But in that sort of context, we've been thinking about that framework and applying it to our work in multiple different ways. So it's not just about what kinds of measures we have in our programs, but it's also about measure development, it's also about thinking about the current measure sets and what really makes sense to keep in those measure sets, what might be right for removal in the context of the criteria we have laid out.

So in that sort of framing, we thought it would be a great opportunity to take those in-

person meetings and get some feedback across the work MAP workgroups about some potential removal criteria for us to consider as we look at our existing measure sets.

And so we did these, they did this session or this agenda across those workgroups, and wanted to actually bring some of that discussion to the Coordinating Committee as well.

And so have wanted to bring this forward.

So, if you'd move to the next slide.

The question, which I have already said but, is, what criteria should CMS consider as it reviews its measure sets across its programs for its quality reporting and value-based purchasing programs?

So, in the next couple of slides, if you advance to the next slide please, outlines some high-level criteria that we put forward to the different workgroups and would love your feedback on those.

The first is that the measures are meaningful to patients and providers. That they

really are patient-centered, that they focus on the high priority quality areas that we've identified in the framework, and that they are current with clinical guidelines. And we had some discussion around guidelines yesterday.

Of note, sometimes we do have statutory specifics, statutory requirements that we need to meet with particular programs and have to develop particular measures for programs, so we need to consider that as well.

Measure type is actually something
that we have had lots of discussion across the
MAP workgroups as well as in the endorsement side
as well, but really do prefer this movement
towards outcome measures. That doesn't mean that
there isn't a role or place for process measures.

Sometimes there aren't outcome measures available in particular quality areas, and so, in that certain situations process measures can be very valuable. Particularly if they're proximal really and tied to the outcome of interest.

Given that we are moving increasingly towards value-based purchasing programs where payment is tied to performance on quality measures, I really do want to look at the variation in performance across a particular metric. And so that's another consideration we identified.

If you move to the next slide please. Performance trends, some of these measures have been in these programs for many years now, so one of the things we want to look at is looking at the trends and performance burden.

Rate mentioned the patients over
paperwork initiative that we have been working on
at CMS to really focus on decreasing the amount
of burden placed on providers and clinicians.
And so the amount of burden associated with
recording a measure, reviewing measure.

Also, it's a burden related to any particular measures is another consideration for us.

Unintended consequences is something

that's come up yesterday. We talked a little bit about that with the A1C measure and the diabetes composite, but certainly want to consider unintended, potential unintended consequences that have arisen from the use of any particular issue.

We also at CMS have significant operational considerations that come into play when we consider both implementing a measure, but also if there are challenges that may influence whether or not we decide to keep a measure in the program.

If you'd move to the next slide.

Alignment is something we heard a lot about
yesterday as well. These are, I think, all these
sorts of elements are things that are surprising
to anybody, but alignment of similar measures
across, not just our programs within CMS but also
with private pairs.

This also ties back to the burden issue that I mentioned on the prior slide.

De-duplication. So we want to try and

minimize unnecessary duplication of measures and measure concepts within measure sets to really focus on measures at the measure set, and they're individual measures, on particular high impact quality areas.

Quality improvement. So, the ability of a particular measure to drive quality improvement is ultimately what we want to see happen.

If you move to the next slide. And this is the last slide. Consideration of that measure and the context of the overall measure set.

So, certainly if you remove a measure, is that going to leave a huge gap in the measure set that's going to be creating an important gap that we don't want to create or will there be other measures that sort of remain that are more important to the overall measure sets.

So those are some overall initial draft considerations that we discussed across the individual workgroups. We got a lot of great

feedback.

Some of its reflected here in the slides I just went over, but would welcome any sort of feedback or reactions or comments based on what we presented. And I don't know if Erin or anybody else from Staff want to comment about certain discussions we had in December.

MS. O'ROURKE: Sure. So, to reiterate what Pierre said, this is something we've brought to each of the individual workgroups to get their input on.

I think overall people agreed with what Pierre put forward here. Some suggestions on additional areas to look at, ways to refine.

So I think we want to get your input.

We know measure removal is a topic MAP has wanted
to take up.

We may not necessarily have the statutory charge to do that right now at a measure-by-measure level, so we wanted to bring you, when we were discussing this with our colleagues at CMS, at least a chance to weigh in

on the input that they are using to make these decisions.

So, again, this is our chance to inform what CMS will be, how they'll be thinking about measure removal and what they'll be doing. And I think overall agreement at the workgroup level with some minor refinements. So we, I think, welcome any thoughts the Coordinating Committee has.

CO-CHAIR KAHN: I have a question.

When the measure developer, and this may be a
naive question, maybe I should know the answer
already but I don't think I do, when the measure
developers make a presentation to you of whatever
their thing is, do they, do you require any
quantification of burden?

Obviously it's different, depending on which, the claims are different then obviously claims versus IT versus something that requires to be done by hand, and then obviously it varies from setting-to-setting too, but is that, is there any attempt to quantify that?

I mean, this has been a constant discussion about burden since we've been doing this thing. Sort of going into the input.

And then the output is, I mean, into the trail would be looking at measures to see whether, gee, we've discovered over the time that the burden may not be worth the price of what we get out of it.

DR. YONG: Yes, thanks, Chip, it's a great question. I think when we consider any measure for any of our, both the MUC list and potentially for any other programs, we look at multiple aspects of any measure, right?

It's not just whether or not the measure is a good measure, has it been fully developed and tested and specified and reliable and all that kind of statistical testing, but we also consider things like the burden of that measure and whether we can operationally feasibly implement such a measure.

And in our regulations, when we proposed a measure, we actually do need to assess

the estimated burden of implementing any sort of change. And so -- and that we do quantify, both in hours and then dollars impact, the estimated impact of that measure.

So, I don't know, Kate, if you want to add anything.

MEMBER GOODRICH: The only thing I'll add is, I mean, it's impossible to be able to, with great precision, right, estimate the burden going in. We do our best based upon what we hear from stakeholders, based upon now our number of years of experience and implementation of measures.

With this being such a priority for this administration, one of the things that Pierre and his team has undertaken is trying to get at that issue with a little bit more detail so that we can have a better understanding of what the actual burden is for, for example, a chart-abstracted EHR registry measure.

So we have plans to sort of go out in the field to understand that a little bit better.

So I think our understanding of burden will increase over time.

And what's very important about that though, to be very clear, is to understand what you mentioned Chip, which is that tradeoff between burden and the importance of the measure to clinicians and patients.

absolutely heard from, I would say in particular the clinician community, is that when it's a measure that's actually helping me to improve care for my patients, the burden may be worth it.

And so, it really is about understanding that tradeoff from, I would say, certainly from the provider's perspective but also from the patient's perspective.

CO-CHAIR KAHN: One more question. I guess, this question of funding of NQFs activity in this area, is there anything authority wise that would prevent you from starting a project where our process could begin to wash through these existing, particularly the more aged

measures and make recommendations in terms of whether they still are relevant or meet the kind of, in how they compare with the criteria you have listed for removal?

DR. YONG: So I can certainly start, and maybe, Kate, you might want to add in. So, I think we'd have to go back to actually look at the statutory language, I don't know it off the top of my head.

But my recollection of the intent or the language that sort of created the, the MAP essentially, was to make recommendations about measures before they get implemented into programs. And that was assumed to be the focus, but we'd have to go look back at the exact language to be sure.

MEMBER GOODRICH: Yes, exactly right.

And the only thing I would add to that is, for
this contract in particular, because the
activities are required by statute, unlike a lot
of our other contracting activities, it goes
through a fairly significant, our scope of work,

goes through a fairly significant review by lawyers and OMB and people like that to make sure that we are doing what is within the statutory purview. So it would require all that.

So the answer is not definitely yes or definitely no, we'd really have to go back and understand that better.

CO-CHAIR KAHN: I guess I want to sort of, I want to broaden your thinking. Which is that I don't, I mean, the MAP is an entity that works with you based on the contracting based on the law, but that doesn't mean there are other authorities within CMS that wouldn't allow you to contract with NQF.

And if this is a priority of the administration then it seems to me that, and some of the issues that come up here really relate to this issue of what's the feedback loop. seems to me that if we were creative in looking, I mean, there are different things you can contract for.

So I just want to open your mind a

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little bit to that.

MEMBER GOODRICH: No, you're right about that. And not to get too into weedy government contracting stuff, it has to do with the funding source, right?

So if our lawyers, and we were to feel like we could not use the current contract, you're right, we could compete another contract for that work under a different funding source.

So, I don't want to get into the details.

So that is something we absolutely could think about. We probably couldn't do it under the existing contractual structure, but we certainly could think about that through another stream.

CO-CHAIR KAHN: Well, at least from my standpoint I wish you would.

MEMBER MULLINS: So, I want to preface some of my comments with some survey results we recently had. We recently surveyed our membership about value-based payment contracts and found some interesting results.

And one of the reasons I spoke so passionately yesterday about burden reduction was because of some of this.

So, 37 percent of our members have ten or more payers that they report to and 60 percent of them have at least seven payers they report to. So that's a lot of reporting and a lot of burden placed on them.

In that same survey, we found out that 62 percent of our members don't believe that quality measure and quality measurement improves patient care at all. So, we have a huge gap in what we're doing and what we're asking them to do and what they believe that does for these patients.

So, as we're trying to improve measurement and move measures in and out of programs, we need to be very cautious that we do it intentionally and we do it with plenty of warning to clinicians. So you can't just have a measure there one year and take it away the next, because the infrastructure it takes to report

that measure is timely and costly.

And so you build it up, you build the process up, you get your EHR ready to do it and then the next year that measure is gone. And that is like pulling the rug out from under the clinicians that are doing that work.

So I would caution against that.

There is the process, you know this, Pierre,
there is a process in MIPS to remove topped-out
measures over a four year process.

so that is probably a good time frame and a good process. I don't know if that could be applied to other programs, but the measure removal process in MIPS is the topped out process. So perhaps that could be something that could be applied across other programs.

MEMBER BOTT: Yes, so a couple questions. So, I noticed patient-centered was at the top of your list, so how are you planning to ascertain if a measure is patient-centered?

And a follow-up question. I noticed, again, it was at the top of the list, is there

any formal or informal prioritization amongst these criteria and measure selection/removal?

DR. YONG: So, maybe I'll answer both.

Or I want to respond to Amy and then I'll answer

John's questions as well.

So, taking John's questions first. I mean I think, yes, certainly in terms of patient-centered we do think in all of our sort of thinking, and that's reflected, I think in our materials as well, the patient is always is at the center of what we do.

In terms of sort obtaining feedback about measures that we might want to put into preference, I think we can get input in multiple different ways. Certainly, we've had lots of input, at least in our development process, we actually explicitly incorporate patient perspectives and care giver and family perspectives as part of the measure development process from the very beginning, as we develop measures.

We also make sure, and we work with

NQF to make sure there is explicit representation of patients and consumers across the MAP workgroup so that we get that perspective.

That's reflected in both the, and across all the workgroups at the workgroup levels as a coordinating committee level.

And then certainly we actually, as we put through all of our proposals they go through a notice and comment rulemaking process. And so do actively look for the comments submitted from patient and consumer groups as part of that process in addition to the rest of the stakeholders that we work with.

In terms of the overall criteria, they're not in any explicit order. Certainly if you have suggestions about things that are more, you think are higher priority for us to consider we'd welcome that sort of feedback on that.

And on Amy's comment about sort of timing and not wanting to, like, that it takes time to build infrastructure and not being careful and cognizant about the impacts of

pulling measures away, once that infrastructure is created we certainly are aware of that.

And certainly as we go through our processes, you know we, to remove any measures generally we go through this notice and comment rulemaking that is usually forecast out at least a few years. And so it's not usually immediate in terms of impact.

There are times when we suspend a measure immediately, but that's usually for safety concerns or some, you know, that we've done that. I remember a case where we did that where we had a diabetes, a post-op glycemic measure that we started to hear concerns of overtreatment or under-treatment of the patients because in order to maintain their glycemic control post-op we suspended that measure immediately.

But that's usually the only case, at least in recent recollection, that we immediately suspended a measure. Usually there is some forecasting out a few years.

MEMBER GIFFORD: A quick clarifying question. This is, we're looking at criteria for CMS to use not criteria that we would adopt and use with individuals. The --

CO-CHAIR KAHN: We have no role in removal, although I guess I'm suggesting that maybe we should have a role if we could work it out.

MEMBER GIFFORD: Yes. I guess then some feedback on it. And it may sound unique to the PAC setting, but on the burden side often

I've seen a discussion around measures that have high performance, low burden and they should be dropped.

But even when they have an average high performance, take some of the immunization measures, or, in our setting restraint measures, you have still a significant number, several hundred or a thousand providers, who are performing really poorly on that measure.

And so what you may want to think about in this programs is how you weight the

measures. So a low burden measure that meets all the other criteria you may not want to get rid of because they're also, they have other benefits outside the program, they are used by other individuals.

So even though doctors have multiple people, they often are using the same measures. Hopefully. Not always.

But, there may be more innovative ways to do that. And certainly I think, in any of the programs, you may want to know how you weight the measures. Because you really often do a disservice to the measures by equally weighting in almost all the programs.

and then the other thing is you're talking, I think, just to add in a clarification point on alignment with the programs, in several settings, particularly the PAC settings, there is a number of other measurement reporting programs or measuring programs that are not coming through the MAP at all that have similar measures that are specified differently than what's in the MAP

program.

And so there is a value to starting to think about how you align, not just there but across the board and other programs that don't come through the MAP that are not statutorily required.

CO-CHAIR KAHN: Carl.

MEMBER SIRIO: Yes, just a couple comments. So the first is, and, Kate, this goes go to the conversation we had about a couple measures yesterday where it looked like things over the course of the next 12 months might change in terms of the guidelines that underpin the measures.

And I was struck by your comment that you've got a process where you can suspend things relatively quickly. It just seems to me that, and if you have it, I apologize for not knowing about it, there needs to be a process where if in midstream, in the middle of a year, that in fact the world changes in terms of a new set of guidelines, that there's an internal process to

in fact determine whether or not, whether suspension is the right word or not, but some process to basically, to take a pause on whether or not there's enough concordance with what's currently going on when compared to what's changing in the world in terms of new guidelines.

It's just a thought in terms of being a little bit more speedy or timely given the fact you have to change the battleship with all your rules in terms of process.

You know, picking up on some of the things that Amy and you have said, and in particular I'm struck by the discordance between what people believe is actual helpful or not, I think there is value in thinking about maintaining measures even if there is an intuitive sense that you can take it away because performance is high in two spaces.

One is certainly the public health space around tobacco cessation and the like, screening for alcoholism. Because those are just important things to do.

And there is I think pretty good evidence among human beings. It's not just the medicine, that if you take your eye off the ball you're apt to start to have decrements in your performance.

So I think there are some baseline things that you just want to kind of keep your eye on.

But the other thing is, as you start to remove measures, in particular if in fact the data, or the survey data suggests that people actually don't believe this helpful, is that as you take things away to see what the change over time is, because if people don't believe this is actually helping, they may not be doing it other than for checking a box. Which is actually not the intent obviously.

so my only plea would be that as you remove things, keep the high priority things in place because they're just the right thing to do. And secondly, make sure that we have some process to actually measure decrements in performance

over time. I'll probably have some other stuff 1 2 to say but I'll think about it. CO-CHAIR KAHN: What's the volume of 3 4 measures that have been removed? I'm mean, sort 5 of how many? I mean, is it a lot, a little, a few, 6 7 I mean, is it a few, many? 8 I don't know the answer to DR. YONG: 9 I mean, every year we make proposals to 10 remove measures in programs. 11 It's usually on the scale probably 12 between, for most part on average, it's probably 13 between like, I'm going to guess, between like 14 three and six maybe per program, per year. And this number measures per program varies. 15 16 As I know for IQR it's a 50 but for 17 PAC it's, I think they're nine measures or seven 18 measures. So it really varies. 19 CO-CHAIR KAHN: Okay. 20 CO-CHAIR PINCUS: I quess this follows 21 up a little -- so, I mean, this follows a little bit of, I think in terms of thinking about how 22

the process of removal could inform our process so that we begin to learn what things have either haven't worked or have developed some problems, it would be useful if we actually got some kind of feedback so that when you're removing stuff like, so we know how many measures were removed and the reasons why.

I presume that you have some way of gathering data on all these criteria, and so if there is some way that we could also share in that information so that it gets fed back to us in some kind of report or some discussion sort of early in our process, hey, here's what we've learned about things that haven't worked or that things have worked for a while but no longer work, that would, I think, help inform our process.

CO-CHAIR KAHN: Erin. Sorry.

MEMBER MACKAY: Yes, I wanted to pick up on John's point about patient-centeredness and suggest that as we are thinking creatively about whether and how to leverage this group to provide

input or to help quantify burden, that we're similarly leveraging the patient and consumer reps that, Pierre, you mentioned are integrated, not only on this committee but in the workgroups, to provide that perspective about the value from the patient and care giver perspective, to be able to inform your decision making around this balanced point that you referenced.

Because it may be that there is a delta between measures that providers don't think are useful to patients and families and measures that patients and family themselves prioritize.

I'm sure there is much overlap but I think it's important to just to be making sure we're having both perspectives represented.

CO-CHAIR KAHN: Kate.

MEMBER GOODRICH: I think --

MEMBER KAHN: Hi, this is Maureen.

MEMBER GOODRICH: -- be helpful for us, and one thing for the NQF Staff to ask, because I just don't remember, in the reports that come out, about all of these deliberations,

as you mentioned, you do say the committee supported conditional, whatever the determination is, some members said this, some members said that.

