

# FORENSIC BIOLOGY QUALITY ASSURANCE / QUALITY CONTROL MANUAL

Approving Authority: Meredith Rosenberg, Quality Assurance Manager

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## Table of Contents

<b>Audits and Assessments</b>	<b>3</b>
I. External Audits/Assessments	3
II. Internal Audits	5
<b>Control of Data</b>	<b>9</b>
<b>Control of Non-Conforming Work</b>	<b>11</b>
Identifying Non-Conforming Work	11
Non-Conformity Reporting Form	12
Response to Non-Conformities (Corrective Actions)	15
<b>Control of Reference Collections</b>	<b>19</b>
<b>Court Testimony Monitoring</b>	<b>22</b>
<b>Equipment Calibration and Maintenance</b>	<b>25</b>
A. Critical Equipment	25
B. Non-Critical Equipment	26
C. General Preventative Maintenance	26
D. Equipment Decontamination	26
E. Instrument Irregularities	27

<b>Exogenous DNA Prevention .....</b>	<b>28</b>
A. Facility.....	28
B. Laboratory Clean-up .....	28
C. Sample Processing.....	29
D. Personal Protective Equipment (PPE).....	29
E. Contamination Prevention Equipment (CPE) .....	29
F. Identification .....	29
G. Interpretation and reporting.....	30
<b>Lab Types Database .....</b>	<b>31</b>
A. Sample Collection .....	31
B. Sample Processing.....	32
C. Sample Disposition .....	32
D. Database Maintenance.....	33
E. Lab Types Reference Databases .....	33
F. Searching the Lab Types Database .....	34
<b>Preventive Action .....</b>	<b>35</b>
<b>Proficiency Testing Program.....</b>	<b>37</b>
A. DNA Open-External Proficiency Testing Program .....	37
B. Serology Open-External Proficiency Testing Program.....	40
C. Blind Re-analysis Proficiency Testing Program .....	40
<b>Quality Incident Review .....</b>	<b>42</b>
A. Quality Incident Reporting.....	43
B. Quality Incident Review.....	44
C. Close-out of Quality Incident Reviews .....	45
<b>Reagents.....</b>	<b>46</b>
A. Reagents Prepared In-House .....	46
B. Commercial Reagents .....	47
C. Reagent Quality Control Testing.....	48
D. Reagent Records.....	49
E. REAGENTS USED BY THE DEPARTMENT .....	50
<b>Validation .....</b>	<b>54</b>
A. Developmental Validation.....	55
B. Internal Validation.....	55
C. Review and Approval of Validation.....	56
D. Training.....	57
E. Storage of Validation Records .....	57

[Quality Control Reagent Sheets](#)  
[Quality Control Procedures](#)

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 3 OF 57

## Audits and Assessments

### GUIDING PRINCIPLES AND SCOPE

Audits and assessments are conducted to improve the quality of the laboratory, as well as to maintain compliance with accreditation standards such as ISO 17025, the ASCLD/LAB-*International* Supplemental Requirements, and the FBI Quality Assurance Standards for Forensic DNA Testing.

An **Internal Audit** is an audit conducted by qualified and trained auditors employed by the Department of Forensic Biology. An **External Audit or Assessment** is an audit conducted by qualified and trained auditors/assessors employed by agency external to the Department of Forensic Biology.

This document describes the external audits/assessments to which the Department of Forensic Biology is subject and the internal audit program of the Department.

### PROCEDURE

The management system of the Department of Forensic Biology is designed to conform to the following sets of standards:

**ASCLD/LAB-*International* Standards:** The ASCLD/LAB-*International* Standards encompasses the ISO/IEC 17025 requirements and the ASCLD/LAB-*International* Supplemental Requirements.

**FBI Quality Assurance Standards for Forensic DNA Testing:** The FBI Quality Assurance Standards for Forensic DNA Testing is issued by the FBI Director and is a set of standards specific to Forensic DNA Testing (mitochondrial and autosomal). ASCLD/LAB-*International* also requires compliance with these standards as a condition of accreditation.

### I. External Audits/Assessments

4. The Department is subject to **external accreditation assessments/surveillance visits** as required by ASCLD/LAB.
  1. Assessment scheduling and the assessment process are the responsibility of ASCLD/LAB.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 4 OF 57

2. Level 1 non-conformities (as defined by ASCLD/LAB) must be corrected to the satisfaction of ASCLD/LAB before a recommendation for accreditation is made.
  3. Corrections for Level 2 non-conformities may commence immediately upon discovery. Otherwise, the Department shall obtain approval from ASCLD/LAB to correct the non-conformity prior to the next, annual on-site Surveillance Visit.
- B. An **external DNA audit** to ensure the Department's conformance with the FBI DNA Quality Assurance Standards for Forensic DNA Testing is conducted at least once every two (2) calendar years.
1. An external DNA audit could occur as part of an ASCLD/LAB accreditation assessment or could be a stand-alone DNA audit such as those provided through the DNA laboratory audit program of the National Forensic Science Technology Center (NFSTC).
  2. For an external DNA audit to "count," it must occur at least 6 months, but no more than 18 months, after an internal or external DNA audit conducted during the prior calendar year.
  3. The audit must be conducted with the version of the FBI DNA QAS audit document in effect at the time of the audit.
  4. The audit document and any Quality Incident Reviews stemming from non-conformities identified during the audit are submitted to the Quality Assurance Manager and the DNA Technical Leader(s) for review and for approval of proposed follow-up actions.
  5. A copy of the DNA audit documentation and laboratory responses to non-conformities is provided to the NDIS Custodian at the FBI within 30 days of the laboratory's receipt of the audit report.
  6. The laboratory maintains the following records from external DNA audits:
    - a. Audit reports
    - b. Self-verification forms completed by the members of the audit team to certify their qualifications as auditors and experience with the DNA technologies and platform(s) used by the Department.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 5 OF 57

- C. External audits outside of normal accreditation assessment or external DNA audit schedules may be required by ASCLD/LAB or the New York State Commission on Forensic Science as a response to very serious quality incidents.
- D. The Quality Assurance Manager (QAM) is the point of contact for any external audit of the laboratory that concerns the technical operations of the laboratory.

