

Post Event Follow-Up Report

Instructions

This form can be used when follow-up is required on a Proficiency Testing Evaluation Report. This can include failures, not scored, non-consensus, self-assessment, use of an exception code, SDI bias or trends, etc.

Note: This document and its process are not intended to replace any existing protocol used by your facility or to replace any quidance or instructions given by an accrediting agency.

Documentation Retention Requirements: PT records and related documents (follow-up reports, calibration, daily QC, maintenance, etc.) must be retained for two years unless longer retention is required for specific analytes (such as Blood Bank for 5 years).

- Facility Name: If your facility is part of a large system, include a unique identifier such as city, WSLH PT Lab ID, or CLIA
 ID.
- Year, Event Name, and Event Number: This information is located on the top right of the Proficiency Testing Evaluation Report. On the fillable form, choose year, event name and event number from the drop-down menus.
- Module/Analyte/Sample: List your module number, analyte, and sample ids that require follow-up.
- Instrument/Method: Include a unique identifier for the instrument if necessary, such as a serial number. Kit name and lot number can be documented here.
- Reason for Follow-Up: This is displayed under "Status" or "Comments" on your Proficiency Testing Evaluation Report (such as Fail, Non-consensus, Not scored, etc.) On the fillable form, choose the reason for follow-up from the drop-down menu. The "Additional Comments" box can be used to document the reason for the use of an exception code or choice of "Other" and also provides additional space to list analytes/sample IDs that need follow-up.
- **Identified Cause:** The cause of the deficiency must be determined in order to implement corrective actions. For assistance in identifying potential causes and troubleshooting actions, see the **WSLH Troubleshooting Guide**.
- Troubleshooting or Self-Assessment Actions Performed: Document the steps taken to identify the root cause of the issue. Include with this report any documents that pertain to this such as retesting results, recalibration documentation, maintenance logs, service call logs, peer data and data calculator spreadsheets, split patient sample comparison data, etc.
- **Corrective Action:** Identify actions your facility can implement to prevent similar issues in the future. Include with this report any documents that pertain to this such as retraining documents, forms, updated SOPs, meeting minutes, etc.
- Person Completing Report: This lets any accrediting agencies know who to contact for any follow-up questions or concerns.
- Lab Director/Technical Supervisor/Consultant/Designee: All post event actions must be approved, signed and dated by the appropriate personnel.



Post Event Follow-Up Report

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Facility Name								
PT Event								
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Reason for	Follow-Up							
As stated in the evaluation report								
Additional	Comments							
Identified (Cause							
Troubleshooting or Self-Assessment Actions Performed								
Include any necessary documentation including retesting results, recalibration, service call log, etc.								
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Troubleshooting or Self-Assessment Actions Perfo	ormed continued from page 1					
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Corrective Action						
Include any necessary documentation such as forms, retraining documents, etc.						
Person Completing Report						
Printed Name	Signature	Date				
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Lab Director/Technical Supervisor/Consultant/De	rianco					
Lab Director/ recrimical Supervisor/ Consultant/ De	signee					
Printed Name	Signature	Date				