

**Minutes of
THE NELAC INSTITUTE'S
Whole Effluent Toxicity Testing (WET)
Subcommittee for Acceptance criteria
March 23, 2011**

The meeting was called the order at 1:00pm EST by RaeAnn Haynes.
Members present are listed in Attachment B.

This group has been on hiatus due to the lack of a Chair for over one month. The purpose of the meeting was to review our status and what direction we need to move toward.

Two thirds of the Proficiency Test Providers have submitted their data. Jeff Lowry agreed to contact the remaining Provider for the data. The group was directed to the current WET Table that is posted on the TNI website for a better understanding of the data the is currently available and being used by the PT Providers.

Theresa Norberg-King has agreed to present the group with an WET 101 paper to further the understanding of this test. She also commented on the Tables use of both MHSF and DMW matrices. EPA views these as no different and any time we can reduce the number of PT test available to a relatively small group of laboratories, our statistical results will be more robust. Finally Theresa is interested in collecting more data from the laboratories including conductivity and pH. RaeAnn Haynes agreed to contact both the PT executive Committee and the AB Committee for a discussion about how we collect data and how the AB view data collected that may not have a score (Acceptable/Not Acceptable) associated with the data.

The group agreed that we need to finish walking through the comment Table (see Appendix C) since this will help scope our current direction with review and improvement of the WET FoPT Table for use by both the laboratories and the AB's.

Our next teleconference will return to the regular meeting time of April 13th at 1:00pm EDT.

Attachment A

ACTION ITEMS
TNI PROFICIENCY TESTING COMMITTEE

Item	Action Item	Assigned To	Due Date	Date Complete
1	Keep the PT Board informed	RaeAnn Haynes	On-going	
2	Send original comments to this subcommittee for evaluation	Stacie Metzler	1/12/2011	Not Complete
3	Request data from PT Providers for evaluation	RaeAnn Haynes	1/12/2011	2/5/2011
4	A summary for non-biologists about the stream-lined WET test proposed by EPA in 2002.	Teresa Norberg-King	2/9/2011	
5				

Attachment B

TNI WET FOPT SUBCOMMITTEE MEMBERS 2010-2011

Member	Affiliation	Email	Phone	Category	Present on the Call Y/N
Raeann Haynes (Chair)	Oregon DEQ	Haynes.raeann@deq.state.or.us	503-693-5757	AB	Y
Stacie Metzler	Hampton Roads Sanitation District	smetzler@hrsd.com	757-460-4217	Lab	N
Ginger Briggs	Bio-Analytical Laboratories	bioanalytical@wildblue.net	318-745-2772	Lab	N
Pete De Lisle	Coastal Bioanalysts Inc	pfd@coastalbio.com	804-694-8285	Lab	N
Robert Kelley	ETT Environmental Inc	bobkelley@ettenvironmental.com	864-877-6342	Lab	Y
Jeff Lowry	Environmental Resources Assoc.	jlowry@eraqc.com	303-431-8454	Other (PTP)	Y
Jamie Mitchell	Hampton Roads Sanitation District	jmitchell@hrsd.com	757-460-4220	Lab	N
Rami Naddy	AE Com	rami.naddy@aecom.com		Lab	Y
Bob O'Brien	RT Corporation	bobrien@rt_corp.com	307-742-5452	Other (PTP)	Y
Faust Parker	PBS&J Environmental Toxicology Lab	frparker@pbsj.com	713-977-1500	Lab	Y
Steven Rewa	Environmental Resources Management	steven.rewa@erm.com	616-738-7324	Lab	Y
Chuck Wibby	Wibby Associates	cwibby@wibby.com	303-940-0033	Other (PTP)	N
Brian Krausz	USEPA	krausz.brian@epa.gov	202-564-2970	Other (EPA)	N
Teresa Norberg-King	USEPA	Norberg-King.Teresa@epamail.epa.gov		Other (EPA)	Y

