

SOP Hot Topics

Panel Discussion

Environmental Measurement Symposium 2016

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Panelists

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- Scott Siders, PDC Laboratories
- Bob Pullano, GEL
- John Gumpper, ChemVal Consulting
- Pam Schemmer, TestAmerica





- Electronic vs Hardcopy
 - How are the documents stored and accessed at your lab?
 - If electronic, where can the staff access them? Do they print copies for use at the bench?





- Review and Revisions
 - What triggers a routine review?
 - Who is responsible?
 - If revisions are needed, what is the flow?





- Tracking the need for Edits
 - If an audit indicates the need for a change to the SOP, how is this started?
 - If someone identifies a minor edit that is not needed immediately, how is this tracked?
 - Do you have a formal feedback mechanism for lab staff?



- Ensuring accuracy
 - Versus the referenced method?
 - Versus actual lab practice?
 - How do you handle when a new version of a referenced method is published?





- Planned Deviations from the SOP
 - Do you have a process for handling a planned deviation that may be needed for a particular project, sample, etc.?





- Handling Proprietary Content
 - Do you provide copies of your SOPs to clients?
 - Are there any that are considered proprietary?
 - What if they are needed for documents that will become public (state audits, EPA QAPPs, etc.)?



Common Audit Findings

- What are examples of issues that were cited during audits (internal and external) regarding SOPs?
- What steps did you take to address these?





- What are the hot topics from the audience?





Thank you

Thank you all for participating.

Any other questions?

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