

NEMC 2021

Implementing ISO 17025 at an Oil Refinery

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Introduction

- Quality Framework
- Describe Monitoring System
- Organization of Project
- Tasks performed
 - How
 - By who
- QA and validation





ISO 17025

- General requirements for competence for testing and calibration laboratories
 - Covers:
 - Standard methods
 - Non-standard methods
 - Laboratory developed methods
 - Traceability to Primary Standard (NIST)
 - Execute method in a consistent manner
 - Continually improve

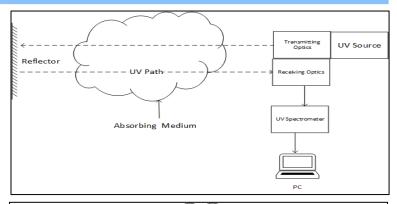


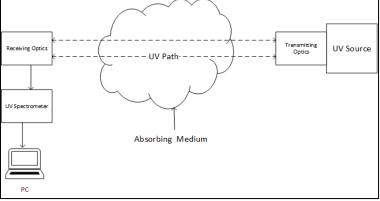




UV Method

- The system should be capable of making spectral absorption measurements along an open-air optical path.
- The system must be able to produce and save a single beam spectrum.
- The system must be able to operate at 0.14 nm wavenumber resolution over the range 185 to 300 nm.
- The system must be capable of acquiring data by co-adding individual, single-beam scans in single-scan increments. At a minimum, the system must be able to co-add single beam spectrums, so that a five-minute average can be obtained.
- The system must have a mechanism where a gas cell of a known concentration can be installed in the UV path, so that the entire beam passes through the cell.

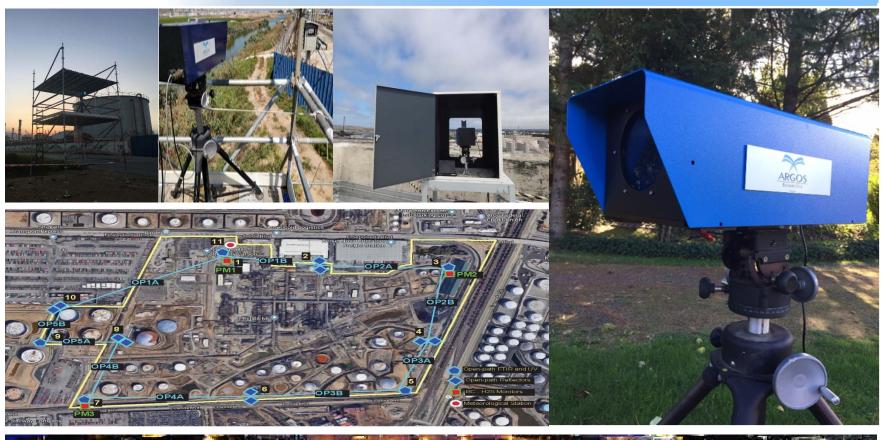








Open Path System: Field Laboratory

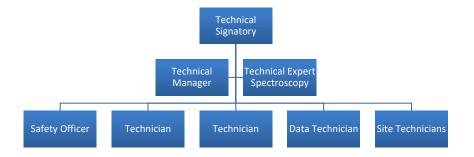






Project Organization

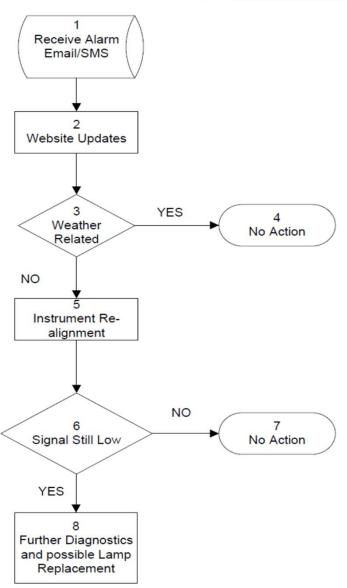
- Make sure the system is operating within its parameters (Level 0).
- Validate system detects and perform single point QA checks (Level 1).
- Validate the method (US EPA 301) (Level 2).
- Validate the data (Level 2)
- Service and maintain the equipment (Level 3).
- Conduct final Data Validation (Level 3).

