

Stanley Tong - EPA Region 9

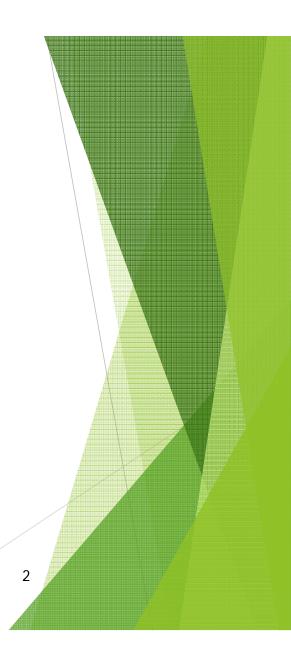
Michael Klein - NJDEP

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January 27, 2015

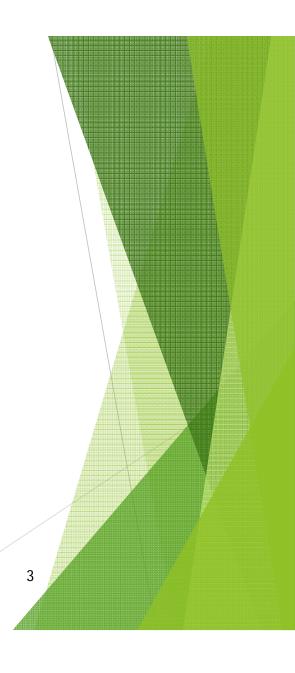
# **Topics**

- Updates
- ► Audit Sample Program Statistics
- Options for Expanding the Audit Sample Table
- Open discussion



# **Updates**

- ▶ 2011 EPA approved TNI's program
- ➤ 2013 Audit samples became available
- ► States signed up to access the program



## Updates (cont.)

- ► TNI's Stationary Source Audit Sample (SSAS) documents: 3 modules
- Vol 1 Mod 1 (Sample Providers) 3/2/14 (not yet submitted to EPA)
  - ▶ Manufacturing Lot: A group of audit samples made at one particular time in one particular place for one particular method at one particular concentration.
  - ▶ Provider shall not send the same an audit sample from the same manufacturing lot twice to the same Facility or Laboratory consecutively, or more than once in a calendar month, or more than eight (8) times in a twelve month period.

Reason for this change: running out of available audits

- Vol 1 Mod 1 (Sample Providers)
  - ► Providers shall supply audit samples that <u>are within</u> reflect the concentration ranges in the SSAS Table.

If requested by the Regulatory Agency and/or the Facility, ranges that are not listed in the SSAS Table may be included in an audit sample if the purpose and technical justification are documented, and if, where appropriate, the Regulatory Agency and/or Facility are notified in advance.

- Samples outside of the SSAS Table concentration range:
  - ▶NOTE: A Provider may, upon request, supply samples, similar in composition to audit samples, which have concentrations outside the ranges in the SSAS Table. By definition, such samples would not be considered audit samples, and are, therefore, outside the scope of this Standard.

- ► Providers shall contact the Regulatory Agency (RA) within 5 business days for any specific RA requests
  - ▶ e.g., changes to audit sample concentration/shipment address
- Providers assign audit samples based on lab-specific information at the time of the initial order
- ▶ Provider shall notify the RA within 2 business days if the Facility modifies audit sample <u>order</u> (including Tester and/or laboratory)

- Audit samples are laboratory specific
  - ▶ If a different lab than the one on the original order <u>submits</u> the audit results, the Facility shall inform the Provider prior to sending the audit sample to a new or alternate lab.
    - ▶ Provider shall:
      - Notify the Regulatory Agency within 2 business days for approval to submit the evaluation results.
      - Submit the results if no reply from the Regulatory Agency after 3 business days.