## II. Internal Audits

The internal audit program is a critical component of the Department's management system. It is designed to ensure that the Department's management system is functioning correctly and that the Department is operating in compliance with its own procedures as well as regulatory and accreditation requirements.

The internal audit program consists of two parts: (1) audits to evaluate the laboratory's conformance with respect to the management system, including the testing activities, and with the ISO 17025 and ASCLD/LAB-*International* Supplemental requirements and (2) DNA audits to evaluate the laboratory's conformance with respect to the FBI's Quality Assurance Standards for Forensic DNA Testing Laboratories.

### A. General Internal Audit Information

1. The Quality Assurance Manager (QAM) is responsible for scheduling and planning the internal audits of the laboratory. Scheduling is done in consultation with the Technical Leaders, Deputy Director(s), and the Director.
  - a. Should an audit require personnel from external organizations, the QAM will take into consideration the schedule and availability of these external auditors prior to agreeing to a date.
2. "ISO" audits and DNA audits are scheduled to occur each calendar year.
3. The QAM selects auditors to ensure that an audit team is "qualified" as per the requirements for each type of audit.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 6 OF 57

4. Each audit team has a lead auditor/team leader. The QAM and the lead auditor/ team leader develop an audit plan that, at a minimum, contains the audit schedule, the activities to be audited, and the audit team(s) assigned to audit the specified activities.
5. The general process for any internal audit is as follows:
  - a. The QAM notifies the laboratory that an internal audit will be conducted, the general scope of the audit, and provides an approximate timeframe.
  - b. The QAM schedules an opening conference with the auditors to discuss the audit objectives, assignments, timing, and report format and distribution.
  - c. The auditors perform their audit activities to assess the soundness of the quality system, management system, and technical operations.
  - d. The audit teams provide the QAM with their audit findings, including potential non-conformities and observations.
  - e. The QAM discusses preliminary observations (if any) with management.
    - 1) Non-conformities that are non-systemic, are easily corrected, and do not indicate serious deficiencies in the management system can be corrected prior to the completion of the audit.
    - 2) The correction is documented in the audit records, but is not included in the final audit report.
  - f. The QAM, Technical Leaders, and other manager(s) as requested by the QAM review the audit results submitted by the audit teams and verify the findings that are true non-conformities supported by objective evidence.
  - g. The lead auditor/ team leader writes the audit report. The QAM reviews the audit report. The laboratory managers are informed.
  - h. The CONTROL OF NON-CONFORMING WORK procedure and/or the QUALITY INCIDENT REVIEW procedure is used for follow-up on audit non-conformities identified in the audit report.
  - i. If audit non-conformities show that laboratory results may have been affected, the laboratory must notify its customers and accreditation agency of the results, in writing, within thirty (30) days of discovery.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 7 OF 57

- j. Audit reports may need to be submitted to the NYS Commission on Forensic Science, ASCLD/LAB, or the board members of the National DNA Indexing System (NDIS). The Quality Assurance Manager shall ensure timely submission of audit reports when necessary.
- k. Audit reports are a form of records and shall be retained according to the guiding principles of the laboratory. See CONTROL OF RECORDS for further information.

## B. Information Specific to Internal “ISO” Audits

- 1. The scope of the internal audit must ensure that all elements of the management system are addressed. The QAM or designees may develop checklists to be used by the audit teams.
- 2. Auditors are “qualified” in any of the following ways:
  - a. Documented completion of an ASCLD/LAB-International assessor training course.
  - b. Documented completion of an external ISO 17025 training course and auditor training conducted in-house by a qualified auditor such as the QAM.
  - c. Documented completion of ISO 17025 and auditor training conducted in-house by a qualified auditor such as the QAM.
- 3. Only qualified auditors will be selected to lead an internal audit team. Staff that has not completed the required training may be used as team auditors, but they must report directly to a qualified auditor.

## C. Information Specific to Internal DNA Audits

- 1. DNA internal audits are conducted using “The FBI Quality Assurance Standards Audit for Forensic DNA Testing Laboratories.”
- 2. Auditors are “qualified” to conduct DNA audits if they have successfully completed an FBI-sponsored DNA Auditing Workshop/Course.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

AUDITS AND ASSESSMENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 8 OF 57

3. The DNA audit team must contain at least one qualified auditor and at least one person that is, or has previously been, a qualified analyst for each specific DNA technology (*technology is used to describe the type of forensic DNA analysis performed in the laboratory, such as STR, YSTR, or mitochondrial DNA*) performed in the laboratory. This may be accomplished by having a single auditor who meets all of the specified qualifications or through a combination of various members of a multi-person audit team.
4. Internal DNA audits are optional in calendar years when external DNA audits have been conducted.