What I don't remember is if it is outlined where the stakeholder perspective came from for members who said this and members who said that. I know there have definitely been times, internally for us when we're considering a measure for proposal, where we wanted to understand better who said what, I don't mean the person's name but the perspective, and I think we maybe didn't have that.

And that would have been helpful for us in sort of crafting our justification for proposed lever measure or internally for us to not propose a measure. That can be really helpful for us for our rulemaking process actually.

So to the extent that's able to be captured, that would be really, really great.

CO-CHAIR KAHN: And I'm sorry, now

I'll go to the phone. So anyone on the phone have comments or questions?

MEMBER KAHN: This is Maureen. I would tell you I echo a lot of the comments that have been said and really, it's very positive to see the work that's going on.

I would just suggest, and I'm sure CMS has seen this, as I sit on a regional policy board, the AHA has done a lot of work looking at the cost of collecting quality data and the administrative burden that came in there.

And if I recall it correctly, I think when they did their study it was about, the average hospital spent about \$700,000 a year just in collecting the data. Not even acting on the data, and taking it to make improvements.

And the larger institutions certainly exceeded the millions of dollars to collect the data. So I would say, I'm pleased to look at ways of how we're, you know, you're looking at making better alignment in the quality and even eliminating some of the measures that may not

have value.

But I do believe, as the last two speakers mentioned, the voice of the consumer in quality indicators and data, I think it's important and we shouldn't lose sight of that.

CO-CHAIR KAHN: Thanks, Maureen. Any other comments on the phone?

MEMBER QASEEM: Yes. Hi, this is

Amir. Very nice presentation, Pierre, I enjoyed
your criteria.

And there is not much to add to the criteria but more of a comment that you mentioned, that there are a certain number of measures that are getting removed. And I think it will be very helpful if, again, I'm a very visual person, something simple that you can, if you would be able to share, here's the list of measures and here is the criteria and this measure was removed because, a simple checkmark yes or no that this criteria, it said this measure failed on this criteria.

I think that's going to help not only

the MAP committee thinking but, for example, if I bring it back to ACP member physicians I can say, look, CMS is listening and this is why this measure was removed. And this information might be there, probably, because it should be public knowledge anyways.

And if it's not if you can help us, like he said, identify where this information is located and if not, if you can ask your staff to create something similar that will be, I will find it very helpful.

CO-CHAIR KAHN: Great, thank you, Amir. And then Tricia.

MEMBER ELLIOTT: Yes, thank you. Good morning. One, first, a comment. I know there is work obviously outside of NQF and there is work being done.

I sit on a committee that's looking at the blueprint and incorporating the patient perspective. We just met last week. So that's been really informative.

Just so others are aware, that looking

at the whole measurement development process and how do you get that voice of the patient, voice of the care giver into that process. So that's been very enlightening and eye opening to work with that group.

Second is just a clarifying question.

I notice that you mentioned in your kind of list of criteria a burden but then you call out operational issues separately.

From the joint commission's

perspective, we have always kind of thought of

those part in parcels so I'm just kind of

curious, from a clarifying, why you felt to pull

out that operational piece separate from burden

instead of using the umbrella term of burden.

DR. YONG: I guess one way to think about, I guess I am sort of thinking about it a little bit sort of external versus internal. One way to think about it, so burden may be more external. Sort of the burden placed on the provider or the patient to fill out the survey for example or to administer a survey or to

collect data.

Operational concerns I think primarily is more, the way I'm thinking about it, is more internal sort of consideration. Do we have the ability to collect, to receive the data, to calculate the rates to, whatever it is internally or there budgetary concerns with, you know, are there updates that need to be made to the system, can we make them in time in order to collect the data, those are the kinds of concerns when I refer to operational considerations.

MEMBER ELLIOTT: Excellent. Thank you.

CO-CHAIR KAHN: Other questions on the phone?

MEMBER SIRIO: Actually, I was chewing over what Amy had said earlier and I think that it warrants some restatement in terms of the stability of the systems over time. If there's stuff that's changing you got to take it out, if it's something that's not helpful it's not taken out, but to the degree that there is some

consistency year-to-year.

As I think about helping to run a health system, the cost associated with actually changing the IT systems alone, never mind reenculturating folks as to what's important from the perspective of the organization, the patient and the country in terms of your measures I think cannot be overstated. So I would just suggest that stability is really helpful.

CO-CHAIR KAHN: Great. Any other comments, thoughts? Going once, going twice.

Okay.

Now, I guess we're going to move to the, what's on your agenda is the 10:15, which is potential improvements to the pre-rulemaking process and our voting process. And I guess, are you going to say anything?

MS. O'ROURKE: Yes. Can I say -
CO-CHAIR KAHN: And I'm going to turn

it over to Erin and then David, Sam and Marissa

will be lead discussants here on this --

MS. O'ROURKE: Okay. So I've got a

few slides to go through but I just wanted to provide a little bit of context about what we're doing, how we've gotten here.

I think every year we want to make this a better process for you all and make sure that you're feeling that your voices are heard and we're doing this in the most efficient way, but in a way that allows you to express your opinion and feel we truly came to consensus.

I'll kind of harken back to some of Bill's comments yesterday about the best way to do that and how to ensure that the conversation is heard.

From our perspective, we have made this more structured over time. I think some of you have been on MAP since year one.

I do remember when we did not take formal votes and it was a purely conversation based process. We did hear some concerns that that led to the chance for people's opinions to be misrepresented.

People could have a, hear the same

conversation and have a different determination of what the consensus position was. So we implemented this voting process to ensure that we, everyone was clear on what the recommendation would be how we got there.

I think we also want to emphasize, we don't actually send the vote counts in the report that Kate was referencing to CMS, it's just what the decision was and the rationale.

We also try to capture any dissenting opinions, minority opinions. Make sure that's all in the document too so that the CMS Staff has the breadth of what MAP had to say and can consider all of those points when they're making their rules. Not just we conditionally supported this measure and they don't actually see any of the counts of the votes and things like that. So I did want to clarify some of that.

So if we'd go to the next slide I just want to briefly go over some of the ideas we had about how we can make this better.

We want to avoid some of the

frustration that perhaps people felt yesterday when the processing to be hampering the conversation you all wanted to have but ensure that we're coming to clear consistent decisions and that everyone is in agreement of about how the meeting proceeded.

So, some ideas that we had. We'd love to get your thoughts.

We don't want to force you to come to any decisions today, this is something we're going to continue to iterate on but we just want to get all of your ideas out on the table. And we can work through the off season, if you will, to bring back something when we bring you all back together next fall.

So there is no need to feel like we need to make decisions today, we just want to get all of your ideas.

So for the voting process, some things we thought about. We could revise the process to more closely follow Robert's Rules of Order.

Some things we did this year was to

add a vote to accept the consent calendar rather than just letting everyone know that it would be accepted. So we did have that affirmation from the committee that you accepted the calendar and there were no concerns with what was going through on that.

We also did want to clarify. The idea of how many motions to handle at one time and the role of the lead discussant isn't necessarily to put forward an alternative motion rather to share their thoughts, react to what the conversation is, their feelings on the measure, but making sure that we're handling things one motion at a time so that it's clear to everyone what is on the table.

We also thought we could, we heard that people do want a larger staff role and for us to be more active in the conversation, helping to clarify how we got to the results of the preliminary analysis, if people have questions, have the chance to ask us why we, or what we did or what lead us to that decision. As well as to

jump in and help our co-chairs theme your conversation more so that we're, people can know what the threads are, what type if input we're going to put into the report.

We also want to address the technical issues. I apologize again for the frustration of the voting clickers, I think we want to have a voting conversation there.

Do people like voting anonymously, are you comfortable voting publicly. Also, would you be comfortable moving to a system where perhaps you vote, we had a little app on your phone or through the web that you could vote that way rather than our big clunky blue clickers. So if people could share some thoughts on that topic too.

We want your thoughts on, should there be a default position. So for the workgroups, this year we used the Staff preliminary analysis decision that if people couldn't come to consensus that would be what would come through to the Coordinating Committee.

For the coordinating committee, we used the workgroups recommendation. And the idea yesterday was you needed a 60 percent majority to overturn that.

Or do people want to keep voting until there is consensus on a decision category. And then similarly for the Coordinating Committee, are you comfortable with split decisions coming to you all. We worked to try to eliminate that on some input from this committee in the past that you did want a solid recommendation from the workgroup to react to.

We also heard some feedback that we should clarify abstentions prior to each vote.

So, Bruce, to some of your concerns about what's our quorum number, what are we looking for here so that the math is clear to everyone of how we're getting to what we're getting to.

So, next slide. So, this is just the first bucket of topics that I've got for you, but we wanted to break them a part. First think about the voting, next we're going to transition

and think about the decision categories.

And then finally, we had had it on your agenda as the algorithm, but I actually hope you would indulge me a little and we could broaden it to think about what kind of information MAP needs generally.

We started to have some conversation about the feedback loops, what type of analytics you like about what happened the prior year, what measures are currently in the program, how what CMS did track with MAPs recommendation, as well as any input you have on how we could change the algorithm to make sure that we're bringing the workgroups the information you all think they need to have.

Also, if you've got any thoughts on our measure selection criteria. We haven't really changed those since year one so anything to freshen that up.

But just to keep things manageable.

If we could start maybe talking about the voting process and what you would think about some of

the changes on the prior side, are there
additional changes you suggest, your thoughts on
a structured voting process rather than a more
loose consensus driven process.

And then finally, this is something our Co-Chairs asked us to put on the discussion items, started to come up yesterday, is, how should we define consensus. It's currently greater than 60 percent, and that's something we did to align across NQF, but we want thoughts on, is that right benchmark, what does the Coordinating Committee think?

CO-CHAIR PINCUS: I just want to clarify a few things. One is that, after this discussion we're going to have a discussion about the voting categories. So this is the voting process focus.

And secondly, with regard to your first question, how are those suggestions that you proposed different from what we involved to over the course of yesterday?

MS. O'ROURKE: So, that's a good

point. Some of them are things we've actually implemented with the Coordinating Committee that came out of the workgroup discussions this year that we heard people wanted, the workgroups wanted that vote on the consent calendar.

They did want some more input -
CO-CHAIR PINCUS: Maybe I want to go
back to the next slide.

MS. O'ROURKE: So if you want to go back, yes. So, some of these we have started. I think we'd still like some guidance on what you all see as a good role for this Staff and how you'd like us to use that preliminary analysis that we put together.

Right now it's really for the workgroups information to inform their conversations. If you want a larger role for a chance to question the Staff on how we got there, what you would like us to do as far as theming. And then obviously the whole bucket of IT technical issues we'd love some input on.

CO-CHAIR PINCUS: And, Marissa, do you

want to hold up Robert's Rules of Order? 1 2 (Laughter) CO-CHAIR KAHN: 3 Amy. 4 MEMBER MULLINS: So, I know that, you 5 know, when I give lectures a lot there is 6 technology built into the lecture that you can 7 actually ask questions and people can answer 8 their phone and it registers on the screen and 9 it's really easy. So, there's an app for that. 10 (Laughter) 11 That can be so much MEMBER MULLINS: 12 easier than the point and click clicker. 13 think that that's an easy fix. Just download the 14 I don't know what it's called but everybody app. 15 has it. 16 (Laughter) 17 MEMBER MULLINS: So I think that can 18 be an easy thing to do. And then I think it's 19 interesting, this question about should we ask 20 the workgroups to come to a decision or should 21 they come to us with a split decision.

I think some of the things we talked

about yesterday would of almost been easier to discuss if we had known how split the decision in the workgroup was to begin with.

So I think maybe coming to us with a split decision wouldn't be bad. Because knowing how much controversy the workgroup had may have helped frame our discussion instead of thinking, well, they said this, they must have really believed that it was okay or it was not.

Knowing that they struggled with it and they couldn't come to a decision may have helped us not just flounder in our discussion for so long thinking, well, why did they think that we should do revise and resubmit, or whatever it was, if they just come to us with no decision at all then we could have started from there and then work to a better conclusion. Just my opinion.

CO-CHAIR KAHN: It seems to me that we need, and one of the keys to conditional I mean is this endorsement. And so we really need, I think, information about pathway on endorsement

when we're talking about measures that have not been endorsed or measures that started through the process and then got sent back.

I mean, I just was always struck by every time this issue came up there was a vagueness about where it was in the process and what the implications of that conditional work really was. And so I think we need a little bit more information about that up front. And I think it would save us some time in our discussions frankly.

CO-CHAIR PINCUS: Can I just add to your point about endorsement?

I mean, we'll talk more about this under the categories discussion, but it's also unclear how many of us around the table have had the experience or direct knowledge of what the endorsement process is. In terms of participating in it and sort of understanding both the process and the timing and the criteria.

CO-CHAIR KAHN: Bruce.

CO-CHAIR PINCUS: So therefore we

should probably have some more orientation about that.

CO-CHAIR KAHN: I think that actually we could do that and then have everything systematized so that on each of the measures where that's an issue, we'd sort of like see the timeline. Where it falls in the timeline. Okay, Bruce.

who have sat on committees and standing committees before, we're used to the vote tallies that you see coming out of those committees. You know, eight high, five medium. And so that's probably built into our mind set of expecting a vote where we actually formally vote and see what the numbers are.

I do think that, again, if you've sat on committees and done the endorsement process before, I think you also realize that it's not a linear or a straightforward and a given. And that's why I repeatedly raised my concern that I think yesterday a preponderance of the measures

that we move forward under conditional have not been NQF approved. A preponderance of them.

And so for someone who has sat on the endorsement committee process before, I'm not sure that those are going to go through unrevised or they're going to go through at all. And I feel like we are kind of advocating a responsibility, in a sense, by just saying, if someone else says this is okay, we'll say it's okay.

And again, I would just emphasize that even for measures that have entered that endorsement process and are specified at some level in that endorsement process, we're not sure if there will be a request to revise those specifications during that process.

And again, for a preponderance of our decisions that just, I feel like I'm not expressing this very eloquently but it feels like we've missed a beat somehow there.

On quorum, you only have to establish a quorum for the meeting and then that's it, your

quorum is established and then you just adhere to your voting process. So, the NQF pending, endorsement pending issue was what caused me the most the concern.

CO-CHAIR KAHN: Bruce, could you say that again, I didn't hear it very well. The last point.

MEMBER HALL: Us moving forward on a preponderance of measures that are pending endorsement was what caused me the most concern.

CO-CHAIR KAHN: Leah.

MEMBER BINDER: I was actually interested in having an opportunity for us to say, when we wanted something to have a high priority endorsement or a high priority on the calendar for endorsement, I really think this group should not be replicating the endorsement process, I think that we should be relying on it.

But I do think that there were some measures that came up that were, to my mind, just urgent issues for which there is a major gap in measures. Example being ambulatory surgical

center, whatever, readmissions to the hospital, revisits to the hospital. To me that's an urgent issue.

Because there is virtually, there is no data right now. There is just virtually no information on safety and quality in ambulatory surgical centers, and this is the kind of data CMS can get that others cannot.

So, anyway, that, for me, was a very high priority. I would have liked to be able to say, well, let's say that we want that one to be on the top of the calendar for endorsements. So, I would like that opportunity.

And the second issue I have though, is just a general issue, is perhaps NQF could have a parliamentarian or something. I feel like the chairs are put in some very difficult spots in many cases where they have to sort of referee and say, well, this is the rule and that's it and we're, you know, it sounds like we're kind of making it up as we go along, which I don't think is all that useful.

And especially when there is contentious issues that can be just very awkward, and I don't think it's fair to the chairs. So I would say, if you could get somebody who can say, well, this is it.

I mean, I've been to plenty of committees where you just have a parliamentary or you have somebody who is the rule, this is the referee and he says here's how it's going to go. I think that would really help to move things along and I think make us all a little bit more comfortable about knowing where the lines are drawn in terms of our arguments.

CO-CHAIR KAHN: Okay, Cliff.

MEMBER GIFFORD: So, I'd say in general I think we put too much emphasis on this voting process because it's not really binding on CMS in any way and it's more the recommendation. But I do think having different -- but I don't think we should throw the baby out with the bathwater.

And I do think having different

categories in voting does help with that. But it's really the recommendations that sort of go along with it.

Building a little bit on what Bruce has says, I see too often, both in the MAP and the workgroups and even in the endorsement process, difference to the timelines that CMS is put under, put under themselves, put under the process or put under by Congress, in the voting process.

I don't think that should be a factor in our voting. We are charged with saying whether it's ready for rulemaking. If it's not ready for rulemaking but CMS has to issue it next month in a rule to comply with Congress, then CMS will issue it next month in a rule and they will say, we are doing it against MAPs purpose.

But I don't think we should say, oh well, you have to do it. And Congress says you have to do it, we don't really like the measure, go ahead and do it.

And I've seen that happen a number of

times over the years. Both in the endorsement side and the other side. So I would agree with you, Bruce, on this side of how that.

And there is going to be measures that are going to be put forth that are not NQF endorsed and I think we should comment on that.

But, we spend too much time on the measure specifications.

Both in the workgroup, the coming up to us. I usually sit in and listen on a couple of them and they just get way down.

And even in our own group we use the word endorsement when we're talking about this.

And we're not endorsing these measures. There is another process for that.

So to that leads me to, I think, Erin, your point. I would love to see you guys more involved and more emphasis on some of the criteria that are about the role and rulemaking.

Because you've really thought them through.

And there is no one measure that's going to mean all the things. And so I think we

have that.

So otherwise we do sort of, I think

Amy you started to say, well, okay, I actually
think the voting would be better if we had the,
whatever categories we had where we just, down
those apps, we just pick what they are.

And then you, CMS sees what the spread is and they'll say whether they're, you know, what it is. Because it's really about feedback to the secretary for their use.

And I was struck by the conversation yesterday, I think in the afternoon, I was coming in and out when I had the calls about, well, if we vote it, will they have to bring it back or anything. Once it goes, once they put it on the list, it doesn't matter what we do we're advisory.

