Microbiology Expert Committee (MEC) Meeting Summary

March 12, 2019

1. Roll Call and Minutes:

Robin Cook, Chair, called the meeting to order at 1:30pm Eastern by teleconference on March 12, 2019. Attendance is recorded in Attachment A – there were 8 members present. Associate Members: Elisa Snyder and Hunter Adams.

Robin reviewed the following sets of minutes with the committee: December 11, 2018 January 29, 2019 February 19, 2019

A motion was made by Jessica to approve the minutes from 12/11/18, 1/29/19 and 2/19/19 as written. The motion was seconded by Michael Blades and the vote will be completed by email.

(Addition: The following votes were received by email:

Cody 3/13/19 - For Christabel 3/13/19 - For Vanessa 3/13/19 - For Jody - 3/13/19 - For Enoma - 3/13/19 - For Deb Waller - 4-12-19 - For Lew - 3/12/19 - For Gary - 4/10/19 - For Jessica 3/12/19 - For Mike 3/12/19 - For Michael Carpinona 3/13/19 - For

The motion was approved.)

2. Standard Interpretation Request Summary

Robin started reviewing the SIR Summary Table that was recently sent to all Expert Committee Chairs. She added her notes to the comments column to figure out whether the SIR was already addressed in the 2016 Standard.

She went through each SIR with the attendees and updated the table using Webex as she worked through the table (Attachment D – only the 5 columns the Committee worked on

are in this attachment). The entire table was sent to Committee and Associate members for review and comment.

3. SIR 301

The committee reviewed the SIR and decided on the following language to vote on:

The requirement is every 10 samples. This would apply to every filter set up on the manifold if there were more than one being used. There is no intent to do a blank in the middle of a serial dilution series for any given sample as may be the case if it were to be done every 10 plates.

A motion was made by Michael C to approve the above language in response to SIR 301. The motion was seconded by Cody. There was no further discussion. The motion passed unanimously.

(Addition: Robin distributed the language and voting results to all Committee members and a concern was raised. The language was updated and will be voted on again during the April meeting.)

4. Action Items

See Attachments B and C for updates to action items.

5. New Business

None.

6. Next Meeting and Close

The next meeting will be held by teleconference on April 9, 2019 at 1:30pm Eastern.

A summary of action items and backburner/reminder items can be found in Attachment B and C.

Robin adjourned the meeting at 3:01pm Eastern. (Motion: Jessica Second: Mary Unanimously approved.)

Attachment A

Members	Affiliation	Balance	Contact Information
Robin Cook	City of Daytona Beach	Lab	cookr@codb.us
(Chair) (2019)	EML		
Present			
Michael Carpinona	NJ DEP	AB	Michael.Carpinona@dep.nj.gov
(2022*)			
Present			
Ron Coss	Orange County Sanitation	Lab	RCoss@OCSD.COM
(2022*)	District		
Absent			
Cody Danielson	Oklahoma	AB	Cody.Danielson@deq.ok.gov
(2022*)			
Present			
Lew Denny	Flowers Chemical	Lab	lewdenny@comcast.net
(2021*)	Laboratories – North		
Absent			
Jessica Hoch	TCEQ	Other	Jessica.Hoch@Tceq.Texas.Gov
(2019*)			
Present			
Lily Giles	Louisiana	AB	Lily.Giles@LA.GOV
(2022*)			
Present			
Mary Robinson	Indiana	AB	mrobinson@isdh.IN.gov
(2022*)			\bigcirc \mathbf{c}
Present			
Michael Blades	ERA	Other	mblades@eraqc.com
(2021*)			U I
Present			
Jody Frymire	IDEXX	Other	Jody-Frymire@idexx.com
(2022*)			
Present			
Kasey Raley	Eurofins Eaton	Lab	KaseyRaley@eurofinsUS.com
(Vice-chair) (2020*)	Analytical, Inc.		5 50
Absent	5		
Vanessa Soto Contreras	Florida DOH	AB	Vanessa.SotoContreras@flhealth.gov
(2020*)			\bigcirc \bullet
Absent			
Gary Yakub	Environmental Standards,	Other	gyakub@envstd.com
(2020)	Inc.		
Absent			
Enoma Omoregie	NYCDEP	Other	eomoregie@health.nyc.gov
(2021*)			
Absent			
Christabel Monteiro	ESC	Lab	cmonteiro@esclabsciences.com
(2021*)			Ŭ Ū
Absent			
Ilona Taunton	The NELAC Institute	n/a	Ilona.taunton@nelac-institute.org
(Program Administrator)			b
Present			