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## Revision History:

February 9, 2010 – Initial version of procedure.  
May 1, 2015 – Updated section to reflect current practices.  
December 24, 2015 – Removed phrase “as needed” from Section II. A. 2.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF DATA		
DATE EFFECTIVE 07-16-2012	APPROVED BY Quality Assurance Manager	PAGE 9 OF 57

## Control of Data

### GUIDING PRINCIPLES AND SCOPE

When computers or automated equipment are used for the acquisition, processing, recording, reporting, storage or retrieval of test data, the laboratory shall ensure that:

1. Calculations and data transfers are subject to appropriate checks in a systematic manner.
2. Computer software developed by the laboratory is documented in sufficient detail and is suitably validated as being adequate for use.
3. Procedures are established and implemented for protecting the data; such procedures shall include, but not be limited to, integrity and confidentiality of data entry or collection, data storage, data transmission and data processing.
4. Computer and automated equipment are maintained to ensure proper functioning and are provided with the environmental and operating conditions necessary to maintain the integrity of test data.

This section describes the procedures to achieve these guiding principles.

### PROCEDURE

Only Department of Forensic Biology staff members have unlimited access to the Forensic Biology network drive. Exceptions may only be granted by the Director or designee. Access is controlled by the OCMF Management Information Systems (MIS) Department. Unless otherwise authorized by an existing standard operating procedure, only the Quality Assurance Manager may authorize the release of data to any party (via any means) external to the Department of Forensic Biology.

Computer software may be used during the processing of case work; however, the results will be incorporated into the case record.

Any calculations and data transfers made using computer software are reviewed for its accuracy by a supervisor prior to its incorporation into a case record and/or are reviewed for its accuracy during the final technical review process of the case.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF DATA		
DATE EFFECTIVE 07-16-2012	APPROVED BY Quality Assurance Manager	PAGE 10 OF 57

Computer software or software modifications developed by the laboratory are suitably validated depending on the purpose of the modification.

1. The appropriate Technical Leader must be consulted prior to validation to ensure that suitable validation tests are carried out.
2. If the software is used to streamline/transfer data, sufficient proof must be furnished to document that the intended purpose of the software is achieved. This may be accomplished by entering a simple set of data to ensure that the streamline/transfer of data is accurate.
3. If the software is used to calculate data, sufficient proof must be furnished to document that the intended purpose of the software is achieved. This may be accomplished by inputting a simple set of data and comparing it to hand-calculated results to ensure that the calculations made are correct.
4. Computer software developed by the laboratory must be approved by the appropriate Technical Leader prior to its use in casework.
5. Validation records are stored by the Quality Assurance Unit.

Once calculations and data transfers have been reviewed by a supervisor, they may be deleted from the Forensic Biology network drive. For some data, such as DNA electropherograms, the electronic data will be maintained indefinitely.

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## Revision History:

February 9, 2010 – Initial version of procedure.

July 16, 2012 – Specific terminology was removed and replaced with generic terminology to accommodate LIMS.

Controlled versions of Department of Forensic Biology Manuals only exist electronically on the Forensic Biology network. All printed versions are non-controlled copies.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 11 OF 57

## Control of Non-Conforming Work

### GUIDING PRINCIPLES AND SCOPE

Non-conforming work is any testing work which does not meet the Department of Forensic Biology's stated standards, either with respect to mode of execution or outcome. All non-conforming work must be addressed upon discovery so that the work can be appropriately evaluated, corrected as needed and prevented in the future.

This procedure describes the Department's process for evaluating non-conforming work, performing root cause analyses and taking appropriate follow-up action. Technical problems or difficulties can arise in all phases of Department operations. Listing each potential problem is impractical, therefore this topic is considered in general terms.

**Apparently similar situations may result in different follow-up actions. This is because no two circumstances are exactly the same and the consequences of the particular non-conformity may be very different.**

### Identifying Non-Conforming Work

1. Any member of staff who discovers a technical, analytical or clerical error or realizes that there is a *technical, analytical or clerical problem that may compromise evidence integrity, the accuracy of casework analysis or results reported*, must address the issue immediately. The analyst who discovers the non-conformity or the supervisor to whom the incident is reported becomes the **principal investigator of the non-conformity**. Situations may arise where an analyst them self may discover their own error. When this occurs, the analyst may become the principal investigator and write up the non-conformity for review by their direct supervisor AND the Quality Assurance Manager. The principal investigator is tasked with investigating and evaluating the significance of the non-conforming work.

a. **Technical problems related to the testing of a batch of samples** are reported to a supervisor. As multiple analysts and/or cases may be affected, corrective actions may be documented on the batch worksheets, via LIMS deviations or via emails to analysts' whose cases are affected.

- Detection of exogenous DNA in negative controls is reported to a Quality Assurance Supervisor. **Note:** Determination of the source of exogenous

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 12 OF 57

DNA where less than 8 alleles are seen in at least 4 loci is difficult and may not be feasible.

- b. **Technical problems related to individual case samples** (e.g., possible sample mix-up) are reported to a supervisor, the analyst(s) assigned to the affected case(s), the supervisor(s) of the affected case(s) AND the Quality Assurance Manager.
- c. **Analytical or Clerical errors affecting reported results** (e.g., errors in conclusions drawn from analytical results or incorrect recording or transcribing of observational or analytical results) are to be reported to the case analysts' direct supervisor and/or Assistant Director as well as the Quality Assurance Manager.
- d. **Technical problems identified during routine quality control activities**, such as instrument performance checks, are reported to a Quality Assurance supervisor and are reported on the performance check worksheets.