I mean, Kate and CMS and the team really are good. I don't mean to minimize it but we just need to realize what the process is. We could vote no and bring it back and they don't have to bring it back.

We can vote yes on a QRP program and they could use it in the VBP program in the future. Once it's on the MUC list it's free to use sort of whenever they want.

Now, Kate I think has been really good about that process. So I think, to me I think we need to think about how that voting process is.

And so the 60 percent and everything that helps provide the meaningful feedback to CMS. And then they have to justify why they are going to do something different than that.

And I think the guidance needs to be, we need to think more about how we give the guidance around the recommendations. Because I don't think it's sufficient with just the categories and then hear all the comments.

Because, frankly, there is some wacky comments out there. Including some of my own.

(Laughter)

MEMBER GIFFORD: No. So I don't think that there is, there is a binding nature to how they have to respond to all of these in the

process. And I don't think that's a good use of CMS's time and everyone else's time on that.

And so, just like you are saying on how to prioritize, I don't think we should get into a voting of it all, I don't think we should be writing the comments because we would never get done. That would be like a five week meeting here.

But I think there's got to be a better way to address those comments.

CO-CHAIR KAHN: I think this point about the voting is this dilemma. I guess, if you just, let's say whatever categories we have, if we just had all the categories, you're not ever going to get the 60 unless it's something that's just pretty clear cut. So, it's going to be confusing.

On the other hand, if you went to this app based voting, I mean, it would be pretty simple, you could do it. Part of the problem is we don't have a sophisticated way of doing voting so it's sort of a mess if we try to do anything

other than be linear.

MEMBER GIFFORD: The app program, you have four categories. We all pick which one it is and we see the distribution here, we can then do a little a modify Delphi and decide whether we're going to talk about it.

But then it gives some -- and frankly, the top three right now, I think the four we have are pretty reasonable. The top three basically say to CMS, go forward. The bottom one says no. And then they really have to justify if they wrote forward why it is.

And it's really, the other two just sort of give, the other two, the top one is go and don't bother, the bottom is, you got to really figure it out. The middle two are, you guys got to do something more and they have to justify it.

Which gets into a monitor discussion.

I think you and Harold were talking about how do
we monitor and track that and I think we should
be, I think that is a bigger role that we should

take on now that we've gotten further down the path in the measures that are out there.

CO-CHAIR KAHN: Carl.

MEMBER SIRIO: So, I'm going to give you a potpourri of comments that I think touch on a lot of the questions you asked. First of all, I would strongly urge us not to get too rule bound with the Robert's thing, right.

I mean, you guys aren't, whoever is sitting in those chairs are not experts in programs or procedure. And I've come to learn through some governance process with another organization, actually a misconception I think I had but many people probably have is that you don't necessarily need to be bound by it, right. I mean, you use rules of order to help you with a process not become hidebound to them.

So, again, I think that the more fluid the conversation could be the better. And it's really up to good chairs to manage the meeting not throw in some other person to adjudicate fights. I could not support that notion.

I want to come back to actually a comment Kate made yesterday, as I was getting a little prickly, and that was that we needed votes and we needed counts.

And I was struck by the comment that was made this morning that said that we started this process without that. So I think that even that we need some clarification on what CMS really needs to in fact feel comfortable that what we're doing is a true recommendation as opposed to kind of just a bunch of smart people talking about it.

Because if we've done it differently, it suggests to me that it's not clear as to what's required, what's needed or what would be helpful.

I am particularly struck by the number of times that we heard yesterday that the Committee has overrode the Staff recommendation.

Look, the Staff puts a lot of time and a lot of thought into and frankly our general experts, in terms of living with this material every day, and

I think that we need to have a better process of understanding.

And I think it's reflective in some of the comments, if a decision was changed with respect to recommendation, how, why and what was the discontent with the Staff recommendation so that we can actually be the arbitrators of that.

Which leads to a sense that we don't need to, the Committee shouldn't be forced to come up with a decision. If they can't come up with a decision, that's our job. I mean, we're advisory and they're advisory to us.

I think that, my sense is that the committees need to, to the point about not revalidating endorsed measures or not need to think about the appropriateness of the measure on balance with criteria and reliability issues so that in fact we get a better sense of a more robust and broader conversation.

I'm ambivalent on the issue of public versus electronic voting. I'm not sure how we evolved to a point we're afraid to see each other

raise our hands. But that, to me, implies a level of trust that needs to be questioned as a fundamental point of the group.

If we have disagreements, that's okay.

I mean, that's what the process is for. We shouldn't be ashamed of them we should be proud of the fact that we can celebrate our differences and move on.

I'm concerned that there's inconsistency based on the conversations we had yesterday and how the workgroups are approaching their work and/or their decision making.

So, again, I'm not sure what the right place is, whether it's with the chairs of the committees or with the staff in terms of kind of giving a little speech ahead of time. But I think we, as a MAP, need to have some sense that the work that they're doing on behalf of us or as the prelude to us, is actually this consistent, internally consistent among them, so that we have some sense that we're getting kind of the same materials to look at with the same kind of

thought process.

We're going to talk about the categories because I've got some significant issues with respect to how we seem to be confused by them yesterday.

And lastly, again, it seems to me that the notion of a consensus at 60 percent is somewhat historical. You know, we spent a lot of time yammering about quorums and whatnot yesterday to the point, once you got a quorum, you got a quorum.

But to the degree that this is not decision making this is advisory, it seems to me if you got 51 percent, or 50 plus one you win.

And that's the recommendation.

I'm not sure that 60 percent is any more logical than 67 or 75 in terms of the super majority, so I would suggest that we look strongly at actually just saying, if you've got a majority, the thing moves forward.

CO-CHAIR KAHN: Marissa.

MEMBER SCHLAIFER: Okay. And some of

comments, I was asked to look at this as a lead discussant ahead of time, so some of my comments have kind of been trumped by some of the discussion yesterday and today.

But first, I think what's most important is that, and I think we've evolved a lot from year one in that knowing what we're voting on. And so I just want to, once again, thank Staff for the background information in making sure that we're very well guided.

And so I think that's always been a little confusion on what are we voting on right now, what really does this vote mean. So that's good.

My thought was, one of the things is,

I see this, and I think it's not necessarily that

the way I see it is correct but it's worthy of

discussion from the wider group, of what we're

doing is endorsing a workgroup decision.

And I think it did me some good this year, something I've said I was going to do every year and had never done until required to, but

sit in on all of the workgroup discussions and listen to those discussions.

And while I don't think all of us have three days to put into that every year, I think everyone on the Coordinating Committee really should, at least one time, listen to the workgroup discussions. Because it's given me a lot of respect for the workgroup.

You know, maybe they're not always right, but it really has made me think about that when we're voting we're voting, in my mind, to endorse or not endorse the workgroup discussion.

One of the things that came up in one of the workgroups, and I think it was the hospital workgroup but I'm not, they all started running together, was that horrible up and down vote that I'm sure Staff has nightmares about still. Where they couldn't come to one.

They couldn't reach 60 percent on one and went through, I think, all four before they finally got to something. But there was a lot of discussion right then about, if we can't decide

should we just go with the Staff recommendation.

Which I think in some ways has a lot of merit.

As Carl mentioned, the Staff has really thought this through. What came out of that discussion was someone else on the workgroup saying, if Congress and CMS wanted Staff to be making decisions, they wouldn't have convened all of us around the table and said that these are the experts that should be making the decision and Staff as a recommendation.

So, whichever way we go I think that's something that we really need to think about, where we've said, let's put more weight on the Staff recommendation. I think we need to know if that's CMS, if that jives with what CMS is thinking or CMS really wants to see that it's the people around the tables decision.

I think, and I don't know that I know the answer but I think it's something we really need to think about. One of the things, and Carl and I kind of, the whole move from raising our hand to anonymous voting through the clicker, I

think we made the move because of technology and thinking that the technology was easier.

Without that real discussion, at least to my knowledge, about it being anonymous voting, I don't care whether we vote anonymously or not, but I think that's a significant change that either hopefully Staff and CMS had discussions about that or we need to have discussions about that to make sure it's not just a technology change it is an anonymous decision.

And then back to the 50 percent. If we are truly, something was said earlier in this discussion, I just want to make sure that we've captured it because I hadn't thought about it, someone said at 60 percent to overturn a workgroup discussion.

And if it's just a 60 percent vote,
but 60 percent is what NQF does, or we could go
to 50 percent, or are we saying that because
we're overturning a decision then we need more
than a, we need a steeper majority, if we're
supporting a decision we only need a majority. I

1	think we need to make sure that we hammer that
2	out.
3	CO-CHAIR KAHN: Let me, I think that
4	the basic operating procedure here is there has
5	to be, and that's why Bill brought it up
6	yesterday, a consensus. So part of the problem
7	is that this consensus thing is a little
8	nebulous.
9	And if you listened to Bill yesterday,
10	and I wish Bill was here today, he sort of said -
11	-
12	MEMBER KRAMER: I'm actually on the
13	line.
14	CO-CHAIR KAHN: at key. The key
15	was getting to the consensus. But the trouble
16	is, but, Bill, you could speak for yourself, if
17	you're able.
18	MEMBER KRAMER: Sure. Thanks for
19	letting me join in. Sorry, everybody, that I am
20	not able to be there in person today. I came
21	down with a bad bug when I woke up this morning.
22	First, my understanding is that

Measures Applications Partnership is supposed to be a consensus organization similar to the National Quality Forum.

It's a multi-stakeholder group in which stakeholders appeal from all different parts of the healthcare industry and the healthcare sectors expect to come together and work out a consensus.

No one said this could be easy, but I think it was felt that that was the appropriate way for decisions to made. Consensus, we need to be very sharp and clear what our definition is and I think there are a number of working definitions around there that we could use.

What happened, expanding a little bit on the discussion yesterday that I brought this up in the morning, this issue, the NQF Forum wrestled with this issue several years ago, and Chip and maybe some others may have been a part of that conversation, we formed an ad hoc workgroup to look at the consensus development process for decisionmaking at NQF, not just at

the board, but at CSAC and various committees as well, with the understanding that the goal was to find a way to reach consensus.

We struggled, frankly, and, you know, as part of that workgroup, I feel that, I'll say our work product was, though I was 100 percent satisfied with, but it got us past some sticking points.

One of the mechanisms that was in there to help move the process along was to say if there seemed to be, a consensus was not clearly emerging from some group discussion that there would be a pause and then a straw vote taken and if it was more than 60 percent then the group would say like, well, you know, it does seem like there is a pretty strong majority here, let's hear from the folks who are still, you know, opposed to the direction we're headed here, the minority, let's see if we can convince you, modify the motion, or figure out a solution here that could get a broader consensus.

Not that this would be your first

choice of the way we should go, but it would be acceptable to you and you would support it, or if it fell below 60 percent then you say, oh, it looks like we've got some pretty serious conflicts here, we clearly don't have consensus, this needs a lot more work, and then it could go either direction.

But, again, the goal is to get, give everybody a chance to hear what the issue is and be heard on their point of view, listen to each other, try to convince each other but also being open to being convinced, and in the end find a solution that works good enough.

Again, it might not be the first preference or first choice for everyone, but it would be acceptable to the folks in the room, and people would also come into this knowing that they would not exercise their veto power to block consensus except in extreme circumstances.

So that -- What unfortunately happened after that was despite a verbal commitment to a consensus things kind of devolved, and I think

particularly we have seen this here in the MAP process over the last couple of years, in which the search for consensus was kind of not emphasized.

Instead we started using the 60 percent guidance as an indicator of whether consensus existed as the final decision making process and then we came up with all these fairly rigid voting rules and, you know, what's the denominator and what's the quorum and all that kind of stuff. It was never intended to be used that way.

Now MAP could decide to use that if they wanted to. I, frankly, it would be hard to get me to participate in a consensus decision around that, but I think in the spirit of formation of the Measures Application

Partnership, in the spirit of NQF, and I think in the spirit of most people in the room that we are trying to, we have a common goal here.

We do not have necessarily a bipartisan or -- I'm sorry. We do not have zero

some kind of interests. This is unlike another institution up on Capitol Hill.

Here we have the interest of trying to improve quality and affordability, appropriateness, the care, and the patient experience, and so on, and make it work for all stakeholders, and I think that in that spirit that is possible and preferable to try to achieve consensus.

It is harder work. It takes training by the staff, it takes training and hard work by a leadership large groups or small groups, and it takes a training of the participants and a strong commitment to make it work.

And it won't always work but I think in most cases it's preferable and I would argue when it is done well it can be more efficient than voting.

In the end more people will support the direction that's been achieved because people have had a chance to have their say, listen to others, be influenced by others, and more likely

to have a solid front within our organization, or our committee, and to the outer world.

So those are my thoughts and I hope those are helpful in the session today.

CO-CHAIR KAHN: Bill, they are,
although they are troubling to me because clearly
when you describe that we have taken it doesn't
comply with what you were trying to accomplish on
the one hand.

On the other hand I think that we are balancing a lot of things and we are also, this each stakeholder, and we can call it interest, but we could also call it, I mean there are different world views, and particularly now that we are dealing with payment programs it's one thing, you know, not to just be dealing with professionalism, but we're dealing with payment programs, boy, the notion of -- I mean I guess the question, if you look back at yesterday on the decisions that were made based on votes in my mind I don't think we got to a consensus.

I think that one side won and one side

lost and I don't know how many hours we would have had to spend, I don't think it would have been physically possible.

I think everybody agrees with the process there, you know, and you paid your money and you take your chances, but I guess I am little concerned.

I don't know, I guess I'm just not sure, we have the time and it's necessarily realistic frankly to be thinking about consensus versus, you know, sort of general view.

I don't know, let's go around the table. Marissa?

MEMBER SCHLAIFER: I think, and I am sure even the people who did not get their way -- I would not want to portray it as one side won and one side lost only because I think one thing we did see yesterday, whether people were excited about it or not, but when we took a vote and it failed to reach 60 percent, when we took the vote the next time to go to either the higher or lower pretty much more or less we got to 90 percent of

the people changing their vote and going along with the group I think to show that this was a successful consensus process.

So I think where people lost or didn't get their way they still indicated that, okay, I want to go along, show we have a consensus, and let this move forward.

It didn't happen every time but it did happen every vote I remember yesterday, and that was something that maybe I was a little surprised to see, but I think we need to be careful not to say it was won or lost.

CO-CHAIR KAHN: I guess, and I think we need to be careful with our words, maybe I shouldn't have said won or lost on the one hand.

On the other hand I think people buy in to the process.

MEMBER SCHLAIFER: Yes.

CO-CHAIR KAHN: Saying people buy in to the process is not saying they have a consensus, because I think when you have the votes at the end I think if you went back and

looked at the people they weren't necessarily, I
mean that were --

MEMBER SCHLAIFER: If we measured how excited they were.

CO-CHAIR KAHN: Yes. I mean, to me, consensus is actually, as Bill is defining it, is spending the time to bringing everybody to yes, and I guess what I saw yesterday was, you know, on the ones that were controversial disagreements and then at the end acquiescence to the process.

That's not bringing everybody to yes in my view. Now that is my perception and I am interested in other people's view, but I would say, and I don't want to name names, but I could name a couple of names that I think would probably, you know, voted yes, I mean voted at the end but they, I don't think that they consider it a consensus, I think they just acquiesced. I don't know.

CO-CHAIR PINCUS: Just to apply some broader technology, we could attach brain scans to everybody and determine whether they were

excited or not, but the problem is it is hard to differentiate between excitement and anger.

CO-CHAIR KAHN: Yes. Carl?

MEMBER SIRIO: So, Chip, I think you put your finger on the difference between, and we were having a little sidebar here, between consensus and acquiescence, right.

I mean, I think you are right. I mean to the degree that there were positions that were supported or not, I won't say won or lost, but to the degree that we are time limited, we are process driven, and we can't spend hours adjudicating all of these, different process, different methods, we might get to a consensus.

But that is why I would advocate just go with a 51 percent because at the end of the day it is not an consensus in terms of -- It's consensus with a process, not with the outcome.

So I would submit that 60 percent is as random a number as any other number north of 50 percent.

Washington DC

CO-CHAIR KAHN: Giff?

MEMBER GIFFORD: If I was Kate I would be chuckling over there at this discussion because I mean not to make our vote irrelevant but in many ways the vote doesn't matter.

I mean I read the rules on everything that comes out of this and, you know, whether -The vote categories aren't what drives this.

What drives it is their need to meet the programs in the rules and whether it is statutory or not and what they have to go forward for.

And unless we just say no it's -- I have yet to see something that hasn't come through not show up in a rule and it gets imposed.

And so there is just way too much over emphasis on what this voting is and it's more about the guidance and I think -- Now with that said CMS, and Kate has been very good about trying to honor these and you will see stuff show up in the rules as to why they are trying to explain it all the time, but it's not -- Well, I'm not Kate, I am chuckling at the conversation

because I mean, yes, I really don't care what the vote is.

I mean to me it is what is the feedback and the wealth of the discussion. I have heard Kate say it multiple times over the years, it is the discussion that they find really valuable.

The question is how do we capture that discussion and get it in there and I think the vote is, this is why I say don't throw the baby out with the bath water, the vote does help with that, but we've got to think about it that way and that I think will help easier get to the consensus sort of view out there.

But to try to get to 60 percent I think is not, it's missing the point of what our role is and how CMS uses the results from that and I think when we start looking at the actual data of how they use this you'll see it.

I can't recall a single time that anyone has ever said bring a measure back that it's actually come back to the MAP. I have seen

them say it should get NQF endorsement, I have seen it shown up as an NQF endorsement later on, but I have never seen a measure come back to us, ever.