Participants Microbiology Expert Committee (MEC)

Attachment B

Action Items – MEC

			Expected	Actual
	Action Item	Who	Completion	Completion
1	Review Method codes and send comments to Robin for Dan Hickman.	Deb	TBD	
19	Provide EPA interpretation on temperature readings to Ilona. She will have it posted on the website.	Robin	1/31/14	
74	Send questions for ABs regarding method codes to Robin.	ALL	3/15/18	
76	Provide an update on what has been done with the databases after Jennifer's review and internal EPA meetings.	Jennifer	4/10/18	
78	Forward link to PDFs on DW website with rule, method and analyte information.	Jennifer	3/31/18	
81	Addition: Forward response to SIR 331 to Lynn Bradley.	Robin	11/13/18	
83	Send out resumes for all applicants to the committee.	Robin	12/10/18	Send before 1/8/19.
84	Send out copy of Charter.	Robin/Ilona	12/10/18	
85	Send out updated Technical Director Language	Deb	1/8/19 or week before Milwaukee	
86				
87				

Attachment C

Backburner	/	Reminders – M	EC
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	Item	Meeting Reference	Comments
1	Update charter (if needed) in November 2018.	n/a	Ongoing

Attachment D - SIR Status Table

	Comment	Applicable	Applicable	Applicable	Addressed/Clarified in
#	666	to 2003	to 2009	to 2016	2016 Standard
27	This language is virtually unchanged in the 2009 and 2016 standards. The SIR is still valid. Some reorganization of the QC section was done in the 2016, addressing the basic premise of the response.	Yes	Yes	No	Clarified in the 2016 Standard by adding in 1.7.3.1.a) "All materials and supplies that would be needed to process"
42	This language is virtually unchanged in the 2009 and 2016 standards. The SIR is still valid. The basic issue with this SIR is that until the 2016 standard it was not made clear that sterility checks and method blanks are different things required for different reasons. Section 1.7.3.1 is about sterility checks, but this question is about method blanks.	Yes	Yes	No	See section 1.7.3.2 for method blanks. See SIR 301 as well.
44	This language is virtually unchanged in the 2009 and 2016 standards. The SIR is still valid.	No	No	No	This question applies to the method and/or program, not the standard.
97	This language is virtually unchanged in the 2009 and 2016 standards. The 2016 standard added in the PV= nRT language and called it sufficient. The SIR is still valid.	Yes	No	No	Clarified in 2009 as note and added to body in 2016.

#	Comment	Applicable to 2003	Applicable to 2009	Applicable to 2016	Addressed/Clarified in 2016 Standard
98	As this is something that is somewhat static and the source is not changing over time as an In- house DI system may be an initial check that mirrors the DI water requirements is reasonable with some relief regarding frequency as noted in 1.7.3.1.d) v. Refer to 1.7.3.1 e) REVIEW FOR NEXT REVISION OF STANDARD	N/A	Yes	Yes	clarified in 1.7.3.1.d.v. Specified what and when for purchased water.
132		N/A	Yes	Yes	clarified in 1.7.3.1.d.v. Specified what and when for purchased water.
135		Yes	No	No	This requirement comes for Chp 5 of 2003 NELAC, That requirement was removed from V2 in 2009 std and remains as such.
138		N/A	Yes	No	Definition of source added in 1.3.1 of the standard. The 3rd question is a business decision as to how a lab would like to prove the bottle came from their laboratory and therefore not a question of clarity with regard to the standard.
285		N/A	Yes	No	clarified in 1.7.3.3 to include any quantitative method.
301	The requirement is every 10 samples. This would apply to every filter set up on the manifold if there were more than one being used. There is no intent to do a blank in the middle of a serial dilution series for any given sample as may be the case if it were to be done every 10 plates.			No	standard says plates

#	Comment	Applicable to 2003	Applicable to 2009	Applicable to 2016	Addressed/Clarified in 2016 Standard
331	The standard requires that the labs generate QC on site as indicated by the words "The Laboratory shall" What has been described in both of these instances is no different that a certificate of analysis/purity or the like. While the standard requires the lab to have those certificates on file, it also requires the lab to verify both sterility and performance. It has always been assumed the lab should be doing this themselves and many assessors used some parts in the body of Chp 5 for 2003 and V2 for 2009, but it was not super clear. So the committee made the decision to add, "The laboratory shall" to the appropriate section to help clarify that intent.		Yes	No	Clarified by adding the words "The laboratory shall"
338	See SIR 331.		Yes	No	Clarified by adding the words "The laboratory shall"