## Non-Conformity Reporting Form

- Some non-conforming work can be easily corrected, such as by reanalysis. An example of this would be a sample that fails to give interpretable peaks in an electrophoresis run, but when re-injected an acceptable result is obtained. In such cases the action is documented on the batch worksheets, in case notes, or on performance check worksheets, as appropriate to the situation, but no further investigation is likely to be needed unless the incident was part of a pattern. In such cases, a **Non-Conformity Reporting Form does not need to be filled out.**
- Some non-conforming work, such as contamination incidents or analyst errors in reporting case results require more investigation as to the scope and cause of the non-conformity. The incident and its evaluation are **documented on the Non-Conformity Reporting Form.**
  - Should the cause of the non-conformity be attributed to an individual, the **Non-Conformity Reporting Form** must be completed by both the principal investigator and the immediate supervisor of the individual (if not already involved as the principal investigator).

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 13 OF 57

When a non-conformity has been discovered, the investigation shall proceed in one of two ways. **It MUST first be determined by the principle investigator in which manner the non-conformity investigation shall proceed.** The two routes a non-conformity investigation may take are as follows:

1. It must be determined whether or not a “**Significant Event**” has occurred. Such “Significant Events” (as indicated in Section 17-207 of Chapter 2 of Title 1 of the Administrative Code of the City of New York) include:
  - Intentional fabrication of work product, evidence examination, analysis or test results.
  - Significant error(s) by an employee, or deficiency in a system or procedure that may have affected the accuracy of reported results of evidence examination or the accuracy of the reported results of analysis in one or more cases.
  - Failure of an employee to follow protocol such that it may have affected the accuracy of reported results of evidence examination or the accuracy of the reported results of analysis in one or more cases.
  - Statements made in the course of testimony by which an employee significantly misrepresents or misstates his/her education, experience, training or qualifications, or the reported results of any evidence examination or analysis.

**IF it has been determined by the principle investigator that a “Significant Event” has occurred, the non-conformity investigation must then proceed as follows:**

1. The principle investigator of the non-conformity **MUST IMMEDIATELY** inform the Quality Assurance Manager that a non-conformity on a “Significant Event” level has occurred. The Quality Assurance manager will assess the non-conformity to determine if indeed a “Significant Event” has occurred.
  1. If a “Significant Event” is found to have occurred, the Quality Assurance Manager **MUST IMMEDIATELY** inform the Office of Chief Medical Examiners’ Root-Cause Analysis Officer of the “Significant Event” and the principal investigator shall begin to document the non-conformity on the Non-Conformity Reporting Form.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 14 OF 57

- b. If the Quality Assurance Manager has determined that a “Significant Event” has not occurred, the principal investigator may carry out a standard non-conformity investigation.
2. The Root Cause Analysis Officer shall follow the procedures mandated by New York City Legislation and the Root Cause Analysis Procedure of the Office of Chief Medical Examiner. This involves convening a Root Cause Analysis Committee to investigate the “Significant Event”.
3. The Department of Forensic Biology shall ensure that the Office of Chief Medical Examiner issues the report of the Root Cause Analysis Committee to the following entities **within 7 days of the report being generated:**
  - The NYC Mayor’s Office/ NYC Council
  - The NYS Commission on Forensic Science
  - ASCLD/LAB
  - Relevant District Attorney’s Office
  - Relevant Defense Council of record

**If it has been determined by the principle investigator that a “Significant Event” has NOT occurred, the non-conformity investigation shall proceed in the following manner:**

1. A Non-Conformity Reporting Form shall be completed. The principle investigator must determine several factors and document the results of their investigation in the Non-Conformity Reporting Form. Factors to be investigated include, but are not limited to, the following:
  - a. The nature of the detected non-conformity.
  - b. How the non-conformity was detected.
  - c. The cause of the non-conformity. **This will require an in-depth Root- Cause Analysis of the situation.**

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 15 OF 57

- d. How the non-conformity was resolved (immediate corrective actions taken).
- e. Recommendations on future preventive actions to avoid the non-conformity from recurring.
- f. Parties notified of the non-conformity.

Once the non-conformity investigation has been completed and the Non-Conformity Reporting Form has been filled out, the form must be given to the Quality Assurance Manager for review. The Quality Assurance Manager maintains the right to request further investigation into the non-conformity on an as-needed basis. Further investigation may be performed by the principle investigator, the Quality Assurance Manager, or another designated employee.

If a particular analyst is found to be the cause of the non-conformity, they must sign the Non-Conformity Reporting Form, as well as their direct supervisor. The direct supervisor shall track performance issues to ensure that repeated occurrences of similar issues are corrected through counseling, retraining, or other measures appropriate to the situation.

Any corrective actions taken to rectify the non-conformity are documented on the Non-Conformity Reporting Form and on the patch worksheets, in case notes or on performance check worksheets, as appropriate to the situation.

If the initial corrective action taken fails to correct the problem, the issue should then be referred to the Quality Assurance Manager for further investigation.

## **Response to Non-Conformities (Corrective Actions)**

1. Based upon the initial investigation into the non-conformity by the principle investigator, the following may occur:
  - The DNA Technical Leader(s) have the authority to suspend DNA analytical operations for the Department or an individual until such time as the technical issue has been resolved through corrective action.
  - The Serology Technical Leader has the authority to suspend serology analytical operations for the Department or an individual until such time as the quality issue is resolved through corrective action.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 16 OF 57

●The Director, Deputy Director(s) and Assistant Directors are notified as soon as practicable when actions to suspend testing are proposed or taken.