Kate is about to prove me wrong, but they are scratching their heads, too. I'm not saying that we have to have had measures come back, I am just saying I haven't seen them come back. We act like they are something --

MEMBER GOODRICH: You're good if you can remember every measure --

(Simultaneous speaking)

MEMBER GIFFORD: I'm just saying we're acting, overemphasizing something that doesn't matter as much and we are missing our point on there and I think that is the message I am trying to deliver to you.

And so to me whether it's 59 percent, 60 percent, 51 percent, I don't really care on it. I think our role is to, if you look at the statute, is to give feedback to CMS on what to do and I think Kate and CMS takes that very

seriously and has done a really good job with that, but I think that, you know, we spend too much time hung up on it.

Did you remember, Kate, when we brought back --

(Simultaneous speaking)

MEMBER GOODRICH: The only thing I am trying remember there was a measure that we brought back as part of the feedback loop pilot, but it didn't go back on the muck list but it came back for discussion. I think that's what I was just trying to clarify with Pierre.

And we have talked about trying to do more of that so that there can be feedback to the MAP about decisions we have made, et cetera, that's all.

CO-CHAIR KAHN: Leah?

MEMBER BINDER: Well I think one other role, to be very blunt, is political cover. I mean I think so. When we have a strong -- I think really our biggest role is going to come when we have strong feelings one way or the other

about a particular measure.

So if this group really feels strongly that a measure should not be in rulemaking and then CMS puts it in rulemaking, they have, you know, it's very difficult for them to politically put it into the final rule because it will immediately come back, well, you know, the MAP strongly did not, you know, approve of this.

again, but I do think it's on the outliers when we have strong feeling one way or another. So that gets to the point of the 60 percent being probably a good metric to give CMS a real indicator that this is a strong feeling, or strong enough feeling on the MAP that it's relevant for them to take into consideration from a political point of view as well as substantive point of view. So that's why I would support that.

I just wanted to make one clarifying remark about what I said earlier about the role of the Chair as sort of a parliamentarian or

somebody who is kind of a referee.

I don't mean that the Chair should not actually oversee the discussion and controversy, those things are all valuable and I think healthy and important for the Chair to oversee and for the conversation to take place.

So I in no way mean to say the Chair should have no role in that by no means, but I do think that one of the things we get caught up in is like the procedural voting issue, you know, do we take a vote on that or do we, you know, do we need a second, or when does the discussion take place, when does a motion happen.

Those kinds of things I think can really trip us up and that's when, and it just doesn't, I don't think it's healthy for the group or the conversation to have the Chair kind of caught up in those kinds of procedural issues where I do see a role for, a staff role to really have somebody there that can just clarify it, to move on, and not get us caught up in things that take on a life of their own and derail the

conversation in a way that is not healthy.

MEMBER QASEEM: Hi, Chip and Harold, this is Amir. Can I just say a couple of things?

CO-CHAIR KAHN: Yes, we're going around the table here, but why don't you speak.

MEMBER QASEEM: Sure. So the issues with Measures I think what I am hearing is that it is an accountability. They are being used for accountability, it's a high stakes environment, right, and my worry is that yesterday what happened and many measures that came through that were not NQF endorsed.

So either we believe in NQF's process or we don't believe in NQF's process and if we believe in NQF's process the way to simplify things if a measure hasn't been endorsed by NQF we need to come up with a set of rules that then need to be just exceptional circumstances that we really need to look at this measure.

Yes, there are many urgent emerging issues, there are tons of them in clinical care, but will it be okay if we wait another year since

we have waited for 20 years to address that issue and let a measure go through the NQF process, at least we start feeling comfortable that some smart people around the table have spent enough time looking at the measure making sure it is a good measure or a not so good measure.

Many times yesterday I think, at least
I felt like is we didn't have enough chance to
dig deep into whether this is a good enough
measure and if it can be used for accountability
purposes or it's just for quality improvement,
and those two are different issues, right.

NQF endorsement what is happening in clinical, you all know, Carl mentioned yesterday about the who attribution issue.

We are endorsing measures at the health plan level but they are being applied at clinician levels without any evidence whether it should be or if it works out or not.

We are all aware of those issues. I think there is some fundamental process issues that till they get resolved, and I have seen

deviation over the years.

I think when MAP started out we used to rely more on NQF endorsement and now we are slowly moving completely away from NQF endorsements.

And I think to be honest we don't have enough time at MAP meetings to go through reviewing and figuring it out whether a measure is a good enough measure one just general comment I think we need to start thinking about it.

Otherwise, I am not really sure what are we doing. As I said yesterday there is a lot of smart people, we are spending a lot of time, but I had rubber-stamping things as well.

The second thing is I think the voting, the votes do matter, but I am not a big fan of the 60 percent/50 percent number. I think what CMS will benefit more is to see the spread and there are various ways to convey that and you can use RAND process, you can do whatever process, but if they are, they are a certain percent -- Like use the example of that A1C

measure.

If the spread is 64 versus 36 percent we should automatically categorize a measure like that as a condition measure. What if the measure is getting 90 percent, then it should be fully endorsed.

I mean there are ways to address that.

A lot of people have done research on this and
more scientific, what does this random 60

percent, at least I feel like that we are using
to overturn the committee's decision, which I do
value a lot as well, and I think that is exactly
the spread that we should also be seeing.

But what happened in the clinician subgroup? What was the spread or help us make that decision. But I think some of these process issues, is MAP's coordinating committee's function is to look into the evidence and look into the quality of the measure or are we going to rely on someone else who has spent enough time to go about doing that and that brings me to the discussion.

I mean I value NQF's staff
recommendation a lot. They have very smart
people and I think we need to take those into
consideration as well as the NQF endorsement
process and I feel like that CMS might have a
deadline but we are gathered around that table
with a different function that we wholeheartedly
believe the measure is a good measure.

And if you believe it is not a good measure, to be frank, I don't care what happens with what Congress is saying and what -- My obligation is to my patients and that's what we should all care about clinicians were sitting around the table, as well as the patients who should worry about what kind of care they are getting.

(Simultaneous speaking)

MEMBER GIFFORD: Kate, I don't want to put you on the spot, is feedback of whether it's an absolute yes or no a black and white vote or the distribution, because you have heard, I think there has several that suggested, what is more

helpful for you guys?

MEMBER GOODRICH: Like I said before, it's really helpful to us, and Pierre should also weigh in, I mean the yes or no in the distribution in terms of numbers does matter less than what the actual comments were and by which kinds of stakeholders.

That's much more useful information to us. I don't quite know how to resolve this dilemma, but I will say at the end of the day that's what we want to be able to use when we are justifying why we are proposing a measure, or internally for ourselves to make a decision about when not to propose a measure as well.

that we do play a role that reinforces in the endorsement process, because if MAP didn't exist and you just had the endorsement process, not that CMS wouldn't worry about the endorsement process, but it wouldn't, there wouldn't be any transparency -- I mean you could look at a measure that was in a reg and see whether it was

endorsed or not on the NQF list, but it wouldn't have been part of a process and I think that's important.

To me that is important because there is a synergy there, I mean, you know, people are talking about it. If people weren't talking about it, if the MAP wasn't here, and I think that is your point, Giff, it really does make a, it probably does make a difference and without the endorsement process we don't have what Amir is talking about.

MEMBER GIFFORD: When we get into a later of discussion about the steps and the criteria for review we are evaluating the measures --

CO-CHAIR KAHN: Well speak into your mic just because the people on the phone.

MEMBER GIFFORD: Well when we get into the discussion later on this morning or this afternoon on the criteria for the review we are looking at the measures form a different vantage point than the endorsement process.

1 CO-CHAIR KAHN: Yes. 2 MEMBER GIFFORD: Endorsement is looking for liability, validity, and all the 3 other stuff. We're looking at is it ready for a 4 5 rulemaking and the rulemaking process. And if you look at the criteria that 6 7 are there, that's why I think the staff have gone 8 through it, it would be helpful to give us more 9 input on that because that's I think --10 CO-CHAIR KAHN: Right. 11 I would like to see MEMBER GIFFORD: 12 more of the discussion on that than on the 13 reliability, validity, or even the burden. Ι 14 mean the burden is important part of the rule, 15 but we don't talk about a lot of those other 16 things at all when we are talking about the 17 measures. 18 CO-CHAIR KAHN: Chris, did you want to 19 say something Chip, I think we might 20 MEMBER QUERAM: 21 be nearing that point in the meeting where

everything that can be said on this topic has

been said but not everybody has said it.

(Laughter)

CO-CHAIR KAHN: Yes, Chris.

MEMBER QUERAM: I find myself agreeing with comments you made probably 20 minutes ago in response to Bill's articulation, a very thoughtful articulation of the consensus process of the NQF, and I would just say that I think that approach to decision making is designed for a different type of structure than we have here.

It works well with workgroups, it
works well with the board. Groups that have a
frequency and a duration to their interaction
with one another where trust, not that trust
isn't important here, but where trust is critical
to ensure commitment to an outcome.

And I think we have to balance expediency with good process because we are here for a day. We consider a tremendous amount of work that has gone into the preparation for a voting process and there just isn't time to be able to follow faithfully the approach that Bill

so nicely articulated for us.

I worry about Kate's comment about relying more on people's contributions and comments than on the outcome of a voting process.

I have very little faith that there is a good recording, a good capture, of the points of view.

It is too subjective, it's too ephemeral. I just don't see the -- I mean I hear you, I accept it, and I think you are being genuine in your comment, but I, it's a thin reed for me to feel we should rely that CMS is going to take to heart the comments that they have heard from people.

So I do think some type of a voting procedure is important. Boards have requirements to protect changes to bylaws that necessitate super majorities or some higher level.

I don't agree with Carl's comment that it should be 50 percent plus one, I think that is too casual. I don't know that there is any magic with 60 percent. I would actually like to see a little higher number so that there is some

confidence that a strong plurality of people around the table agree with a particular point of view. So those are my comments.

## CO-CHAIR KAHN: Kate?

MEMBER GOODRICH: Yes, I think for us the good thing is for the majority, I think, of measures the MAP is pretty clear, there is a pretty good consensus on whether the measure should be support, conditional support, or do not support.

So we are talking about, in my mind, a minority of measures, or if I harken back to internally when we are trying to make decisions about whether or not to propose a measure that got one, you know, ruling, if you will, or another.

We talked a lot about, well, so what was the discussion, because I think for us when we are putting something out into a proposed rule for a measure where there is not consensus really.

I mean maybe it got voted in one

1	direction but we know from the conversations that
2	there wasn't consensus. What can helpful to
3	elicit helpful public comment in a proposed rule
4	is to be able to say here is measure $x$ , we want
5	to propose it for this reason, it went to the
6	MAP, it got, you know, conditional support, or
7	whatever, and, you know, some of the viewpoints
8	were this and some were that, you, public, tell
9	us what you think.
10	So it is in those situations which
11	thankfully are a minority of situations where I
12	do think that especially having that richer
13	detail would be helpful because then we can say
14	to the public here is what we have already heard,
15	tell us what you think, and that can then, I
16	think, engender more specific and more helpful
17	public comment for us.
18	CO-CHAIR KAHN: Sam, I was going to
19	hit Joe and then
20	MEMBER LIN: Well
21	CO-CHAIR KAHN: Well you may leave.

Yes, why don't you go.

PARTICIPANT: You haven't spoken for a day and a half.

(Laughter)

MEMBER LIN: Thank you. Well, I had written Leah, Giff, Carl, and Marissa's remarks. Following Chris's new rule for the group here, do not repeat what has been repeated, I will try to say something new.

I am thinking, you know, Giff had the comment about the discussions and whether it is important and I certainly buy in to that except I have to think back to yesterday where we spent time needing to count the numbers for two different programs because the numbers had to be there.

That's not following just the discussion, that's spending more time on numbers. So, you know, with English as a second language I always go to Wikipedia as my source of information.

So what is a consensus decision making process, and this may give us either help or more

headaches. Consensus decision making is a group, a group, decision making process in which group members develop and agree to support a decision in the best interest of the whole, a decision in the best interest of the whole.

Consensus may be defined professionally as an acceptable resolution, one that could be supported even if not the favorite of each individual.

So if what we are trying to do, and I am following Bill's lead at this point about consensus, is that we take our vote. We find the vote is six to four, whatever that is, fine.

So then we have to decide whether or not we are going to go into a consensus and whether or not the four can agree as part of the group, because it's this group thing, to be supportive.

They don't have to support, but they have to be supportive of the other six. We come up with a consensus of ten out of ten. Now I know all that is very simplified, but that's, if

we are going to do consensus or we shouldn't be talking about consensus, we ought to just be talking about rule of law or something.

CO-CHAIR KAHN: Joe?

MEMBER KRAMER: This is Bill, if I could jump in maybe with a process suggestion. I appreciate the last person who spoke. I couldn't quite recognize who it was, but it is consistent with my thinking.

CO-CHAIR KAHN: Sam.

MEMBER KRAMER: But listening to the conversation I think it's pretty clear, my observation is that we don't have consensus yet in the Coordinating Committee about what decision making process we should use.

It warrants further discussion. We actually don't have to resolve this until, before next year, the next cycle. This is in some sense the beginning of the next annual cycle.

I suggest that leadership of the

Coordinating Committee and maybe a small ad hoc

group work with the NQF staff and maybe others at

NQF maybe on the, you might want to bring in a board here that has been doing some work on this, to try to look at the range of options and see if that workgroup can come back with a consensus about the appropriate decisionmaking process.

It may be that it is the classic consensus, it may be the opposite voting, it may be a hybrid where we seek consensus and resorts are voting only as a last resort.

But the suggestion is that we work on this over the next six, nine months, so as we go into the next cycle we are more clear of how we are going to make decisions and do the preparatory work so that people understand what the process is going to be at the workgroup level as well as in the Coordinating Committee.

CO-CHAIR KAHN: Thanks. Let me go to Joe and then I'll come back to Sam.

MEMBER BAKER: Thank you. I guess, and I certainly -- and this my first meeting so I have more questions than answers and I agree with Chris that a lot has been discussed and I agree

with I think we have reached a consensus and agree with what Bill just said.

A few observations from the newbie as it were. One is I think the technology of voting certainly frustrated us yesterday and was part of the problem and people, you know, whether that frustration was real or just displaced frustration because they couldn't be frustrated with the measure itself or other processes making the voting easier by hand or by technology I certainly endorse. It should be as smooth of a process as it can be given the importance and the feelings involved.

I mean secondly I do think that there is this gap between endorsed and non-endorsed and that creates a lot of, you know, discomfort it seems for some folks and for others there is a feeling like we need to get this out and we need to get it out quickly and we trust the endorsement process.

It seems to me the cycles are misaligned and that's never maybe going to

resolved. So on some level we can't -- and it seems like this discussion has been had before, so we either have to say we had that discussion in 1962 and we are done, you know, yes, thank you new person, this is a rule, you can't talk about this anymore, you know, because I'm not going to, okay.

I don't really care, okay. Others care about this issue, maybe I don't, or I'm not going to care about it now because it doesn't look like it's going to change, and that's maybe, you know, there is not only a parliamentarian, as Leah has, you know, recommended, but there is also like, no, here is the rule keeper, you know, you can't discuss that part.

And then, three, I have a question, or, you know, I also saw this idea that I need to pull a measure in order to make a comment about it, right.

So all of a sudden right then I am like, everybody is like, oh, you know, it's been pulled, that's a thing, you know.

And then you've got to do a lot of explaining, as Carl did yesterday in some instances, that I am not pulling because I hate this or because I think it's all wrong or because I am a communist or, you know, I am a radical.

MEMBER SIRIO: But I am.

MEMBER BAKER: Well you are, okay.

But, you know, you're not doing it because you are a communist, you're doing it because you have other concerns.

You know, it creates this contentiousness and this confusion, right, around that. So maybe there could be more briefing up front about why some thing is being polled so we know up front here is the things that are being polled so the discussion can be had and here is the things being pulled because a nuclear option is being debated, you know, or because there is a -- and then we are prepared.

Like, okay, there is nine pulls, but six of them are really just so people can discuss it and three of them are, you know, really to

kind of drill down into some of these issues because, for me, I would certainly need that to get those three things in my brain as a consumer advocate and I think it would all focus our attention and calm us down on the other pulls, if you will.

So those are my observations and ideas.

CO-CHAIR KAHN: I think one of the solutions I mean here is we could -- I didn't like it yesterday, we don't need to use the word "pull." I mean we can just use identification of -- I mean we don't want to discuss every one on there because some we don't have to discuss.

So you need to find another word. You don't need to find it now that just talks about raising it. And the trouble is you don't know necessarily, some may be pulled, well, may be identified for discussion that could end up with a motion or it may not end up with a motion.

They can't necessarily predict that.

I mean in some cases, you know, Amir maybe had

his list that he wanted motions on, but that could have changed in the discussion, but I think we do need some nomenclature there that's a little bit more neutral.

So we are almost at that point, I'm going to let Harold close out, summarize, and then -- I'm going to pass the baton to him as chair.

CO-CHAIR PINCUS: So it sounds that there has been a lot of discussion about this and from my point of view I think it's hard to, number one it is hard to distinguish the process from the decision categories and I think we are going to come to that after the break.

But I think that there is a kind of back and forth between adjusting the process to the decision categories and the discussion categories to the process, and so we need to sort of think about them on both tracks.

But my inclination is very similar to Giff's in the sense that the main purpose of this is to give advice and whether the advice is

dichotomous advice, you know, yes, no, or whether the advice is more qualitative, or both.

And so the issue for me for process is what process will provide the most useful advice and to think through that and I think some of the suggestions that we made I think make a lot of sense in terms of, you know, maybe we should, you know, I think being more clear about, you know, what pull means and how to distinguish between clarifications and questions as compared to I really have a disagreement with the proposal that has been given to us by the workgroup, so to clarify that I think can help to do that.