- Evaluation Report
  - ▶ Inadvertently deleted requirement for Provider to include unusual details of the audit sample (e.g., need to change an assigned value or delete an analyte from evaluation) (Section 11.2.1 i) will be restored)
- Regulatory Agency Responsibility
  - ► Clarified items RA should review during the 15 day period:
    - ► Evaluate method, container, matrix, analytes, requested value.
    - Notify Facility and Provider if sample must be changed so the Facility can modify its order with the Provider.
  - ► NEVER notify the Facility about a change in concentration

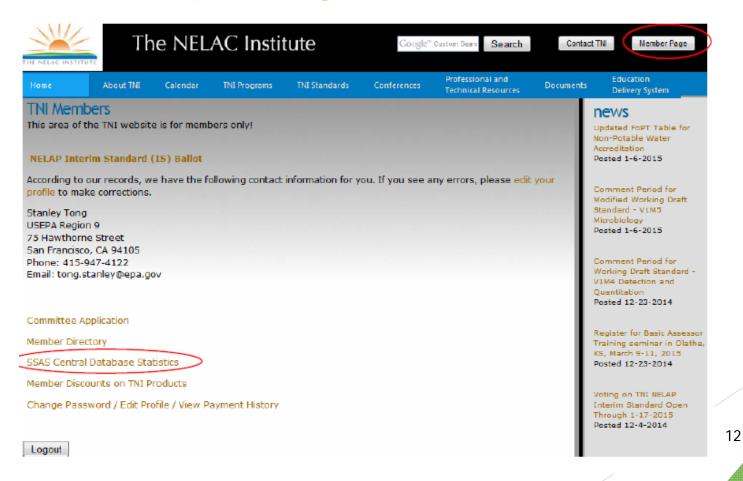
- Complaint Procedure (e.g., inaccurate audit sample value)
  - Provider include with reporting forms:
     Regulatory agency contact information
     Webpage link to submit a compliant
- Important that States respond ASAP to Provider's communications
  - ▶ i.e., audit sample order / audit sample order changes

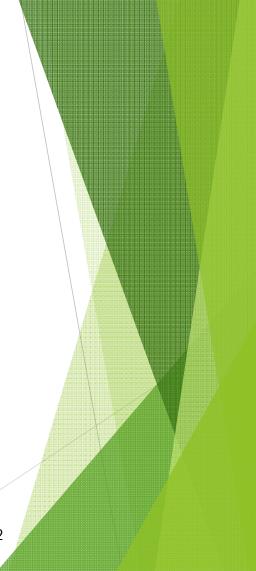
# Audit Sample Program Statistics

- ▶ Audit samples analyzed to date (~1000 samples ordered to date)
- ▶ Pass rate generally 90% or higher
- TNI members page statistics
  - ▶ <a href="http://www.nelac-institute.org/index.php">http://www.nelac-institute.org/index.php</a>
  - ► click on "member page" tab at top right

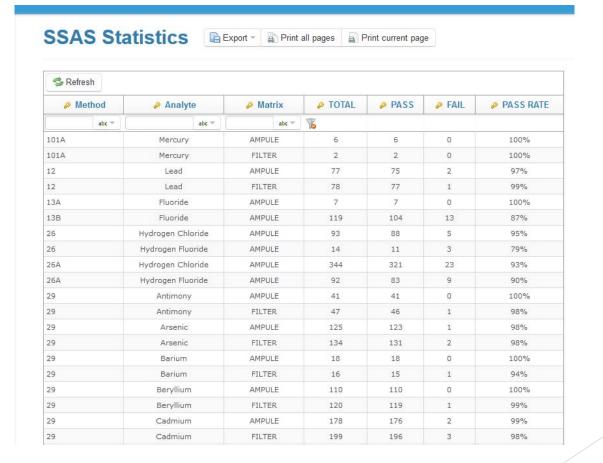


# Audit Sample Program Statistics (cont.)





Audit Sample Program Statistics (cont.)