2. If it has been determined by the principle investigator or, subsequently by the Root Cause Analysis Officer, that a “Significant Event” has not occurred, appropriate corrective actions may be implemented by the laboratory in order to ensure that the non-conformity does not recur.
3. Once the Root Cause Analysis Committee has issued their report concerning a “Significant Event”, any recommendations they make concerning corrective actions may be implemented by the laboratory in order to ensure the non-conformity does not recur.
4. The Quality Assurance Team analyzes Non-Conformity Reporting Forms on a regular basis in order to track issues so that trends can be identified.
  - a. Non-Conformity Reporting Forms are assessed at least quarterly to determine if similar events occurred (such as those in the same area of testing or caused by the same individual) within an unreasonable timeframe.
  - b. The Quality Assurance Manager will determine if any trends pose additional concerns to the Management System of the laboratory.

Document Control Coordinator 06/20/2016



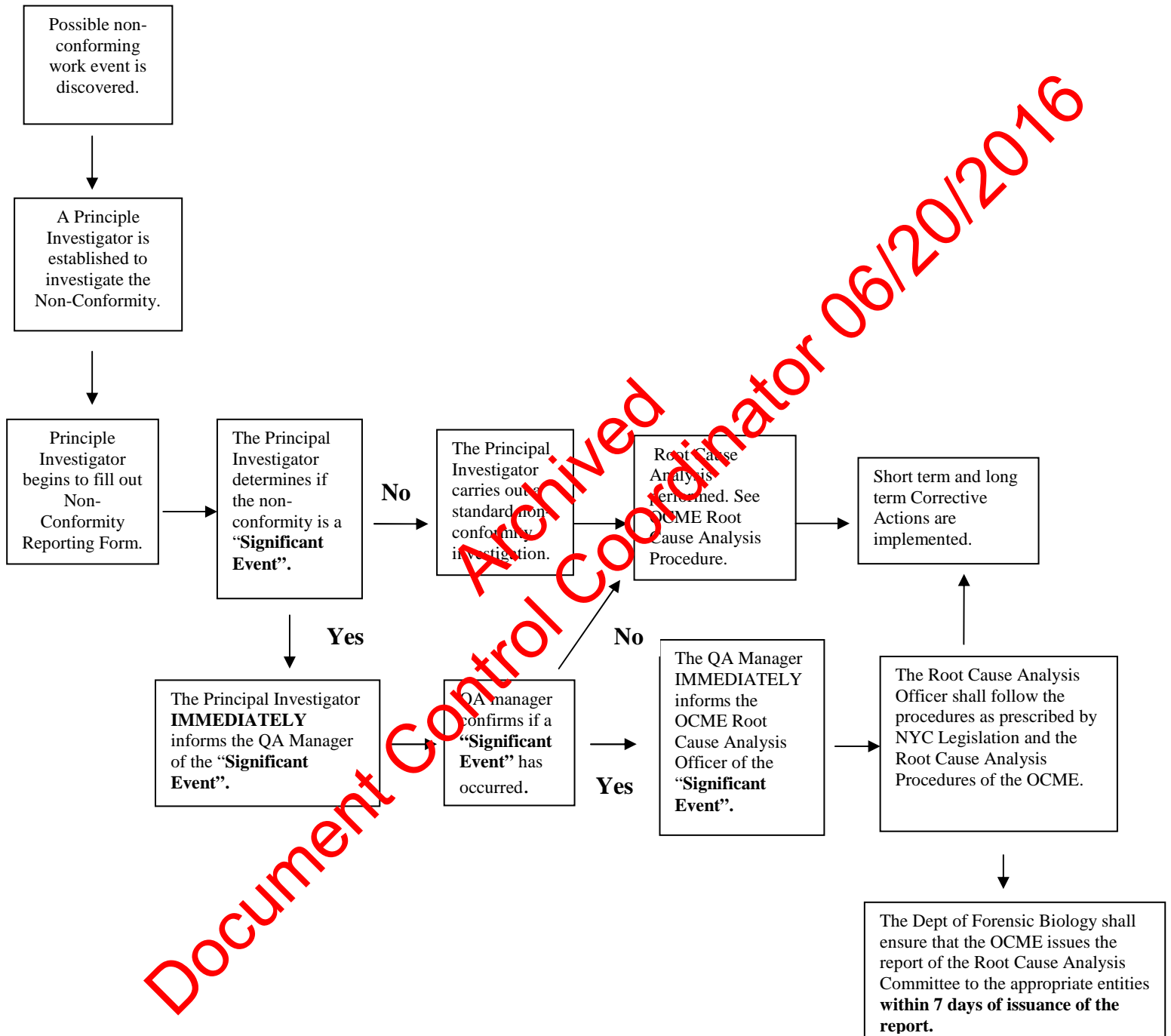
# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

## CONTROL OF NON-CONFORMING WORK

DATE EFFECTIVE  
04-15-2016

APPROVING AUTHORITY  
QUALITY ASSURANCE MANAGER

PAGE  
17 OF 57



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF NON-CONFORMING WORK		
DATE EFFECTIVE 04-15-2016	APPROVING AUTHORITY QUALITY ASSURANCE MANAGER	PAGE 18 OF 57

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## Revision History

- April 1, 2014- Initial version of updated procedure. This new version is necessary to account for new NYC Legislation concerning Root Cause Analysis. Previous version archived.
- September 1, 2014 – Clarification made that the analyst who discovers a non-conformity may also become the Principle Investigator.
- April 15, 2016- Added city code which dictates what constitutes a “Significant Event”.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF REFERENCE COLLECTIONS		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 19 OF 57

## Control of Reference Collections

### GUIDING PRINCIPLES AND SCOPE

Reference standards and reference materials shall be stored in a manner that ensures the prevention of contamination or deterioration in order to protect their integrity. Procedures for safe handling, transport, and use of reference standards are outlined below.