I think achieving a consensus is not necessarily the purpose of this because we don't have the time or the, you know, the time or really the resources to go through a whole RAND Delphi process.

I do that a lot and it's a lot of work. And so it's really not a process to achieve a consensus but it's a process, again, to provide advice and I think we can talk about

what's the best one, but I think, you know, giving -- My own view is that I think coming to some kind of vote puts people to the test of actually having to collect their thoughts and decide how they want to go and then provide some justification for that.

And that process does provide more qualitative as well as kind of a more categorical response. So we are going to take a break, okay.

CO-CHAIR KAHN: Let me just say one more thing. This was my idea, this whole thing, and when the Hospital Quality Alliance had run its course, and Chris was on the Hospital Quality Alliance, and we needed something that was multistakeholder and met all the criteria and to me, and to sort of to bolster what Harold said, the one thing about this idea that has worked is that they are sitting here and they sat here through the whole process and they hear everything that we talk about, and so at the end of the day, you know, I don't want the perfect to be the enemy of the good, I think we do accomplish that.

Now whether or not we actually move them on all the things and sometimes, you know, we may not be, you know, maybe it's not possible, but I think we do accomplish that we have their ear, which I think we lost in the Hospital Quality Alliance for a whole set of reasons that were as much process as anything else.

CO-CHAIR PINCUS: So we're going to take a break. Let's get back together at ten after 11, okay.

(Whereupon, the above-entitled matter went off the record at 10:52 a.m. and resumed at 11:09 a.m.)

CO-CHAIR KAHN: So the categories that we have had as voting categories have kind of evolved over time and there is various interpretations, even as they have evolved.

Even as the categories have been stable there is different interpretations of what they mean has kind of evolved over time and, you know, I think some of the issues that we have dealt with have really came to light yesterday in

terms of thinking through and people being clear about what they were voting on.

Let me just sort of set this up just to give you sort my way of thinking about this.

I think in some ways we have been confusing several different issues as we deal with the categories.

Number one I think we have been confusing content and process in some ways, so that we have one category that clearly is a process category of conditional typically for some condition being met most typically in endorsement and then we have more of a content category of refinement and resubmission, which is more about, you know, particular issues that need to be dealt with, and I think there is some confusion about that.

And then even when we talk about some of the content issues we have some confusion about that also. Some of it has to do with issues around the nature of the measure itself, some content issues pertaining to the nature of

the measure, for example, the measure needing risk adjustment as being something that the people have put forth.

And in some cases it has to do with the nature of the measure as it applies to the program in which it's being placed that it hasn't been, it's been attested and endorsed at a health plan level but not at a provider level, and so we have some confusion about that.

And from my point of view I think
that, again, going back to our discussion that we
just had in terms of really what we really want
to get at is what is the best way to provide
clear advice is that we need to clarify these,
the distinctions, especially between the two
middle categories.

And it seems to me that we have two options. One is to simply go with three categories that we support, we don't support, and something needs to be fixed in the middle, or we make some clearer distinction between the two categories about conditionally support and revise

and resubmit and really clarify the distinctions between them in a much more effective way.

But ultimately what we want to come out of this is that for those, whether it's one or two middle categories, that we are more specific about what we expect.

So why don't I turn it over to the staff to go through sort of their initial thinking about that.

MS. O'ROURKE: Sure. If you could go to the next slide. So Harold went through these, but just to reorient everyone to our current decision categories.

Right now we have support for rulemaking, conditional support for rulemaking, refine and resubmit for rulemaking, and do not support for rulemaking.

And then in the second category, I know it's a little hard to read, it's basically just to our decision algorithm how everything maps to show that at least for the staff assessment when something doesn't hit one of the

assessments that we do where it would fall down in the algorithm, if you will, or how we got to the decisions that staff put forward for the workgroup's consideration.

So that's maybe not necessarily where we need to focus for this conversation. I think on the next one we could maybe talk about that algorithm and what information people need, but I think Harold already really laid out, if you go to the next slide, some of the concerns we have heard and at least some ideas we had of how we could potentially improve.

I think we wanted to get peoples
thoughts on is there value in preserving this
refine and resubmit category. I think to kind of
echo Harold's thoughts of I think the intent is
that it signifies there is a larger change
needed, do we want to keep that and clarify some
of the language, and I think certainly one of our
ideas is to change the name.

We were thinking something around support continued development or -- that's

obviously not anything we are wed to, but an idea is it just in the naming of this resubmit, is that the word that is causing problems, or have we just created differences without distinctions, if you will, and we should just go back to the three category system where there was support, conditional support, and do not support.

so I think we just really -- Again, nothing we need to decide on today, but we want to get your thoughts about what you think would work and what's the most valuable for you, do you like the four categories, three categories, so if we could just get everyone's ideas on the tables like we did for the voting we'll bring it back in the fall.

(Off-microphone comments)

MEMBER GOODRICH: So I have said this before, refine and resubmit, and I will say even the former, support continued development, because we had on like that at one point, I think, or encourage continued developed.

MS. O'ROURKE: It was a two-measure

pathway and then we collapsed the two pathways into the four.

MEMBER GOODRICH: Yes, right. I will say for us at CMS it was, kind of plagued us a little bit, and that was nobody's fault. We were trying it, right.

So one of the thoughts that I have around this in terms of just thinking about what would be helpful for us, too many categories gets just challenging I think for lots of reasons.

However I think there does need to be something that the MAP can say to us when -There is a significant, you know, right direction, but there is a significant concern that you really need to address.

Part of the way we have tried to address that is by not bringing measures forward or not very many anyway unless they are like a significant public health priority like the opioid measure from before that are too early in their development process.

So they really wouldn't be ready for

rulemaking next year anyway because they haven't even gone through testing or what have your, right, so hopefully there will be fewer of those kinds of things.

What I think could be helpful to us, and I think just as a general comment for the conditional support category anyway, is I think the refine and resubmit my personal view is we shouldn't have it.

However, because there is still a role for substantive comments that currently may be under refine and resubmit what I might suggest that MAP to consider is some more precision around the conditions in conditional support.

Because refine and resubmit or support continued development really has meant we support the direction of this, we think the concept is important, it fills the gap at whatever, whatever, but here is some problems.

So we always think of conditional support as being conditional on NQF endorsement because that is the biggest bucket of things that

it is conditional on, but there is no reason that it couldn't also be conditional and you got to test this at the clinician level, it's only at the health plan level now.

So until you have tested it at the clinician level and it passes muster, you know, that's our condition, or that's just one example, but that is an example of where a measure has gone to refine and resubmit in the past but could just as easily be a condition.

And I think that, from my perspective, and you should weigh in, Pierre, if you disagree or think differently or whatever, I think that could be a helpful way to resolve this a little bit more easily, at least when I think about what we talk about and think about internally as we are going through the MAP recommendations prior to rulemaking.

CO-CHAIR PINCUS: So it sounds like one of the things you are suggesting is rather than making the conditional one be a process recommendation to make it more of a content

## recommendation?

MEMBER GOODRICH: Yes.

MEMBER MULLINS: So I am going to take your suggestion one step further. I think that we should have two categories, support or do not support.

I think anything else we do gets really into the endorsement process. Either the measure is ready for rulemaking or the measure is not ready for rulemaking and I feel like that is what our charge here is, is is this measure to be ready in the rule or not.

I think that you could scale it after that. So if the measure is -- If you support it I think maybe what would be helpful for you guys would be could we support on a scale of one to ten and then you could some sort of feel from the group of how supportive we are of that, because really the support with conditions, we have talked about it for the last day and a half, it doesn't matter, they're going to put it in the rule or not.

What matters is the comments and the discussion around it. So either we don't support it or we do and whatever condition or category we put it in, unless it was a do not support, it's probably going to end up in a rule somewhere.

So we either support it or we don't and if we support it how supportive are we and what are our comments around that measure.

CO-CHAIR PINCUS: Okay. So I know people put their cards down, but, Giff?

MEMBER GIFFORD: I would just sort of add that I think the one thing that gets us hung up is when CMS brings a measure that has not been NQF endorsed and whether -- I don't think our role, it struggles because at that point how do we discuss whether it's ready for rulemaking if it's not been NQF endorsed, and some of us were offline should we just blanket it and say it's not ready for a rulemaking do that.

I think that's probably a little bit Draconian, but we almost want to have a default recommendation and I think that's almost where

you may want to have some sort of subcategory which is, yes, we understand it has to go forward with rulemaking, yes, I think we buy it, but you better get, you really need to get NQF endorsement and what Leah is saying is it's a high priority out there.

But I would agree with you, Amy. I think it's how do we communicate that, and I don't think this third category in the list is adequate communication there, a resubmit or anything else because then it gets into this whole thing.

It's just -- We need to have that rulemaking, or that category. And a better way of capturing our support and our endorsement, but you're right in the end it's really black and white.

## CO-CHAIR PINCUS: Carl?

MEMBER SIRIO: Yes, I actually am persuaded also that kind of an up or down is probably the wisest thing, but I think I need some affirmation of my institutional memory.

We got to this place if I recall because of the fact CMS was in fact considering materials that were not NQF endorsed so we kind of jerry-rigged this process to create some criteria.

So my sense is that going to an up or down it takes the onus off of us in terms of worrying about NQF endorsement or not because we can factor that in but this work around in my view was a construct of the fact that measures were coming forth that didn't have NQF endorsement.

It seems to me that we can have the up or down vote and then probably a conversation around measures that are worthy of continued development but I would be reluctant in terms of the comments that were made about support continued development because it's really support continued development as a code word for don't pass along, right.

So there is positive verbiage but it's really a negative vote. So I would just be

careful that whatever you come up with in terms of the next iteration, whether it's an up or down or a three category thing, that it's pretty clear unless you get an affirmative yes moving forward everything else is a no.

CO-CHAIR PINCUS: Marissa?

MEMBER SCHLAIFER: I'm going to leave any conversation about the conditional support to Sam, my other lead discussant, and I think he's going to make some comments there. I will just say I agree with him.

On refine and resubmit I think both some comments here and I agree what Giff just said about this being confusing but it was something that he said earlier that was going to cause me to disagree.

Earlier he said there is three levels of support and one do not support and I very much see refine and resubmit as saying this isn't ready for prime time therefore we don't support it, however we like the concept, we like where you are going, we want you to continue going in

that direction, but we don't support this going forward for rulemaking now.

And, you know, maybe we will in the future, maybe we won't in the future, we don't know for sure because we are not seeing enough. So whatever this turns out to be I would not use the word "support" in any way in that, you know, I think we talked about encourage.

Because when Giff said earlier there is three levels of support I think that showed the level of how we all interpret this differently and I think it needs to be really clear that we are saying we don't want you to go forward, we like what you are doing, but we don't want you to go forward, and I think that needs to be really clear assuming we continue with this.

CO-CHAIR PINCUS: Sam, did you have a comment?

MEMBER LIN: Agree, agree, agree.

Well, I agree with Amy's suggestion, but since we are still under the old rubric I will stick with that.

A point of ignorance, other than pending NQF endorsement what other conditional support have we labeled, like it has to rain tomorrow or it has to snow next week, I mean what the conditional other than NQF support?

Now I am not trying to be totally facetious because I am still thinking of Amir's concern yesterday relative to the clinical workgroup and his understanding at that time of the composite.

Before Marissa and I started attending some of these workgroups I would listen in on the webinar and I remember one time, and I think the numbers are accurate, it's been five, six years at least, where there was like some 28 people on a clinical workgroup and there was some 20 folks in attendance for that particular webinar.

And so there was a neurology issue and, you know, there was no neurologist on the thing so they voted for it anyway. That's not a criticism, that's simply how our workgroups cannot include every single specialty, otherwise

we would be and CMSS right now.

On the other hand, when NQF does their endorsement if it's a neurology issue the neurologist are hot into it, you know. If it's a rheumatology issue the rheumatologists are hot into it, so that the scientific evidence base should be pretty clean, pretty clear, and I think we probably, Amir would accept that.

What is our capacity then? I think our capacity is not the evidence base but the experience base, which is all this other context that we have to deal with.

We accept the evidence base because we are not the neurologist, but then we look at the other issues that are involved and recommendations to CMS that has to deal with the politics that maybe we can help on or hinder, depending on where we come out.

So I need some clarification on what other than NQF support, or endorsement, what conditional support would mean. I think the up/down, vote down, you know, again, I go with

Giff saying we are advisory. Again, HHS will do what it thinks is necessary.

I will go with what Leah said the last hour that where there is an emergency issue as in all cases in emergency issues we deal with them on a case-by-case basis as needed, as necessary, they already fall out of the normal run of things.

CO-CHAIR PINCUS: I mean I can imagine, and Kate or Pierre maybe wants to comment on this, I mean to even now we have had cases where there has been an already endorsed measure but it hasn't been endorsed in a way that would fit with the program. So that is one example where, you know.

There also may be reasons for which we want the endorsement committees and CMS and the measure developers to be prepared for, you know, critical issues that we think will come up in the endorsement process that we want to have addressed because of the nature in which CMS, the program in which CMS is proposing to use this.

So there may be different types of 1 2 comments that we want to make and explicitly call out that may be relevant to or, you know, the 3 4 endorsement process. If I can follow up. 5 MEMBER LIN: Thank you for the --6 7 (Simultaneous speaking) 8 CO-CHAIR PINCUS: These guys may have 9 some other ideas about that also. MEMBER LIN: Well, thank you for that 10 11 clarification. My concern then is if we have 12 those hangars on when we send it forward what if 13 the hangars on are not accepted, are we still 14 supporting it conditionally or do we say all bets 15 are off? 16 CO-CHAIR PINCUS: Well, I mean one 17 thought is to, you know, have this so-called 18 middle category not be support, conditionally 19 support, but to say that, you know, that it 20 really needs to be, and these issues need to be 21 taken into consideration specifically.

MEMBER GOODRICH: With that, which I

think is fine, the one thing that this committee will have to manage is the issue we were talking about before of getting too deep into the weeds and recreating the endorsement process.

I just think that's going to be true not matter what, but just to highlight that as well.

CO-CHAIR PINCUS: One of the things, actually if you go back to the slide before this, so -- Actually, the slide before that. So if you read that, you know, actually, right here it talks about the program measure set is adequately addressing the needs of the program, these responses to the program, and sometimes we kind of lose sight of the linkage of the measures to the program.

And I think that that's, you know, and that's really in some ways the heart of what we are addressing, more than the sort of endorsement criteria, but the way in which it is intended to be used.

And so that's something that we need

to sort of, you know, think about as we do that, 1 2 and that's part of the issue and that's irrespective of whether it's endorsed or not. 3 Pierre, did you want to say something 4 about that or --5 DR. YONG: No, I think that's a fair 6 7 point. I mean, something that we also included 8 as part of one of the draft criteria when we 9 talked about measure removal criteria, was sort 10 of, as we think about measures, they are for an entity, whether it's a facility or a clinician, 11 12 it's part of an entire sort of reporting program. So, I think it's a fair point about 13 14 that we -- at least based on the couple years of the MAP that I've been able to attend -- have 15 16 really been thinking more about the MUC list as 17 individual measures, as opposed in the context of 18 the entirety of the set. 19 CO-CHAIR PINCUS: So, let's go around. 20 John? 21 MEMBER BOTT: Yes, I like Amy's initial suggestion of vote a measure up or down, just 22

binary basically. And if it's not yet NQF endorsed and we've been presented with adequate information to vote it up, so be it.

The way it's currently written, it doesn't have to be NQF endorsed for us to support for rulemaking. If we don't feel we have enough information or we feel it's inadequate, then we vote it down.

I like the idea of providing those supplementary comments with do not support to explain why we can't support it at this time.

So, I vote for simplifying our voting that way.

CO-CHAIR PINCUS: Leah?

MEMBER BINDER: I would like to see some way for us to vote for fast-tracking for endorsement, because there are issues that do merge, and there will be more.

So, opioid would be a good example of something that's just a real emergency, and we need to not wait two years to get this through endorsement or something, because then it takes time after rulemaking and everything else.

So, I think that there are going to be emergent issues -- or maybe not emergent, but urgent issues or areas where there's an urgent gap.

Again, I would reiterate my point about the ambulatory surgical center safety and quality measures, for which there is an urgent gap. We don't have enough measures for that, and 60 percent of surgeries are being done there. So, we have to, I think, have the flexibility to be able to put a sticker on something that's around timing.

I actually don't have a problem with refine and resubmit, I think that sounds actually -- that makes sense to me. That makes a lot of sense. I think the conditional, defining that, and -- but again, adding something where we can ask for an urgent process to put through endorsement.

CO-CHAIR PINCUS: We have Rich

Antonelli on the phone, who I think wants to make
a comment. Rich?

MEMBER ANTONELLI: Yes, thank you. I want to sort of raise an issue that I often start my own team around QI, is: what's the problem we're trying to fix?

So, a lot of the really interesting observations and good ideas and comments that I've heard are reflective -- are self-reflective of MAP members. But I'm wondering, other than for us feeling like there is a more harmonized, synchronized approach across our processing, are we missing something?

And I'd like to just sort of put two comments on the table for our consideration, to think of what we would measure success at the end of this conversation.

One, and I don't know if he's still in the room, but Sam, I think, had such a good point yesterday, when we were looking at one of the measures where it actually failed under scientific reliability. And I was thinking about, okay, well, we just spent of time talking about that.

And two, both here in Massachusetts and other conversations, this is kind of in line with the comment about the opioids, there are times where there is a strategic gap that is very time-sensitive when we're going to advance something. And so, I think there will be times where we're going to have to frame shift and do that.

So, I guess, that would be the question I'd like to raise is, what's the problem we're trying to fix? Because then, the solution to these conversations around process may be better aligned.