Attachment C

Comment	Response
The bulk of historic data is based on pre-NELAC labs: is it reliable?	What's the background for the pre-NELAC? Used EPA Tables to generate current NELAC Table. Methods were changed in 2002. Or methods were streamlined but not so much to affect accuracy and precision? Cannot get rid NOEC for the permits, point estimates are necessary. PT study needs a point estimate. However, all NOEC tests include an IC25. Seems appropriate to use an endpoint test rather than a point estimate hypothesis result.
Restrict endpoints to IC25 (reproduction or growth) and EC50 (survival) and do away with NOEC endpoints.	PMSD no data for PTP's, has established limits. Limits not hard. Can be used to repeat tests. May not reflect lab performance. Samples different than controls. No toxicity then no limits for PMSD. If we don't require NOEC then the PMSD is not necessary. 2 SD vs 3 SD? Answer by looking at the data? May depend on the test and study size.
Have each field (test type / species) have two criteria - the IC25/EC50 for "accuracy" and the PMSD for "precision". Both would be reported and a lab would have to fail both for the result to be considered a failure.	
The limits for the IC25 and EC50 should be the 99% confidence interval and the limits for the PMSD would be the upper limit only as defined in the EPA Method Variability document for toxicity. I don't think the lower limit should apply in as much as it is not a bad thing if the lab has very precise results.	
That the WET FOPT table be improvised and easier to understand	
That the WET PT studies be modified to better reflect a laboratory's performance, either by modifying the endpoint (i.e. required reporting of IC25 instead of NOEC) or making AB bodies understand that this is a monitoring tool and not to put so much emphasis on it.	The NOEC test uses the IC25 and the two results are normalized by the control. Do we really need two results for the same test?
NOEC endpoints should not be used and only point estimates (i.e. IC25 or LC50) should be used.	The IC25 is an endpoint. The NOEC is more like a DL (detection limit).
The TNI criteria need to work with the DMRQA criteria. (But I don't see why they have to be same).	

Attachment C

Use of 95% probability limits for TNI WET limits is probably not a good idea. TNI chemistry wastewater limits are based on 99% probability values.

These really need to talked about in terms of standard deviations. EPA has stated +/- 2 std dev is appropriate for WET however, the data evaluation will show which is most reliable.

There are multiple variables in WET testing to begin with (live organisms, which have their own set of issues like age, condition, sex, etc.; multiple technicians handling test and organisms every day; variations in test water between labs (both salt and fresh); different test vessel configurations and on and on. With PT we are adding the most significant variable of all, which is the preparation of the "neat" PT sample for use in the test. A relatively small error in preparing the simulated effluent probably explains most of the differences between lab results; in the Chronic test series there is a daily preparation of the PT sample, almost always by different lab techs (even when you try to have the same person prepare the sample). This step in the process is not part of the normal WET test protocol.

Losing Accreditation for one species could have major ramifications for all tests (either salt or fresh water). In Texas, permittees run either chronic or 48 hour acute tests, in addition to a 0 % & 100% acute. If a lab losses accreditation for any one method in these groups they will lose all clients that have that requirement. If you are a freshwater only lab you are out of business. Even if you run both fresh and salt your revenues could be cut in half with the same result. This is not appropriate for such a variable screening tool.

The reality is, that with the exception of the C. dubia tests, there just isn't any significant difference in the various methods. If you can run one method you can pretty much run them all. So assessing on the OVERALL performance of the lab on PT's (even across methods) makes a lot of sense. And we have to keep in mind that we are, in some, cases grading a method, within a method, that has the same potential for error and variability.

Attachment C

The dilution series used (.5) is inappropriate. A .75 series based on where the end point is projected would provide better data. In the old days, as I recall, we ran a range finder and then determined our own dilution series on the acute tests. An appropriate response curve would make much more sense for PT testing.

The end points being used are dumb! LC 50s are slightly better than point estimates. But if you do a literature search on the toxicity (LC 50's) of various toxicants you will find much variability and in some cases extreme variability. WET is a screening tool for complex effluents. We are only looking for a toxic response at some critical dilution. The agencies understand this variability and allow for additional confirmation showing that there is a REAL issue before holding the permittees feet to the fire.

Both IC25s and NOEC (i.e., hypothesis test) endpoints should be required for ALL short-term chronic WET studies for sublethal data (possibly for survival data as well). This should help evaluate response data and gain a better understanding of the variation of the given endpoint from all the testing labs. Currently, all labs may not be reporting both endpoints.

When reporting sublethal hypothesis test endpoints (i.e., NOEC, LOEC) require the reporting of the percent minimum significant difference (PMSD). This is a requirement for WET testing results and should be required for PT studies. This should help interpret / explain results among various laboratories and explain out-of-range results (as I mentioned on the previous call). It will also help evaluate the PMSD range among WET laboratories and ensure that invalid studies are not being reported.

Attachment C

Suggestion (based on our discussion): For hypothesis test endpoints include results at 95% and 99% statistical significance (i.e., $\alpha = 0.05$ & 0.01). Don't think we can exclude results at 95% significance because most WET dischargers with NOEC requirements have same limitation (i.e., they can't use 99% statistical significance for reporting purposes).

Suggestion: Provide a dataset to all labs to statistically analyze and report endpoints. This may help provide some (initial) feedback on variance associated with data interpretation. Data set should result in anomalous statistical results that would require interpretation of the NOEC endpoint (e.g., USEPA 2000).