### PROCEDURE

#### A. Reference Standards

Reference standards of measurement are to be used for calibrations only and for no other purpose. Since the laboratory does not conduct any calibrations, reference standards do not exist within the laboratory.

#### B. Reference Materials

Reference materials used to conduct intermediate performance checks of instruments and equipment are, where possible, traceable to certified reference materials such as those from the National Institute of Standards and Technology (NIST). Reference materials that cannot be traceable to certified reference materials must be certified according to original manufacturer's specification. Internal reference materials are checked as far as is technically and economically possible.

Where possible, reference materials are obtained from sources that can supply appropriate traceability information such as a Certificate of Analysis.

Reference materials are stored according to manufacturer's specifications. If the manufacturer's specification does not indicate storage conditions, the laboratory determines how similar materials are stored within the laboratory and applies those storage conditions to the reference materials.

Storage conditions must ensure the safe handling and safe transport of reference materials. Furthermore, storage conditions must minimize contamination or deterioration, where possible.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF REFERENCE COLLECTIONS		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 20 OF 57

Checks needed to maintain confidence in the calibration status of the reference materials shall be carried out periodically. Where practical, reference materials must be re-certified on or before the expiration date or they must be removed from use. Any reference material that does not have an expiration date must be re-certified or removed from use after one (1) year of its first use.

## 1. Standard Reference Materials (SRMs)

The use of standard reference materials is essential to reliable methodology. The laboratory will check its DNA typing procedures against an appropriate and available SRM annually or whenever substantial changes are made to the typing procedure. SRMs will be purchased from the National Institute of Standards and Technology (NIST) and shall have an associated Certificate of Analysis available. The laboratory may choose to use other SRMs to check any of its procedures, but it is not required to do so. SRMs must not be used beyond its expiration date unless a Certificate of Analysis is issued from NIST to document its recertification.

Secondary standards that are traceable to SRMs may be created by the laboratory for use in lieu of purchasing them directly from NIST. To create a secondary standard, a "lot" of DNA samples (such as a blood stain) must be run and analyzed in parallel with an appropriate NIST SRM. Documentation must be maintained to demonstrate that the results for the SRM are correct (as compared to the certificate of analysis) and the results of the secondary standard are consistent (as compared to a prior result).

## 2. Weights

The laboratory will conduct intermediate performance checks of balances using Class 1 weights that are traceable to NIST Standards. Weights must be calibrated prior to the expiration date of the Certificate of Analysis, or must be removed from use. Various companies exist that can calibrate weights traceable to NIST Standards. However, the Department of Forensic Biology will endeavor to select a company that is accredited in accordance with ISO 17025 standards for calibration laboratories.

Prior to each use, analysts should visually inspect the weights to ensure that there are no physical defects that would affect their performance.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

CONTROL OF REFERENCE COLLECTIONS		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 21 OF 57

## C. Reference Collections

The laboratory uses a DNA “Lab Types” reference collection for comparison to case work DNA profiles to ensure that no exogenous DNA is present in samples and a “suspect database” to determine if there is an association between a named suspect and DNA profiles from previously tested cases. The use of these reference collections are outlined in the Forensic Biology CODIS Manual and the LAB-TYPES DATABASE procedure. These manuals identify these reference collections and describe how they are controlled.

Archived  
Document Control Coordinator 06/20/2016

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### Revision History:

February 9, 2010 – Initial version of procedure.

Controlled versions of Department of Forensic Biology Manuals only exist electronically on the Forensic Biology network. All printed versions are non-controlled copies.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

COURT TESTIMONY MONITORING		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 22 OF 57

## Court Testimony Monitoring

### GUIDING PRINCIPLES AND SCOPE

Court testimony is the culmination of the work performed by the laboratory's scientists. To ensure that court testimonies are relevant, and presented in a clear and professional manner, the testimony of each testifying examiner is monitored at least once during a calendar year providing testimony is rendered.

This document describes the Department of Forensic Biology's courtroom testimony monitoring program.

### PROCEDURE

When a case goes to grand jury or trial, the Reporting Analyst (RA) will be contacted to testify either by phone or subpoena. An informal request by phone should be directed to the RA's supervisor to gather details of the testimony. OCME counsel should be consulted if the request is via a subpoena. In either case, a pre-trial with the Assistant District Attorney (ADA) or defense attorney is advisable to discuss or go over the line of questioning. The RA should pull the case and all cross-referenced cases and/or suspect files. The RA should also bring a copy of his/her curriculum vitae and a spell-sheet to court.

If this is the RA's first testimony for the year or if the RA is inexperienced, their supervisor should be present at the pre-trial and trial. In addition to answering questions and providing support, the supervisor is responsible for evaluating the RA's testimony at trial. Evaluation of the RA's testimony at grand jury is left to the ADA, since no observers are allowed into court for grand jury.