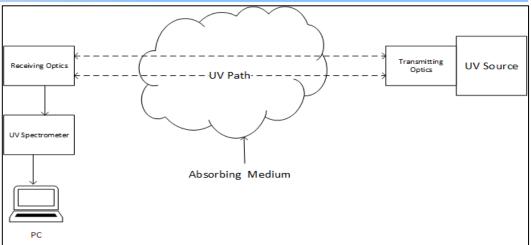






Low Signal Alarm Response (Level 1)

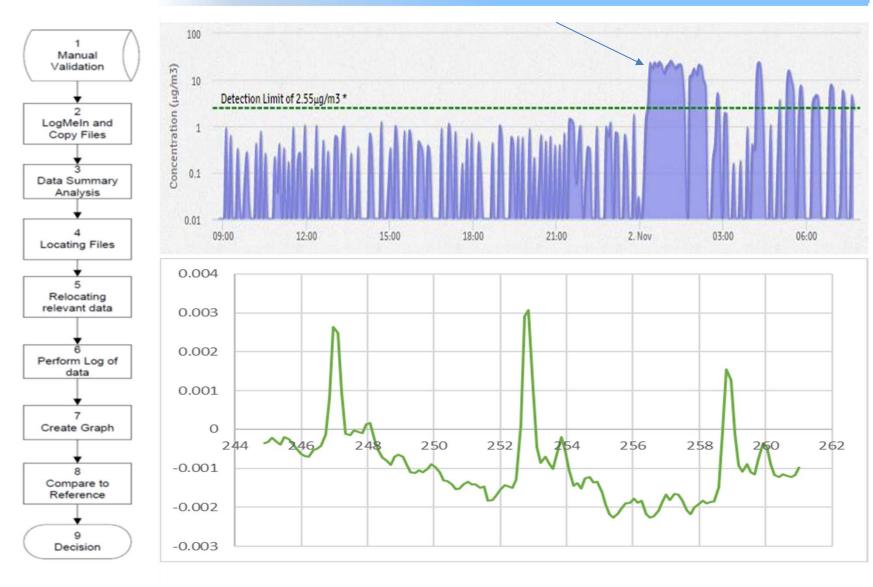








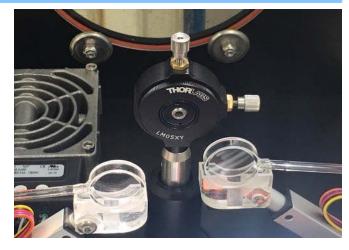
Manual Validation of Data (Level 1)

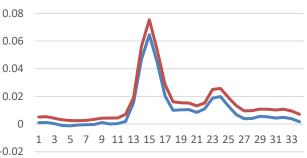




Traceability (Level 2)

- Create reference spectra using reference system.
- Fill sealed cells with gases for field spiking.
- Validate concentration of cell with reference system.
- Validate concentration of cell in the field.









Verification and Validation Model

Parameter		Method Acceptable	Site-specific Method Acceptable	Unacceptable
Relative Bias		<= 10%	Between 10% and 30%	> 30%
Precision		Relative Standard Deviation (RSD) <=20%		RSD > 20%
MDL		N/A	N/A	N/A
Accuracy		<= 15%		> 15%
Linearity		$R^2 >= 0.9$		$R^2 < 0.9$
Robustness Temperature		<= 1%		> 15%
Robustness Strength	Signal	<= 15%		> 15%
Robustness Time	Sample	<= 15%		> 15%



Results

Parameter	Value	Status
Relative Bias	2% over 5 systems	Method Acceptable
Precision	1.9% over 5 systems	Method Acceptable
MDL	0.475	< 0.9
Accuracy	5%	Method Acceptable
Linearity	0.92	Method Acceptable
Robustness Temperature	3% from 9 to 45 deg C	Method Acceptable
Robustness Signal Strength	0.74%	Method Acceptable
Robustness Sample Time	10% from 0.5 min to 30 min	Method Acceptable



Tasks: Level 3

Check Type	Check	Frequency	Reference Doc	Roles and Responsibilities
Level 3				
Instrumentation	Annual service FTIR, UV and OGD	Annual	FLM-QLT-SOP-005 for Planned Maintenance, Critical Spares Tracking List	Technical signatory
Instrumentation	Certification system brought to factory spec	Annual	FLM-QLT-SOP-014 Monthly Reporting	Technical signatory
Data	Full reconciliation of data	Monthly	FLM-QLT-SOP-014 Monthly Reporting	Technical signatory
Data	Supervisor check for data trends	Monthly	FLM-QLT-SOP-014 Monthly Reporting	Technical signatory
Program	Complete system audit	Annual	Internal Audit Plan	Technical signatory
Program	Program evaluation and upgrade	Annual	Annual Management Review	Quality Manager





Project Achievements

- Continuous Improvement
 - Automate Manual Validation of Data
 - Data used to track leaks
- More frequent MDL
- ISO 17025 accreditation for 4 sites







Areas of Improvement

- Proficiency testing
- More frequent MDL
- Lower detection limits
- Expanding accreditation to other sites