CO-CHAIR PINCUS: Thank you, Rich. Giff?

MEMBER GIFFORD: Thanks. So, let me ask Sam and then -- so, a couple of examples of where things were not ready, even though they were NQF-endorsed. The main one, I think, as Bruce's point has been, they've not been tested in the other setting.

The other is, the discussion we had

there, they may have been tested in the setting, but it's an attribution issue. And so, for the program, attribution is important, but for NQF endorsement, it was not important.

The other, as we talked a little bit before, the infrastructure may not exist or the time it takes. So, it might be that we endorse it for rulemaking, but it needs to be one or two years out to build the infrastructure.

Because we've had some rules go
through where it relies on MDS items that CMS has
tested in a pilot, but has not been rolled out
yet. And so, there's just no way you could -- I
mean, as you point out, there's a huge cost to
changing infrastructure and going forward.

And then, I think the other is the unintended impact the measures might have in the program, and that that should be addressed in the rulemaking. So, it's more about, not whether the measure -- some of it's the measure itself, but how it's used in the measure set in the broader issue.

I think the problem we're trying -one of the problems I think we're trying to
identify, having been through this now is,
particularly when measures come through that are
not NQF-endorsed, it causes great angst. And it
really confuses our roles with the endorsement
process out there.

And while I agree with Amy's point,
the one thing I worry about is that, in the past,
if we endorse it not ready for rulemaking, that
-- as you point out, Leah -- that makes it
politically really hard for CMS to go forward.

They have, but it makes it really hard for them to go forward. And I wouldn't want it to be an absolute that, just because there's not been endorsement, we should vote it down. It should be, if there's no endorsement, it's just so early, it can't go forward.

But I think, you almost want to have in each of those categories why you're voting that, because I think if we went back to just two, it wouldn't -- the problem we were always

trying to address was this anxiety of both, not that or that, yes, we understand the measure's not quite ready for rulemaking, but we think this rule and the direction and the measure concept is good, but you need to address attribution and other things.

And then, CMS will either address it in that rule, I've seen them delay and it'll come out a year or two years later, and they have done the homework to address it. So, I think just a straight up and down vote might lose that -- will bring us back to that anxiety.

And we ended up having trouble voting on some of the things because of the labels. So, I think we are looking for a softer label to say, yes, you can use it in a rule, but make sure you address these topics.

CO-CHAIR PINCUS: Rachel? And then Bruce.

MEMBER LA CROIX: I think this follows along with what a few folks have been talking about, and I understand the desire of wanting to

go to just a support or do not support.

One thing that I was thinking about, and this might be too soft of a third potential category, but the workgroups all identified basically priority gap areas, areas where they want to find new measures and keep working on.

And I don't know that that's something we discuss in here, as the Coordinating Committee at all.

But a potential third area would be for those measures or areas that don't seem quite ready or that are missing a key piece for us to fully support them, would be either to set those up as important priority gap areas that we think should be addressed moving forward or to help support just some of those other ideas that aren't ready yet or where we don't have measures.

And maybe that would be a third category to provide food for thought moving forward.

CO-CHAIR PINCUS: Bruce?

MEMBER HALL: I don't want everybody to think I'm a one-trick pony here, but I want to

build on two comments by Giff and my esteemed 1 2 colleague, John. I think that we are not a committee 3 4 that is structured or capable of doing a 5 technical review, so I disagree that if it's not an NQF measure, we can just look at the technical 6 7 specs and move it on or not move it on. 8 I don't think we're equipped to do 9 that in any way, shape, or form. Having done it on the NQF side, where I think it's very 10 valuable. 11 12 So, I would ask, then, for a measure 13 that's not NQF-endorsed, is it in the endorsement 14 process and what is the expected date of completion? 15 16 If it's not in an NQF endorsement 17 process, where we agree not everything has to be, 18 then who is the technical review body that's 19 vouching for that? And if that's not complete, 20 when will it be complete? 21 And perhaps that's the other category 22 we need, to say, this is in the NQF endorsement

process with the following expected date, or this 1 2 technical review is performed by someone else, who is that? 3 4 And then, that becomes another 5 category that we can say, we would move it forward, but we can't vouch for that technical 6 review; that technical review was performed by 7 8 someone else. 9 CO-CHAIR PINCUS: So, you're thinking -- talking about essentially having a third 10 11 category that's focused on a process item? 12 MEMBER HALL: I think, yes, in that 13 I think calling out that category that sense. 14 we're now confused where to put it in the middle, calling it out for what it is and saying, this is 15 16 not yet NQF endorsed, but it's expected on the 17 following date, or this technical endorsement is by someone else, and that's what needs to be 18 19 represented to CMS. 20 CO-CHAIR PINCUS: Sam?

Following up on both Giff and Bruce at this

MEMBER LIN: Thank you, Harold.

21

point, this may be an anathema in the current market, but I'm sort of old-fashioned, I still believe in science. And so, I think the first thing is that the scientific basis is the first item that's got to come out of the shoot.

What we do, and this follows up with Bruce, is: we are not the scientific experts, we're the socialization experts -- if I can use that term -- which is the world or the context in which that science-based evidence is going to be used.

So, that's sort of where I go back to my earlier question about what other than NQF endorsed, what does conditional mean?

of being evidence-based, scientifically-endorsed, then we can sort of look at the wrap-around and the packaging, and I'm not trying to demean our role, but that's sort of what we do, the rest of the world that this fits into, and then, decide if that fits in with that and then, whether or not we can, by consensus, support it.

So, I mean, I think we're sort of getting to some area of changes. I'm still, I don't know, I'm still supporting the up or down, because it's pretty clear, at that point, regardless of what the vote numbers are, if we think about consensus, then it's pretty clear that the group is moving on to the next measure, rather than continuing to drill down.

Before I forget, since we've non-voted the up or down, could we go to that next slide?

Because there's one thing that this -- Item 4,

it's just hit me yesterday, the definition of do not support for rulemaking.

The definition or the criteria, sorry, is that the measure under consideration does not meet one or more of the assessments. I think it's better stated, the measure does not meet any of the assessments.

If it meets, for example, refine and resubmit, that is sort of a conditional support.

It's not a do not support, it's a conditional support requiring refine and resubmit. If it

goes under conditional support, that's not a do 1 2 not support; it's a conditional support. So, I think, hopefully we're going to 3 change this, but if we don't, I would suggest 4 5 that we might clarify that criteria to say that the measure under consideration does not meet any 6 of the Assessments 1 through 3. That's all. 7 8 CO-CHAIR PINCUS: Sam, I appreciate 9 your comments about science and evidence-based. 10 Last year, I published a paper in Lancet 11 Psychiatry on evidence-based science policy in a 12 post-truth era. 13 MEMBER LIN: Post what? 14 CO-CHAIR PINCUS: Post-truth era. Leah? 15 16 MEMBER BINDER: I want to pick up on 17 something that Bruce just said that was kind of a 18 You mentioned setting a timeline, saying 19 we want it endorsed -- or when is it going to be 20 endorsed, by what date? 21 What if we said that the conditional 22 category was actually very specific -- that it is

to be endorsed within one year, period -- go
through endorsement within one year or six
months? That we were specific about what the
condition was and a timeline for it?

And then, it should be brought back to

MAP by X. And say, we want it back. Because we've all talked about that, like what happens to these things? So, that we're like -- so, conditional support is specific and time-bound.

And getting to my point, that gives us a little bit more of an ability to say when something is urgent as opposed to non-urgent.

CO-CHAIR PINCUS: Giff?

MEMBER GIFFORD: It would be almost helpful, and this gets into the next thing about the criteria review, what are -- to Sam's point -- what are the conditions that would, if they don't pass, would lead us to all vote, don't support in rulemaking?

And, clearly, endorsement is one, but not an absolute. But I think attribution would be another, tested in -- we could come up with a

set, and then guidance from the workgroups and guidance from the staff would help us in determining the decision out here, and take us away from the discussion about measurement specification.

Burden would be one, another category that I think we spend a lot of time on. But there is sort of a finite number of things that should lead us, as a MAP for rulemaking decisions, not about the endorsement.

And then, it takes that focus of our discussion on that piece rather than on the measure specs.

CO-CHAIR PINCUS: Okay. Elisa?

MS. MUNTHALI: I just wanted to clarify a couple of things related to the endorsement process. In the old process, there was significant wait time, and so there were -- it was about a two or three year lag time between projects.

In our redesign -- which Erin mentioned yesterday -- we're offering submission

times twice yearly. And so, developers can come in any topic twice to us, and that process takes about seven months. So, it is not two to three years.

And that's why yesterday, during our dialogue, we did the redesign in very close collaboration with CMS and, in fact, they funded it.

So, we were trying to the best of our ability to build in a pipeline, so we can understand what measures CMS needed to come through the endorsement process, but other developers as well.

And so, I'm -- as you've heard, it's really hard to kind of coordinate that timing, but we're still working on it. And so there are more opportunities than there were in the past to come to NQF.

CO-CHAIR PINCUS: Leah?

MEMBER BINDER: Can time be carved out on the next cycle or the next meeting, to launch the endorsement process, that's just for what

comes out of the MAP as -- that should be started on endorsement right away?

So, in other words, if we see a measure come up that we know needs to be endorsed, that we see as a priority, it's coming from CMS for rulemaking, that we as a MAP can say, well, we would like it put into the high-priority endorsement, and then it's reserved.

So, it's not like they have to wait even six months; it's just the next meeting it gets started. In other words, so that we can have some capacity to identify those measures that need that very quick turn.

Because, by the way, rulemaking itself takes another year and then, it takes another three years beyond that to even get the data.

So, no matter what, we're talking about very long lag times for some of these urgent issues.

So, I would like to see if we would have the capacity, really, right now to say, you have a meeting coming up in three months, even though it's not normal that it be that short of

time -- turnaround to get something in, that for that measure, we could say we want it to be put in.

CO-CHAIR PINCUS: I mean, it seems to me that what could be done is to actually put together a small group with staff to actually come up with some proposals that would be brought up at the next meeting.

And I think that that makes a lot of sense. I think there's a lot that came up in this discussion where I think there's some reasonable consensus on, in terms of, number one, we need to clarify these four categories and make some changes and reduce the number of categories.

Number two, we want to make sure that we're able to give very clear, specific instructions to CMS about what needs to be done for measures that we don't support.

Number three, that we need to be very clear that there are measures that we support, and the other measures we don't support, to distinguish that.

We need to also, I think, make sure that we get more feedback about the process, in terms of what's happened after our recommendations, so that we're informed by that and we can learn from the kind of recommendations that are most helpful.

And I think also, the notion of specificity in our recommendations, both in terms of content and in terms of process, would be important.

But I think -- which I think bodes
well for a small group to actually get together
and come up with how to take that into a specific
set of recommendations. Is that something that
you think could be done?

MS. O'ROURKE: Yes. So, I think in the last bucket of information, we actually already started having that conversation.

So, if we could bring up the algorithm, I think this is something, the last thing we really wanted to get some input, because I think we already started having a conversation

about this type of information that you all need to support your decision-making.

And based on what you were talking about yesterday, I think we would also like to hear, not just about this algorithm, but this feedback loop process, what other information you would like to come back through other channels.

But to support some of the conversation you were just having about the fit-for-purpose, and that was one of the things we heard with this algorithm, is that it drives a little too much back to revisiting endorsement decisions or starting to do the work of the endorsement committees.

And to Bruce's point, we're not necessarily constituted that way. So, I think, Giff, you had some great suggestions about how we focus more on the fit-for-purpose of the program. So, I think any other suggestions people have there would be great.

MS. BUCHANAN: And, Erin, if I could just add? So, everyone received a link to the

MAP Member Guidebook and that is on -- Page 27 is what we're looking at, so if you want to pull it up on your own computer or if you're participating through the phone, it's Page 27 of the MAP Member Guidebook.

MS. O'ROURKE: Yes, so any input people have about what should be in the algorithm, different assessments you'd like to see to make sure that the workgroups are getting the type of information that you want them to have.

And then, similarly, I think we'd love any information that you'd like to see through the feedback loop process or analytics that the staff could perform on what went into the rules this year in concordance with MAP's decision-making, what would be valuable to you as a Coordinating Committee.

DR. AMIN: So, just to potentially get the conversation started, there's a few themes that we've already discussed that we can sort of discuss right now.

I think one of the discussions that

we've heard a lot about, in terms of additional data, is what happens with the MAP recommendations over time? So, I think this sort of concordance-type analysis would be helpful, Erin.

And I think the other thing that we've talked about is related to a little bit of the endorsement pathway and also, the endorsement experience for the measure. The way that it's currently represented in the discussion guide is sort of an absolute yes or no, whether the measure's been endorsed.

But it doesn't get to the nuances related to the level of analysis, of whether it's tested at the clinician level, for example. Or whether it's been resubmitted, which version of the measure. Or, to Leah's point as well, what's the pathway for endorsement, in terms of when the developers would be ready to submit that information to an upcoming cycle?

So, those are, I think, some recommendations that we've heard already, which

1	are pretty significant in themselves. But at
2	least to get the conversation started.
3	CO-CHAIR PINCUS: Comments?
4	Suggestions?
5	CO-CHAIR KAHN: So, I think it's a hard
6	one to slice. I mean, I almost wonder whether we
7	should ask the group to come back, if they feel
8	like I mean, with written comments on the I
9	mean, because we could discuss it, but to me,
LO	it's a little difficult. Or if you want to set a
L1	few people to look at it and come back with
L2	recommendations.
L3	CO-CHAIR PINCUS: Yes, I think it would
L <b>4</b>	be good to I mean, and either way, I think
L5	that we should assign a small group to work with
L6	staff that would be charged to review sort of the
L7	process criteria and decision algorithm.
L8	MS. O'ROURKE: Okay.
L9	CO-CHAIR PINCUS: And to as a
20	package, because it's really hard to separate
21	them out.
22	MS. O'ROURKE: Yes.

CO-CHAIR PINCUS: And to come back with a set of recommendations, or even options.

MEMBER GIFFORD: Yes, I think that's a good idea. And if you're going to do that, it would seem to me, part of the charge to the group is how the information is presented to us, how the material -- what the decision-making process was ahead of time.

But given the very tight time lines we're under, to the degree to which we can figure out a way to make the process of our review of the material, I'm not sure easier is the right word, but a bit more streamlined, since getting this morass of stuff a couple of days before the meeting is almost impossible to go through with any clarity of thought.

How we incorporate public commentary and how in fact we either use ahead of time in the agenda books or following, written commentary that might actually make some of the conversations a bit more succinct.

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So, it's a lot of process questions,

Harold and Chip, but I think that we really need to look at the way we do our work and how we get input into the whole process.

Because right now, I think there's a sense, and it gets to, I think, comments you made, Chip, is that, without meeting frequently enough and without having the ability to actually spend the time to actually ponder some of these questions, they're almost done on the, I'm not meaning to be pejorative, but it's almost like you're doing them on the fly, because it's just so much stuff that comes at you, like in a week.

CO-CHAIR PINCUS: Marissa?

MEMBER SCHLAIFER: The one question,
one comment I had when I reviewed this in advance
may have been kind of overridden by some of the
comments we've had over the last couple of days,
but I think one of the -- with one of the
questions being, can you combine the
reliability/validity criteria with the NQF
endorsement process?

I think it's really clear when we're

talking about using a measure in the population 1 2 for which it was NQF endorsed. And then, you could combine them. 3 But we did have at least one measure 4 5 where it may be NQF-endorsed, but we were looking at it -- but it was endorsed at the health plans 6 7 level and we were talking about it at a provider 8 level. 9 And in that case, you do need to pull out, pull apart the reliability, is this reliable 10 and valid, from the NQF endorsement process. 11 I think, yes, sometimes, but, no, other times. 12 13 CO-CHAIR PINCUS: Any comments from 14 people on the phone? MEMBER KAHN: Hi, this is Maureen. 15 16 I will tell you, as a first-time attendee, this 17 is my first meeting, I'm overwhelmed with all the 18 comments on this. So, I'm hanging in there. 19 (Laughter.) CO-CHAIR PINCUS: We all are. 20 MEMBER GIFFORD: Just to follow up on 21 22

MEMBER KAHN: I can't see everybody's faces, but it's been a -- I've appreciated the conversation and the dialogue. And -- because I read the 236 pages and had a couple of sleepless nights to be prepared for the meeting, and the book, and I take this very seriously. And so, I do value the comments that I've heard.

CO-CHAIR PINCUS: Giff, you had a comment?

MEMBER GIFFORD: Yes, I was just going to say, to Marissa's point, it has implications for, I think, CMS providing information on the MUC list to the workgroups, and then to us, to figure out how to do the vote.

Because I think, if it's NQF-endorsed, we don't need to know all the other stuff. What we really want to know is attribution, other stuff. If it's not NQF-endorsed, I think Bruce laid out really nicely what we want.

And then, that information will expedite our ability to read through the 200-odd pages, and I think get more robust feedback from

the workgroups to us as well.

CO-CHAIR PINCUS: Okay. Other comments from the phone or here? So, and thanks, Giff sent us some of the ideas for different categorical kind of the issues that might come up and I think this could be useful in the discussion around the smaller group that we put together.

MEMBER GIFFORD: Question, and this is just my own ignorance, I could go back and look, but are there any members of this Coordinating Committee on any of the four workgroups?

MS. O'ROURKE: Back in the day, there were. I think we purposefully tried to limit it, so that as many stakeholders could participate in MAP --

MEMBER GIFFORD: I understand; the answer is no. So, what I'm getting at actually, from the perspective of the memory of this group, and also the trust that we have in the work, I'm not suggesting we put people from this group on it, I would suggest that we think about creating

a liaison, where in fact there's an expectation, one, that all of us actually at least go to one or participate on the phone in one in their first year of membership, so we know what the devil they're doing.