#### A. Documenting Court Attendance

Staff members who are called to appear in court must have each court appearance documented, regardless of whether testimony was provided and/or evaluated.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

COURT TESTIMONY MONITORING		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 23 OF 57

## B. Testimony Monitoring

1. The testimony of each examiner is monitored at least once each calendar year, assuming that testimony is rendered. It is the responsibility of each testifying examiner to ensure that this is done.
2. Acceptable methods of courtroom monitoring are:
  - a. Direct courtroom observation by a higher-level supervisor (Criminalist Level IV or above).
    - i. This is the preferred method for trial testimony.
    - ii. In most cases the "higher-level supervisor" will be the immediate supervisor of the testifying examiner; however, a peer of the immediate supervisor or a higher level manager may perform the monitoring.
  - b. Direct courtroom observation by an ADA and/or defense attorney present during the testimony.
    - i. Evaluation by the ADA is the preferred method for Grand Jury testimony.
    - ii. For evaluation of trial testimony, the testifying examiner should attempt to get feedback from both the ADA and the defense attorney. The testifying employee can ask the attorney who summoned them to court to provide an evaluation form to the opposing counsel; however, if the attorney is not willing to do so it is not necessary to insist that it be done.
3. The testimony evaluator completes a Forensic Biology Court Testimony Evaluation Form. The form includes evaluations/comments on the following areas:
  - a. Appearance
  - b. Poise
  - c. Effectiveness of presentation (technical knowledge, ability to convey scientific concepts)
  - d. Interpretation of laboratory results
4. Evaluation forms completed by someone other than the testifying employee's immediate supervisor are forwarded to the testifying examiner's immediate supervisor.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

COURT TESTIMONY MONITORING		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 24 OF 57

5. Immediate supervisors review the evaluation with the testifying examiner, discussing areas of strengths and weaknesses.
  - a. The immediate supervisor may prescribe remedial action if the evaluation is unsatisfactory. Deficiencies in knowledge or courtroom presentation may require remedial training that includes one or both of the following:
    - i. Retraining on technical information if the testimony was inaccurate.
    - ii. Moot court retraining if the testimony showed deficiencies in the ability to express the concepts clearly.
6. The immediate supervisor and the testifying examiner sign/initial and date the evaluation form.
7. Completed evaluation forms are forwarded to the Quality Assurance Unit for entry into LIMS and hardcopy storage.

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## Revision History:

- February 9, 2010 – Initial version of procedure.
- January 6, 2011 – Modify section B.2 to allow for courtroom testimony evaluation at trial by defense attorneys and to describe a possible mechanism for supplying forms to both ADAs and defense counsel.
- July 16, 2012 – Specific forms and worksheets were removed and replaced with generic terminology to accommodate LIMS.
- August 14, 2015 – updated section to reflect LIMS usage.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EQUIPMENT CALIBRATION AND MAINTENANCE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 25 OF 57

## Equipment Calibration and Maintenance

### GUIDING PRINCIPLES AND SCOPE

Equipment maintenance, calibration, and performance checks are essential for establishing confidence in the results that are generated during routine testing of forensic DNA samples. The Department of Forensic Biology uses equipment, and associated software, which is suitable for the tests conducted and will not use equipment or software that is outside of its permanent control.

Equipment and its associated software are operated solely by trained and authorized laboratory personnel. Vendor equipment manuals are maintained and are available to all laboratory personnel on the Forensic Biology network.

All laboratory equipment is given a unique identifier by the laboratory. This unique identifier, as well as the manufacturers unique identification, and the instrument's current location, is maintained within the LIMS. Maintenance, calibration, and performance check information specific to each instrument is maintained within the LIMS. Equipment maintenance schedules are maintained by the Quality Assurance Team.

### PROCEDURE

#### A. Critical Equipment

**“Critical equipment” is that which requires calibration or a performance check prior to its use in casework and periodically thereafter.** Such equipment must have records of calibration and/or preventative maintenance. Specific calibration, performance check, and/or preventative maintenance programs and procedures for critical equipment are found in the Quality Assurance/Quality Control Procedures Manual.

Critical equipment must have maintenance usage logs.

The following is “critical equipment” used within the Department of Forensic Biology for DNA testing:

- Balances/scales

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EQUIPMENT CALIBRATION AND MAINTENANCE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 26 OF 57

- Thermal cyclers
- Real-time PCR systems
- Genetic analyzers
- Robotic systems
- Mechanical pipetteters
- Thermal cycler temperature-verification systems.

The FBI Quality Assurance Standards for Forensic DNA Testing (July 2009) lists traceable thermometers used for conducting performance checks and electrophoresis detection systems as “critical equipment,” however, the Department of Forensic Biology does not utilize these items.

## B. Non-Critical Equipment

All other equipment that is not covered under the definition of a “critical equipment,” as per the FBI Quality Assurance Standards for Forensic DNA Testing (July 2009) is considered “non-critical.” Examples of such equipment include pH meters, vortexers, and thermomixers.

The Department shall strive to conduct preventative maintenance on all non-critical equipment whenever feasible; however, it is not required to do so.

## C. General Preventative Maintenance

Maintaining cleanliness of any scientific equipment is the key to preventive maintenance. Spills must be taken care of **IMMEDIATELY**. Some spills may be corrosive to neighboring equipment and cause more damage than necessary. While some spills can be cleaned at the desk, some will require special treatment and/or additional follow-up. It is always best to contact the Forensic Biology Safety Coordinator or the OCME Health and Safety Unit for further information where needed.