# Audit Sample Program Statistics (cont.)\*

- ► Mth 8 Sulfuric Acid Mist: (120/147) 82%
- Mth 13b Fluoride: (104/120) 87%
- ► Mth 26 Hydrogen Fluoride (11/15) 73% (small sample size)
- Identifiable Trends?
- Program is working to find problems

<sup>\*</sup> As of 1/15/2015

# Options for Expanding the Audit Sample Table

- ▶ What we found:
  - ► Custom audit assigned values in the Central Database: 146
    - ▶ 141 of these were for lower concentrations
    - ▶ 112 were for Method 29
    - ▶ 15 failed (Cr, Pb, Ni, Cu, Ba, Zn, HCl, SO<sub>2</sub>) Sb, 6- M13b
  - ▶ What happens if results differ "significantly" from the assigned value?
  - ► Audit samples outside the Audit Sample Table acceptance range will no longer be provided.
    - ► How do we move forward?

# Options for Expanding the Audit Sample Table(cont.)

- Determining the acceptance criteria
  - ▶ 40 CFR 60.8 Based on "well established statistical methods"
  - ▶ Possible options to move forward: Temporary acceptance limit; reestablish after new historical data.
    - ▶ Extrapolate existing data on what labs can achieve; or
    - ▶ Temporary limit of 10x of Provider's repeatability limit
- Goal: Audit samples that challenge both the tester and the lab
  - ► Method 25Z
- ► TNI: Investigate Method 8 failure rate
- ▶ Need new members NJ, NC, EPA Region 9 termed out
  - ► http://www.nelac-institute.org/amember/signup.php
  - ► Generally meet every other Monday 2pm Eastern

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#### **Open Discussion**

- Reuse of residual excess audit sample as a QC sample
- ► Audit results that are beyond the acceptance range
  - ► Biased high
  - ► Biased Iow
- Review spreadsheet of low bias failures
  - ► EPA Region 9 investigation HF (Lab determined its cleanup cartridge caused low bias) / Ag (no regulatory requirement, results accepted)
  - ▶ Open forum to discuss how States investigated failures and the outcome
- ► Feedback for audit sample Providers

## Audit Sample Table - avail analytes/conc. ranges

- http://www.nelac-institute.org/index.php
- ► Click link to Stationary Source Audit Sample Program

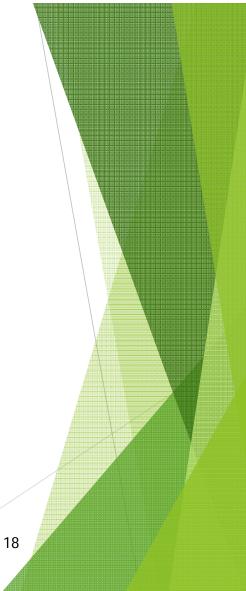


► Click link to Audit Sample Table



Click link to Stationary Source Audit Sample Table

Stationary Source Audit Sample (SSAS) Table
Current Table: Revision 5, Effective April 15, 2014 (PDF, 570KB)



## **Audit Sample Calculation Tool**

Three lines below the Audit Sample Table link is the link to the Audit Sample Calculation Tool - <a href="http://nelac-institute.org/ssas/calctool.php">http://nelac-institute.org/ssas/calctool.php</a>

Audit Sample Calculation Tool

Click here to access the Audit Sample Calculation Tool, used to calculate the concentrations needed when ordering audit samples.

AUDIT SAMPLE INFORMATION PER SECTION 4.1.1 OF VOLUME 1, MODULE 3 REQUIREMENTS FOR PARTICIPATION IN THE TNI STATIONARY SOURCE AUDIT SAMPLE PROGRAM

[CLICK FOR INSTRUCTIONS]	
TEST METHOD: Select Method ▼ MEDIA:	Select ▼
DATA ENTRY:	
Checkmark to include in order: (toggle all)	
Analyte:	
Emission Limit (lb/hr):	
Est. In-Stack Gas Concentration (mg/dscm):	
Stack Flow Rate (dscfm):	
Estimated or permitted as applicable  Sample Rate (m <sup>3</sup> /hr):	
Sample Time (hr):	
Final Diluted/Concentrated Volume per Method (mL): Enter if audit is in terms of mass/liquid sample volume.	
CALCULATE AUDIT CONCENTRATION Reset	
CALCULATED VALUES:	
Analyte Molecular Weight (g/mol):	