And secondly, that in fact there be a liaison so that when in this room, recognizing we have the Chairs on the phone, but to the degree that there's a little bit of a cross-check, especially when there's discontinuity between the staff and the decision, that we've got a voice from this room that at least understands the dialogues.

Just, again, as you're thinking about process, another suggestion.

CO-CHAIR PINCUS: I mean, there was a little bit of that, Marissa and I and Rich, we're sort of hanging around the workgroups, in our roles with the Medicaid Task Force.

MEMBER SCHLAIFER: Yes, to follow what Harold said, we were required to sit in on it, because of trying to get a Medicaid tie, but it

was trying to get a Medicaid tie, not trying to 1 2 get a tie to this group. Although, it did serve that --3 4 CO-CHAIR PINCUS: Yes. 5 MEMBER SCHLAIFER: -- purpose. 6 guess, we can dual --7 CO-CHAIR PINCUS: Yes, no, but I think 8 your point, I think you made a very good point 9 about, as almost part of the orientation process, and sort of understanding what's going on. 10 11 And I would add, it is important also, 12 I think, because -- how many of you have actually 13 been involved in the endorsement process? 14 almost -- about 60 percent. MEMBER SCHLAIFER: We should get our 15 16 clickers out. 17 (Laughter.) 18 CO-CHAIR PINCUS: But some of you 19 haven't, and I think it would be actually 20 important to also give people some experience and 21 also some more background on the endorsement process, so they know what's entailed with that. 22

1	Sam?
2	MEMBER LIN: I just remembered Frank
3	Opelka, remember when he was the Chair
4	MEMBER SCHLAIFER: Yes.
5	MEMBER LIN: of the Hospital and he
6	was
7	MEMBER SCHLAIFER: Yes, he was on
8	MEMBER LIN: on the actual I
9	mean, that was
10	CO-CHAIR PINCUS: Yes, and originally,
11	there were people who were on in fact, it was
12	almost, I think, required that the Chairs be
13	members of this.
14	MEMBER GIFFORD: I would second Carl,
15	I've always gone to the PAC group, and it's both
16	reassuring and not reassuring.
17	(Laughter.)
18	MEMBER SCHLAIFER: Yes, I can say I
19	learned so much this year sitting in on it.
20	MEMBER HALL: I want to say, I replaced
21	Frank Opelka, and I know I'm not as good looking
22	as he was, but I'll tell him you were remembering

1	him fondly.
2	CO-CHAIR PINCUS: Okay. So, we will
3	sort of convene with the staff and sort of come
4	up with a small group to look at this and look at
5	these issues. Are we going now to the Rural
6	Workgroup?
7	MS. O'ROURKE: Yes. So, I think we
8	should do a public comment; we haven't done one
9	yet this morning.
10	CO-CHAIR PINCUS: Okay.
11	MS. O'ROURKE: And then, Amy and Chris
12	are our lead discussants, and I think they both
13	need to leave in a few minutes, so if they can
14	make their comments and then, do Karen's
15	presentation, if that's okay? Just so we let
16	them catch their flights on time.
17	(Laughter.)
18	CO-CHAIR PINCUS: Okay.
19	MS. O'ROURKE: But, if we could open
20	for comment first.
21	CO-CHAIR PINCUS: Okay. Anybody from
22	the room wish to make a comment? Okay. Anybody

1 on the phone? Can we open up the phones for 2 public comment? OPERATOR: Yes, sir. At this time, if 3 4 you would like to make a comment, please press 5 star, then the number 1. There are no comments at this time. 6 7 CO-CHAIR PINCUS: Okay. So, let's 8 proceed to the Rural. 9 MS. O'ROURKE: Great. Amy, Chris, 10 would you mind giving your thoughts? I just want 11 to make sure we get your --12 MEMBER MULLINS: Yes, so, as I was 13 looking through the slides for this presentation, 14 one of the things that struck me is, I was very The Chair is 15 happy to see a Rural Workgroup. 16 actually an AAFP member, and another member of 17 the workgroup is also an AAFP member, so I think 18 it's very important that family medicine is 19 represented in that capacity. 20 One of the things that I just wanted 21 to comment on, and I'm sure that there would be 22 more explanation if I could actually see the

whole presentation, but one of the things it said was that making rural participants a mandatory participation in CMS quality measurements and QI programs, that was on one of the slides.

So, in MIPS specifically, there is a low-volume threshold that has been set forth that is going to exclude, and actually, if you read the rule -- which I've had the pleasure of doing a couple of times -- it actually was set to exclude rural physicians.

That was actually in the verbiage; it was actually set higher the second year to exclude more small and rural physicians. So, just saying you want to make participation mandatory in CMS quality programs can't just happen. That would have to go through a rulemaking process.

What we would ask for, and actually what the Academy asked for, because of this higher low-volume threshold, which we actually liked in some ways and didn't like in others, because as the second year participation comes

around, some physicians are going to be kicked out of the program that were once participants and maybe wanted to participate. Some people are going to be happy about it, because they once participated and didn't want to ever participate.

so, what we asked for and didn't get, and Pierre knows this, we've had this conversation, but is an opt-in. We wanted an opt-in, so if you wanted to participate, but you were kicked out because of the low-volume threshold, that you could opt-in to the program and report if you chose to. So, if we could maybe align our messages to CMS along that vein, that would be helpful.

CO-CHAIR PINCUS: Chris?

MEMBER QUERAM: Thank you. Just a couple of comments. Just for context, in our state -- I come from Wisconsin -- about 50 percent of the hospitals are rural or Critical Access Hospitals.

So, it's a significant part of the landscape in our state, as I'm sure it is in

Amy's state as well. And one of the formidable advocates in our state is a member of this workgroup, Tim Size, who is the executive director of the Rural Wisconsin Health Cooperative.

And in talking with Tim a little bit about this workgroup and invited him to help influence how I might characterize a perspective on this, words that he uses are "stigma" and "backwater."

He's very concerned that on behalf of the constituency that he represents, that fact that rural hospitals are not eligible to participate or find it difficult to participate in CMS's programs, payment programs, perpetuates a stigma that rural hospitals, Critical Access Hospitals, are backwaters -- clinical backwaters.

And to provide another context for this, a political science professor at the University of Wisconsin, Kathleen Cramer,

Katherine Cramer wrote a book called The Politics of Resentment. It's gotten some publicity here

in Washington, because it coincidentally captures some of the themes that emerged in the 2016 presidential election.

She actually did an ethnography model or study of attitudes, beliefs, values, in the State of Wisconsin, based on a longitudinal series of interviews that were done in about 30 different communities in the state, and over a period of 2010 to 2015.

So, predating some of what emerged more forcefully at a national level. And uncovered just a tremendous amount of resentment in rural areas towards elites and how they make decisions and how they allocate resources.

And the takeaway theme from that book was that there's a fundamental distrust that politicians, and maybe I'd reframe that as the establishment -- in this case, CMS acting on behalf of the administration -- any administration, not this particular one necessarily -- will fail to recognize the distinct identities and values of their

communities and fail to allocate resources on 1 2 their behalf. So, as we think about the work of this 3 4 workgroup, I'd put it in that context -- that 5 this is critically important work to counteract some of the burgeoning attitudes, the rural/urban 6 divide that we are experiencing in Wisconsin, and 7 restore some faith in the ability of these 8 9 facilities to participate in as full and an equal 10 manner in the CMS programs as possible. 11 CO-CHAIR PINCUS: Thank you. 12 Karen, you want to give your presentation now? 13 MS. JOHNSON: Sure, and thank you. 14 Kate, are you going to start us off or have you already done the first bit? Apologies. 15 16 MS. BUCHANAN: So, I did an overview of 17 the feedback that the MAP Rural provided on the 18 MUC list, but that's -- so there is familiarity 19 with that. 20 MS. JOHNSON: Okay, apologies for not 21 having my ducks in a row there. So, my name is 22 Karen Johnson, I'm one of the Senior Directors

here at NQF. Thank you, Lisa. I should know better by now. And I have the privilege, actually, of leading this work.

It's very exciting for me personally, because I'm from Central Appalachia, so it's a very rural area, so this topic is near and dear to my heart. And Kate is also one of the folks that are on the team with me.

So, what I wanted to do today is three things. One, give you the background of how we got here today, tell you a little bit about what we're doing in this current project with the MAP.

And then, get some feedback from you guys.

So, I hope -- how much time do I have?

I have 45 minutes or I know, I'm in-between you

and lunch and that's a dangerous place to be.

(Laughter.)

MS. JOHNSON: Okay. All right. I'll try to talk fast. So, back in 2015, CMS, in cooperation with HRSA, asked NQF to do some work on rural. And basically, what they wanted from us was to provide multi-stakeholder information

and guidance on performance measurement and challenges for rural providers.

And it was really in the context of CMS payment programs. So, that was the context that we were working under. And they wanted us to make recommendations about measures that would be appropriate for these kinds of programs, make recommendations to help mitigate some challenges that we know about, and also to identify gaps in measurement.

And that project was limited in scope, somewhat. We were looking mainly at ambulatory and hospital settings, so we didn't really get into the post-acute world. But what was really interesting was that we weren't just thinking about the rural providers who were paid through the PPS system.

It also, by design, included Critical Access Hospitals, rural health centers, and rural FQHCs. And the latter two are paid in a different way and don't actually come under PPS rules.

so, that was interesting, because especially the last two groups pretty much don't participate in the programs that we're all used to talking about and thinking about. And Critical Access Hospitals do participate in some of the programs, but only on a voluntary basis.

So, this project was meant to really understand some of the challenges of those groups, but also small rural hospitals and small clinician practices that do participate in PPS.

So, two different groups, some who actually do participate, others that have never, and trying to bring those two things together.

It was a great group, meeting Tim was great, and he's on the current one. He was on that last one in 2015. We had a lot of people from these different groups that we don't normally get to talk to and a lot of on-the-ground providers as well. So, do I -- I guess this -- yes, there we go.

So, some of the key issues and challenges in terms of measurement, we

articulated in this work. And I'm not going to go into major detail about these. They are clearly related, but they are kind of different concepts.

The idea of isolation that is true of some, but not all rural providers. And that gets into the ideas of having specialists who may or may not be available, having transportation issues, IT challenges, perhaps, other kinds of limited support.

And all these things, again, the idea here was how do these things impact measurement and how do we think about it? Small practice size, where we were getting at with the small practice size really was this idea of limited staff time.

So, one of my favorite stories was, actually our Co-Chair of this project, who talked about not only being the MD of his rural health clinic and working in a CAH associated with it, but also being the IT guy and the plumber when they needed those services. Really small and

tiny.

And when it comes to measurement, having very few people, it makes it difficult to collect data for measurement and to do QI activities.

In terms of heterogeneity, we really talked a lot about differences in rural residents. So, where I'm from, Central Appalachia, is a different animal than the Deep South, and that's different than Alaska and maybe the Western states in terms of social factors and those sorts of things, cultural attitudes.

All those things come into play when you start thinking about risk adjustment, for example. So, heterogeneity, really big. And also, heterogeneity in the types of services that are done in rural hospitals and clinics.

So, especially thinking about small hospitals, and I actually never really thought about this, not all of them do surgery, not all of them delivery babies, or at least maybe only on an emergency kind of situation.

So, when you start thinking about measures and measures that are used in programs, some measures just don't apply. And then, you get into the low case volume problem.

That really gets to reliability of measurement and being able to -- when you have a set of measures that you're looking at in programs, if a particular provider can't report on a few of the measures, that means really more weight is hanging on others.

And that may be good or not so good,
but it can be a problem. So, with all of that -and the other thing is, not all of these things,
of course, are rural-only.

As a matter of fact, we were quite challenged to come up with things, probably other than the isolation piece, that was really a rural-only problem. But it was really -- it exacerbated the measurement issues for rural providers, maybe in ways that it wouldn't for other providers in suburban or urban areas.

So, the overarching recommendation,

and Amy has already talked about this, was that eventually we'd like to make -- and when I say, we, I mean, they, the committee at the time -- would like to make participation in CMS quality programs mandatory for all rural providers.

So, this is not just the ones under PPS, they were talking much more broadly than that. But also, understanding that, given the experience, a phased approach of some sort would be needed. And you'd have to explicitly think about the low case volume challenge.

So, with that -- and it was interesting, it was, honestly, for me, a bit of a surprising recommendation, but it was one that had full consensus around our table, as well as consensus from our commenters on the report.

So, this is something, and it goes back to what Chris had mentioned that Tim really was talking about, which is, by being left out, and that was their phrase, being left out of these programs, they're kind of left behind, as it were.

So, they don't have the opportunity to maybe understand their quality as well as others. They don't have the opportunity to participate in bonus payments, potentially. They don't have the opportunity to publicize how good they can be in providing care.

So, those were some of the ideas that we talked about and that underlie -- I don't know what the right word is right now -- that underpinned the overarching recommendation.

They also provided many other recommendations to help support a potential transition to participation for all rural providers. And many of these recommendations had to do with development of rural-relevant measures, alignment of measurement efforts.

Interestingly enough, that wasn't just about alignment of measures, but also about alignment of data collection efforts and alignment of informational resources and TA, technical assistance, kinds of opportunities.

And they also had some recommendations around

payment considerations.

But for today, I want to highlight the recommendations they provided in terms of thinking about measure selection for programs for CMS.

So, I have these on the board, I'm not going to read them word-by-word, but really, they wanted to use guiding principles, and they articled some guiding principles to be used when it's time to select measures that would work for rural providers.

They suggested identifying a core set of measures, as well as an optional set. So, have core and optional in these kinds of programs.

They wanted us to be sure not to forget PCMH models and the measures that are used in those systems. And they also suggested or recommended creating a MAP workgroup to advise CMS.

So, fast-forwarding two years, I'm sitting here today to tell you, we're really

excited that CMS took us up on that recommendation and we just have seated our MAP Workgroup for Rural Health.

The objectives of this year's work is listed here. Basically, we want to articulate a set of criteria for selecting measures. So, two years ago, we had guiding principles, now we're getting a little bit more concrete, what are the criteria?

We're actually going to identify a core set of measures, rural-relevant measures to address the needs of the rural population. For this year, we are concentrating on the hospital inpatient and the ambulatory settings. So, again, pushing off post-acute care for now.

We, as usual for MAP work, will talk about gaps in measurement, provide recommendations around alignment. And also, kind of an interesting piece that we're going to be doing is addressing a measurement topic area relevant to individuals and measurements for rural residents.

And we actually just yesterday got consensus, I think, from the group, so we think we know where we're going on that. And I'll talk about that a little bit more very soon.

In terms of our interaction with you guys and the other workgroups, we kind of did a meet-and-greet with all the other workgroups, back in November/December. We're telling you a little bit more about us today, the Coordinating Committee.

And in our last call with them, we did have a chance, a very small chance, to look at the MUC list and offer some high level input, that I know Kate has already shared with you.

And then, finally, in August, you guys will have the opportunity to look at what we've done over the year and, hopefully, approve what we come up with.

So, our progress to date, just so you know kind of where we are, we -- this is moving fast for us. We put out our call for noms for this workgroup at the end of September. We had

seated this workgroup by mid-November.

Had our first webinar with them at the end of November. And a second one in the middle of December. And then, our third one yesterday. So, it's moving right along, roughly about once a month, we're contacting everybody.

We're doing it all via webinar and the thinking there was, we don't want to pull some of these rural providers who are really busy on-the-ground providers of care and have them travel to D.C. for a couple of days, so we're doing everything virtually. So, that's been interesting, but it's worked so far.

So, what we have done so far is obtain guidance from our workgroup on the criteria. So,

I mentioned we want to articulate our selection

criteria in order to identify core sets of

measures.

So, our criteria to date are that we want to pick from NQF endorsed measures. And what that does is it hits several of the guiding principles that the committee from two years ago

outlined, because NQF endorsement guarantees evidence-based measurement, room for improvement, feasibility, scientific acceptability, that sort of thing.

So, they checked off several of those principles by saying, let's focus on NQF endorsed measures. For this core set, we also are really interested in finding measures that address low case volume.

Address probably isn't quite the right word, but basically, it's the idea of, you don't want measures that a lot of rural providers won't have enough cases to report. Cross-cutting measures is another thing that seemed really important.

So, and cross-cutting and addressing low case volume are related. They're not necessarily mutually exclusive, but if you have cross-cutting measures, often that will mean you also may not have to worry quite so much about low case volume.

They also mentioned several, what I'm

calling right now, must-have topic areas or conditions. So, the ones that have come up are things like mental health and substance use, medication reconciliation, transitions, moving from one setting to another, diabetes, high blood pressure, COPD.

Also of interest has been at least considering measures of readmission and thinking about how and when to bring in measures around perinatal care and pediatric care.

So, yesterday on our call, we presented a draft core set that we came up with by applying these criteria. And we basically provided the Committee a strawman set of core measures, we came up with 44 of them to cover the inpatient and ambulatory settings.

And over the next week, we're going to continue to obtain feedback from them. There may be more than 44 that they want to think about and then, we'll start winnowing down that 44-plus to something that's reasonable for a core set.

And just so you know, our thinking in

terms of what would be a reasonable core set would be probably ten to 20 measures per set.

So, certainly not -- other things could potentially go in optional sets, if we are funded at some point to do that work.

So, that is my kind of lecture part.

So, discussion questions. Let me just throw

those out for you and we can go in whatever

direction you would like to go. But are there

key measurement or programmatic issues that we

should keep in mind when identifying our core

sets?

What are your reaction to the criteria that I discussed? Again, cross-cutting, addressing low case volume, and certain conditions. And just so you know, the conditions that were brought up were ones that are really quite prevalent overall and particularly in rural areas.

