

#### Proficiency Testing (PT) from the Point of View of the Accreditation Authority

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- Accrediting Authorities:
  - know the value of Proficiency Testing (PT) samples
  - determine the capability of the laboratories
  - want acceptable, consistent results
  - use samples with known target concentrations

- US Department of Agriculture
  - Pesticide Data Program (PDP)
  - All PDP laboratories analyzing water will participate in PT sets designed by MPO and administered by a selected commercial vendor.
  - Technical Director shall be responsible for overall monitoring of the proficiency of PDP laboratories.

- US Department of Agriculture
  - Microbiological Data Program (MDP)
  - Proficiency Testing (PT) samples will be analyzed according to the current MDP Semi-Annual Program Plan and the particular test protocol supplied by the USDA Monitoring Program Office (MPO)
  - PT samples prepared by the laboratory Quality Assurance Unit (QAU) and transferred to the Technical Program Manager (TPM) or designee for analysis

- US Environmental Protection Agency
  - PT vendors then send evaluations of the submitted data to the laboratory and any other designated certifying/accrediting authority.

- USEPA: 40 CFR 141.23 (k)(3)
  - Analyze Performance Evaluation (PE) samples provided by EPA, the State or by a third party (with the approval of the State or EPA) at least once a year.
  - For each contaminant that has been included in the PE sample and for each method for which the laboratory desires certification achieve quantitative results on the analyses that are within the following acceptance limits:

Contaminant	Acceptance Limit	
Antimony	$\pm$ 30 at	0.006 mg/L
Arsenic	$\pm$ 30 at	0.003 mg/L
Asbestos	2 standard deviations based on study statistics	
Barium	$\pm$ 15 at	0.15 mg/L
Beryllium	$\pm$ 15 at	0.001 mg/L
Cadmium	$\pm$ 20 at	0.002 mg/L
Chromium	$\pm$ 15 at	0.01 mg/L
Cyanide	$\pm$ 25 at	0.1 mg/L
Fluoride	$\pm$ 10 at	1 to 10 mg/L
Mercury	$\pm$ 30 at	0.0005 mg/L
Nickel	$\pm$ 15 at	0.01 mg/L
Nitrate	$\pm$ 10 at	0.4 mg/L
Nitrite	$\pm$ 15 at	0.4 mg/L
Selenium	$\pm$ 20 at	0.01 mg/L
Thallium	$\pm$ 30 at	0.002 mg/L

- Laboratories need some type of certification or accreditation to operate effectively in their particular "business arena."
- Customers of laboratories, which may be internal and/or external, want some way to know that the quality of analytical data is the best available from the laboratory they are using.

- Accreditation Authorities:
  - typically audit the laboratories
  - require acceptable performance on PT samples
  - administrative process that requires the laboratories
    - to describe their scope of accreditation
    - submit fees for the accreditation
    - pay for the performance of an on-site assessment

- Scope of Accreditation
  - very important in determining the expertise of the laboratory
  - may need to contact multiple Accrediting
    Authorities to be accredited in all areas
  - focus on drinking water analysis
    - Regulated pesticides
    - Regulated volatiles
    - Metals
    - Radiochemistry, etc.

- During the assessment, Accrediting Authorities:
  - will review performance on PT samples
  - look at the following related to the analysis of the PT samples:
    - Traceability
    - Calibration
    - Record keeping
    - Training

- PT provider
  - inserts the target quantity
  - upper and lower limits of acceptance are calculated
  - using the published regression constants
  - reports results to the laboratory
  - Reports results to the Accrediting Authority

- New methods and new analytes
  - categorized as experimental tests
  - use at least one year's experimental data to establish acceptance criteria constants
  - method includes multiple analytes
    - produce acceptable results for 80%
    - a sample that contains 10 to 14 analytes the laboratory must not produce failing results for more than two analytes

- Corrective action process is required when performance does not meet the criterion
  - analyze supplemental PT samples
  - investigates the cause of the failing result
  - minimum amount of time between failed PT results and the make-up PT.
  - avoid loss of accreditation for an analyte or a method

- Performance criteria are prescribed in federal statutes
  - overlay on top of the criteria determined by the accreditation authority
  - EPA method for volatile organics (524.2) includes criteria for acceptability of quality control sample recoveries at  $\pm$  30% recovery
  - internal standard, surrogate standard and laboratory-fortified blank are all judged based on this 30% window

- Performance criteria (cont'd)
  - 40 CFR 141.24 (f) (17) requires that volatile organic PT samples must fall within +/- 20% of the target
  - when the target concentrations are 0.010 mg/L
  - 40% when the targets are 0.010 mg/L
  - can be "in control" according to the method but fail the PT sample when the recovery is 121%

- Availability of PT Samples
  - not all matrices
  - not all analytes
  - certificate does not indicate whether or not
    PT samples were analyzed
  - laboratory user is unsure of which methods used PT samples and which ones did not
  - without PT samples the assessment may need to be more detailed

- Pt Results are not the only "ruler" to use for accreditation
  - Quality systems
  - Training
  - Equipment
  - Record-keeping
  - Documentation
  - Onsite assessments
    - are all used by Accrediting Authorities to make the final decision