Do you have any advice in terms of what would be important to emphasize as we try to explain our results? We're going to be writing a

report, like we always do here at NQF, and we want to make sure that it's a useful document.

Going forward, what information or input or guidance from them to you would be of help? So, what can we bring back to you?

And then, finally, I didn't put it on this list, but after yesterday and in a lot of our discussions, the idea of access to care was really, really huge. And that's a different way of thinking about measurement and access is not the same as quality measures, right?

So, they're different animals. And so, we are thinking that that will probably be our measurement topic area. So, if you have any ideas on how we could scope that down, because access to care is a really huge topic and we would need to really scope that down.

We were thinking it might be something along the lines of very foundational, almost thinking piece, what are the types of access?

Looking at it through a rural lens, what would be important to consider? What are the pros and

cons? What are the things to be aware of? That sort of thing.

Again, none of that is yet set in stone, but if you have any ideas, we'd be very much appreciative. So, with that, I'm going to stop and, I guess, hand it back. Or I can facilitate, however you want to do it.

CO-CHAIR PINCUS: I think, right now, we're going to open it up for discussion on some of these sort of key issues. And so, one question I had, well, a couple of questions I had, just to kick things off, was in one of the earlier slides, you mentioned using some of the Patient-Centered Medical Home measures.

So, are you including structural kind of measures in here too, in terms of the thinking about this? Is it specific though to the thing that CMS is doing? And also, when you're looking at, asking about suggestions around measures, are you talking about actual endorsed measures or are you thinking about measure concepts?

MS. JOHNSON: So, your second question

first, measures right now, because if we identified a core set, it would be ones that we conceivably could put in use immediately. We could certainly, and we did a couple years ago, talk a little bit more about concepts.

And I think we would get there more in our gaps discussions that we would have. In terms of PCH kinds of measures, right now, we have limited to NQF endorsed. But we certainly are open to structural measures.

MEMBER BINDER: Thank you. I'm really pleased to hear about this work and I congratulate you on it. I actually, myself, worked in a rural hospital for eight years and came from Maine, so I have some familiarity and am excited about this.

I have a couple things to recommend.

One is the framing of it, around the concept that rural people deserve the same quality of healthcare as anybody else. I think that's just an important kind of framework for it as opposed to saying, we're going to come up with a whole

set of measures that are different for rural communities than urban.

They're people and we want them all to get high quality care. And I would say, in terms of measure selection, to try to avoid saying, we have a separate set of criteria for rural populations than we do for urban.

I would just say, there are certain measures that have to be adapted, because of the small numbers. But otherwise, we select measures according to the same criteria we use for everything else.

That would be my suggestion, simply because, otherwise, I don't understand why we're pulling them out with different criteria, that doesn't make any sense.

Other than, of course, that key issue of the small numbers in terms of the calculations and needing to adapt and come up with measures that might be acceptable, given that the small numbers don't allow for certain outcomes to be tracked.

So, to somebody else's point about structural measures, are often a substitute or something like that. So, I would suggest thinking of it as an adaptation of current measures.

And then, the other thing that I think might be worthwhile for the workgroup to think about too is, how rural communities can actually do things better?

I don't know that that's come up in your group, but we, in the rural community that I was part of, we did some very interesting projects that would have been impossible in urban settings. And they were instructive.

So, an example would be, we decided we needed to address domestic violence in a more coordinated way, so we had one meeting and we pulled together the sheriff's office, a judge came, the emergency room department director for the only hospital, and physicians, the shelter staff.

It was like everybody who had

something touched the lives of victims of domestic violence, and the victims themselves came. You can't do that in an urban community. You can't bring in kind of all the key people all at once, usually.

But we could and they did some very interesting changes. And it really, really had a significant impact, I think, on the well-being of the victims.

so, I mean, and that's just one example, but I think that there's ways that you can look, especially when looking at population health as it intersects with acute care, where I think rural communities have done some extraordinary work and there may be areas to build on those strengths.

MEMBER MACKAY: Karen, I just wanted to say, thank you so much for this work. I echo previous comments that it's exciting to see rural communities being brought into this shift towards value.

It's understandable why a lot of times

these providers have been excluded from some of the programs that we've discussed in previous days.

It's hard to live in these communities often, it's difficult to practice medicine, but I do think that we may have been missing some opportunities, particularly because rural providers are oftentimes more familiar with or better at coordinating care and doing some of these more innovative community-gathering exercises or initiatives, like Leah mentioned.

So, it's just great to see. A couple of things that I wanted to mention as you're thinking about measure selection, is just getting more data in the pipeline, that will eventually - about the quality of care provided and provider performance, that will eventually trickle down to patients and families in these communities.

I think easily accessible information for rural residents, even in cases where they might not be able to use that information to shop for care, is nonetheless important and it is a

critical part of their ability to successfully and meaningfully partner with their providers in care and achieve the outcomes that we're all looking for.

I also think that, sort of echoing what Chris mentioned earlier, lack of information about the quality of care that is often provided in these areas may contribute to the incorrect assumption that -- or I screwed up that first part.

But not having enough data sometimes leads us to the incorrect conclusion that high quality care is not being provided. I think that can be dangerous and could drive a further outmigration of patients, because they think that their local providers -- or they aren't aware of the care that their local providers are providing.

And then, lastly, just that, it's my understanding that healthcare providers are often economic engines of these rural communities and they mean a lot more to the residents of the

communities than just the ability to access care.

And so, I think that underscores our need to be selecting measures that are reflective of the things that these providers do so well, like care coordination. So, thank you very much.

CO-CHAIR PINCUS: Carl?

MEMBER SIRIO: Yes, I think, just to build on Leah's comments, and that is, as I drive across Route 80, I would like to think that the care I got in Cleveland than it might be in Topeka, Kansas and in some rural community inbetween, if I got into a car wreck. But I think we all know that it wouldn't be.

That having been said, it seems to me that some of the issues you've raised, Karen, with respect to the problems around small sample size are not generic to just the smaller populations, right?

So, my point is, to the degree that what you do and what you learn cross-fertilizes what I think has been a perennial question, and to some extent, a perennial pushback on the part

of at least the clinical community is the issue of sample size and, therefore, robust validity of the outcomes that are being assessed through measurement.

So, I think that the Task Force is a great idea, but I think it can actually help inform thinking that's actually broader than just the impacts on smaller communities.

CO-CHAIR PINCUS: Giff?

MEMBER GIFFORD: Yes, there's, I think, over 80 percent of the counties in the country have a nursing home. So, we have a lot of small -- and even in larger, so we have a small sample size. And I -- they always want to be involved in the programs and everything, until they get involved in the programs.

One thing, and it's a little bit in the weeds, as you think about the small sample size, don't -- we've suggested a few times to CMS that they just expand the time window for the measures, but they don't feel that they can have measure specs for the same group with different

specifications.

I -- you think about -- you may have to do that. And I know there's -- it depends on what OGC says about statutory issues and everything.

Second, your burden issue becomes very different. So, we've calculated in some of the VBP programs that our rural people really want to get involved with, they discover that, just through sensitivity analysis, you're talking about either losing \$50 or getting \$500, if they play in the program.

And it doesn't require much burden to suddenly make participation in that program more expensive than \$500. And that's all they're going to get by playing in the program.

So, I didn't hear the burden aspect in that and so, you may want to think about that burden, help point that out. Because when we point that out to them, they suddenly are not as eager to be in the programs any more.

CO-CHAIR PINCUS: I just might add,

given Chris's comments earlier, you might want to add to your slide about specific issues, the issue of stigma to that as well. Sam?

MEMBER LIN: Thank you, Harold. And,
Karen, thank you so much for your presentation.

I heard you three times in December and I have to
say, just your presentation alone has advanced in
the last month. So, thank you.

My contribution to the, I was rural when rural wasn't cool, was my time with Indian Health Service, 100 miles from Spokane. And talk about rural and isolated.

That sort of leads to my question, you indicated you had three federal liaisons and these are supposed to be publicly reported programs, and I'm wondering, if you don't, you might think about having someone from Indian Health Service there, for Indian Health Service's sake, not for anybody else's in all honesty, because they still have the problems of access, the problems -- et cetera, et cetera.

The issue of, what do we do about the

rural, goes back to a story, at least I recall, about a former Army Surgeon General years back, when he was asked what he thought about this whole issue of maldistribution of physicians in this country and he says, what maldistribution?

And they said, well, sir, you obvious know about the lack of physicians in this country. He said, there's no maldistribution of physicians, if those farmers would move to the cities, we'd be okay.

(Laughter.)

MEMBER LIN: So, anyway, they're not going to move the city, so therefore, we do have to put them in high consideration. But I really think Indian Health Service needs a lot of help and this is at least a start to get them included in what we'll call the diversity in disadvantaged activity that we here have been pushing for the last two years.

CO-CHAIR PINCUS: Do we have any comments from people on the phone?

MEMBER ANTONELLI: Harold, Rich

1	Antonelli would like to get in the queue please.
2	CO-CHAIR PINCUS: Okay. Sure, Rich.
3	MEMBER ANTONELLI: Is it okay to jump
4	in?
5	CO-CHAIR PINCUS: Yes, speak up, Rich.
6	MEMBER ANTONELLI: Okay. So, Karen,
7	this is Rich Antonelli, Medical Director of
8	Integrated Care in the rural place otherwise
9	known as Boston, but I just, first of all, want
10	to applaud your approach.
11	Second, a lot of the work we're doing
12	nationally in integrated care and care
13	coordination, and be mindful that that work is
14	embedded in other workgroups, is actually
15	strongly focused on quality measures, including
16	experience and the A-word, access, for rural
17	areas.
18	Some of our most profound work,
19	actually, is being done right now up in Alaska,
20	leveraging the pretty precious resources of
21	community health workers.
22	So, that said, my suggestion is as

follows. Every quality measure that we look at across the MAP and across the workgroups, as far as I'm concerned, somebody should make the argument why it wouldn't be appropriate for rural populations.

Second, I think that what I would find would be a tremendous value-add from the folks thinking about rural health are the folks thinking about, okay, what would be some of those unique challenges?

I also think there are access issues in urban environments that would be different logistically and tactically than rural environments, but the access measures, so utilization of telemedicine, for example, care coordination and implementation, et cetera.

So, I guess I would encourage your workgroup to try as hard as you possibly can to think about the core measures at the broadest level, and really challenging to make the case for why we would need any additional measures for this going forward. So, thank you for your

commentary, I look forward to hearing what comes 1 2 from your workgroup. CO-CHAIR PINCUS: Other comments on the 3 phone? Derek? 4 MEMBER ROBINSON: I was just going to 5 make a brief comment. I appreciate the work that 6 7 Karen and the workgroup is doing. I think it's very important and 8 9 certainly, I echo some of the comments around the importance of individuals in rural communities 10 having access to the same high quality care as 11 12 other areas. And I think it's important to have 13 good transparency, as well, from a quality 14 performance perspective. My perception, that historically, the 15 16 challenges that come into play when quality and 17 performance is tied to reimbursement and the risk 18 for penalization or, perhaps, disruption of other 19 supplemental payments that may be going to those

providers. And so, I think that's something

that's important for the workgroup to consider.

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Washington DC

I think the, my perception, I think

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the excitement about getting engaged in this, having been not part of much of the value-based payment arena or progress over the last few years, is very positive to hear.

But I think also, as we get into tying that performance to payment, is really sort of where the rubber hits the road and whether we're able to keep that momentum in a way that folks feel is equitable and fair.

CO-CHAIR PINCUS: So, I had a comment, a couple of comments. One is, probably among the biggest issues into the areas in terms of access in rural areas is access to behavioral health services. And it's something that I think is important for the group, obviously, to look at.

But it also raises the issue of attribution, because there are very few psychiatrists/psychologists. I can't remember, there's some statistic about something like 50 percent of rural healthcare areas have no psychiatrist or psychologist available at all.

And so, who's responsible for that?

It's -- and that's going to come, not just in behavioral health, but it's something to think about in terms of sort of attribution for access in those situations.

And just one other thing is just that, we have a Harkness Fellow in my office who is funded by the Commonwealth Fund, fellows coming from other countries, soon to be career people.

And we have one from France who's actually doing a paper, one of her projects is doing a paper on comparing access to behavioral health services in France versus the U.S. And so, she might be somebody worth talking to.

Other comments, questions?

MS. BUCHANAN: Harold, we received one chat question, asking how geographic isolation is being handled by the workgroup?

MS. JOHNSON: So, I'll answer the question about geographic isolation. I think, in terms of what we're doing in this particular project, in terms of core measures, we're definitely thinking about geographic isolation,

since it is a huge challenge.

We'll have to think about that, when they identify the core measures. So, that will just be one of the things that we think about. Probably, it'll come in a little bit more -- like one of -- what we did yesterday is basically give a list of measures. Here's a strawman, here's a first cut at core measures.

What we're asking the workgroup to do over the next few days is to look much more critically at that first cut and tell us things about, okay, based on your perceptions, is this really feasible to collect these data used for these measures in rural areas? And if not, then maybe it doesn't belong on a core set.

Other things like, are there unintended consequences that this measure may have from a rural viewpoint? And I think that might be a place where the isolation comes in.

And eventually, once we come to a pretty good set of core measures, we need to look at the set itself as a whole and start thinking

about, does that really reflect quality for rural providers, as well as we can, knowing that there would probably be optional measures later on that would come in.

But I think looking at things kind of more as a set will also be helpful, but that comes a little bit later in the process.

Just a couple of things that I wanted to mention, and I probably won't hit everything, but part of the idea of a core set, and I apologize if I've given the impression, the core set of measures for rural providers, if it's ever taken up by CMS and used, the idea isn't necessarily that it would be a completely different set of measures used by everybody else.

As a matter of fact, the ideas of alignment with other programs and with other providers is really very important, because you want to be able to compare, if you can, rural quality to their non-rural counterparts. Again, many are very proud of the quality that they provide.

And so, the core set hopefully would be something that there would be a lot of intersections. It might not be complete overlap, but there would be a lot of intersections.

Going back to the idea of the access measures, I think attribution is absolutely something that we're going to have to hit head-on. A lot of access measures really are more population-based measure.

So, if you think about, okay, what does that do? Does that put our community into a -- can we really -- is our core community, or our whatever, fill-in-the-blank community, is that a reasonable thing to do that you might hold somebody accountable for?

Those are very important issues that we're going to have to address, I think. There were probably a few other things, and I apologize, that I've kind of lost. But I'm happy to take any questions or any advice that you have going forward.

CO-CHAIR PINCUS: Well, thank you. So,

1	I guess that concludes our agenda. So, we'll
2	just hear from the public?
3	MS. O'ROURKE: I think, yes, if we
4	could do one more public comment.
5	CO-CHAIR PINCUS: So, anybody in the
6	room from the public wish to make a comment? I
7	don't see anybody. So, anybody on the phone wish
8	to make a public comment?
9	OPERATOR: At this time, if you would
10	like to make a comment, please press star, then
11	the number 1. And there are no public comments
12	at this time.
13	CO-CHAIR PINCUS: So, I guess we're
14	ready to adjourn? We have lunch there.
15	Let me just really thank and really
16	express my appreciation to Chip, my partner in
17	this. And to NQF staff and the CMS staff and,
18	certainly, to everybody on the MAP Coordinating
19	Committee. So, thank you all.
20	MS. O'ROURKE: Yes, thank you. A
21	special thank you to Harold and Chip for all of
22	your leadership not just over the past two days

but all the work we do to get here. To Kate and her team for all their partnership and working with us through this.

And to all of you for bearing with us for the past two days and all the homework that I know we send you and the short time to prepare, we really appreciate it. Thank you especially for all the feedback on what we could do better next year.

I think we'll be following up about how we can make that happen and if we can convene a small group of you or if people want to send thoughts offline, if anything comes to you on the plane ride home, we'd love to hear it.

I did just want to briefly cover some, just what we'll do to wrap up pre-rulemaking. We will start issuing the reports of MAP's recommendations in the next few days, really. So, February 1, the spreadsheet of all the individual measures under consideration and MAP's recommendation will be published.

February 15, I will release the

guidance for hospital and PAC LTC programs. And then, March 15 will be the guidance for clinician programs.

I did want to highlight, we will be back in touch in the coming months. There will be a web meeting in August of the Coordinating Committee to review the input from the Medicaid and Rural Task Forces. So, keep an eye out on that and we'll be getting that on your calendars in the coming weeks. Giff?

MEMBER GIFFORD: Follow-up for Harold and Chip, I think yesterday it was suggested tracking historically stuff that's come through the MUC, whether it got in rules and what -- would that -- how does that fit in that time line? I mean, it's not urgent, but do we wait to August to have that?

CO-CHAIR PINCUS: I think our next official meeting is a webinar in August. But there is certainly the possibility of being a communication before then. And certainly, the idea of this sort process workgroup would

probably be some phone calls before that.

MS. O'ROURKE: Yes. And I think that's something, if you have thoughts about what information is useful or would be helpful, what intervals and formats you'd like us to provide that information, let us know. Because I think that's on our list of how we can bring you better information for next year.

So, if you have ideas, let us know and we can follow up on that. Otherwise, again, thank you for all of your time over these past weeks and past two days. We know they were long meetings, so thank you. And we look forward to seeing you virtually in August. So, thank you very much, everyone.

(Whereupon, the above-entitled matter went off the record at 12:46 p.m.)

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## <u>C E R T I F I C A T E</u>

This is to certify that the foregoing transcript

In the matter of: Measure Applications Partnership

Coordinating Committee Meeting

Before: NQF

Date: 01-26-18

Place: Washington, DC

was duly recorded and accurately transcribed under my direction; further, that said transcript is a true and accurate record of the proceedings.

Court Reporter

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