## D. Equipment Decontamination

Various Quality Control Procedures have been developed to help maintain a DNA-free environment at the points of sample contact with the equipment used in DNA analysis. A 10% bleach solution is extremely effective in degrading DNA and is thus used for general cleanup procedures of equipment and the laboratory environment (e.g. laboratory desks

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EQUIPMENT CALIBRATION AND MAINTENANCE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 27 OF 57

and benches). Because of its corrosive nature, the use of 10% bleach should be followed by the use of 70% ethanol and/or deionized water. Decontamination of equipment should be done before movement of equipment from one lab space to another or removal from the laboratory permanently.

## E. Instrument Irregularities

Anyone observing any irregularities with any equipment may suspend the equipment from casework use to prevent the potential loss of sample. If this occurs, the Quality Assurance Unit and/or the appropriate Technical Leader must be notified shortly thereafter for follow-up. If the irregularity cannot be repaired and must be taken offline, the Quality Assurance Unit member or the appropriate Technical Leader must properly mark the equipment to prevent further use.

Prior casework performed on an instrument exhibiting irregularities may be evaluated to determine if the equipment irregularities affected casework results. This will occur on an as-needed basis. If casework results have been affected, the CONTROL OF NON-CONFORMING WORK procedure shall be followed.

Should repair and/or re-calibration occur, only designated Quality Assurance Unit members or the appropriate Technical Leader may re-certify that the equipment is available for casework. Instruments taken offline are taken out of service in LIMS so they cannot be selected for casework use. Any equipment taken offline for an extended period of time must either be removed from the bench, or a sign must be placed on the equipment to ensure that it is not used until appropriate repairs are made.

Re-certification requires that a designated Quality Assurance Unit member and/or the appropriate Technical Leader ensure that any required performance checks have been successfully completed, documentation that the instrument is available for casework has been entered in the appropriate maintenance log (hard copy or LIMS), and any signage to indicate otherwise is removed.

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### Revision History:

- February 9, 2010 – Initial version of procedure.
- July 16, 2012 – Specific terminology was removed and replaced with generic terminology to accommodate LIMS.
- May 1, 2015 – Updated Guiding Principles and Scope section, Equipment Decontamination section and Instrument Irregularities section.
- August 14, 2015 – Updated section to reflect LIMS practices and changed “QA Unit supervisor” to “designated QA member”.

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# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EXOGENOUS DNA PREVENTION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 28 OF 57

## Exogenous DNA Prevention

### GUIDING PRINCIPLES

Exogenous DNA is defined as the addition of DNA/biological fluid to evidence or controls subsequent to the crime. Sources of exogenous DNA could be first responders, crime scene technicians, NYPD personnel, or laboratory personnel, to name a few.

It is the goal of the Department of Forensic Biology to not transfer any DNA from employees to any casework sample. Several measures have been taken to prevent this, and this document will cover these measures in general.

### PROCEDURE

#### A. Facility

The laboratory is divided into physically isolated areas for evidence examination, DNA extraction, pre-amplification (amplification setup) and post-amplification (amplification and DNA typing). Each area has its own dedicated equipment. Once samples are accepted into the laboratory, they move through these areas in one direction only. Samples are first processed in the evidence examination area. They are then moved to the DNA extraction area. Following DNA extraction, aliquots of each sample are quantitated in the DNA quantitation area. Following DNA quantitation, aliquots of each sample are moved into the pre-amplification area. Here fresh kit reagents are stored and samples are prepared for amplification. Finally, the samples are amplified and typed in the post-amplification area. This laboratory setup helps eliminate the travel of DNA from post-amplification areas back into non-amplified DNA areas.

#### B. Laboratory Clean-up

In addition to the separation of space between analyses, the Department has implemented a documented clean-up program on a monthly basis. The documented clean-up program may be more frequent in areas where High Sensitivity DNA Testing is performed. The clean-up program involves the decontamination of instruments/equipment, bench/counter tops, sinks, etc. While 10% Bleach is extremely effective in destroying exogenous DNA, it is also very corrosive. Care should be taken so that when 10% Bleach is used, it is immediately followed by 70% Ethanol and/or water to wash off the Bleach from the surface of instruments/equipment.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EXOGENOUS DNA PREVENTION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 29 OF 57

## C. Sample Processing

Exemplar samples are processed separately from evidence samples. Also, only one sample is processed at a time using single-use disposable supplies whenever possible (e.g., pipette tips), and scissors/tweezers are thoroughly cleaned between each sample.

## D. Personal Protective Equipment (PPE)

PPE is designed to protect employees from serious workplace injuries or illnesses resulting from contact with chemical, reagents, or biological hazards. PPE includes a variety of devices and garments such as goggles, gloves, lab coats, etc. Proper PPE must be worn during analysis, and required PPE may vary from location to location depending on the hazards of the area. While PPE is designed to protect employees, it can also prevent the transfer of DNA from employees to work surfaces or evidence.

## E. Contamination Prevention Equipment (CPE)

CPE is designed to prevent the occurrence of exogenous DNA in samples. While all PPE are considered as CPE, not all CPE can be considered as PPE. For example, in clean-rooms of the laboratory where high sensitivity DNA testing takes place, the wearing of booties or bouffant caps is to prevent the transfer of DNA from employees. CPE must be worn when designated and available. If not available, employees must first seek permission to work in that area from the appropriate Technical Leader and exercise extreme caution to maintain a clean environment.

## F. Identification

Exogenous DNA may be indicated by 1) the presence of signal in reagent blanks, 2) the presence of extraneous alleles in positive controls, or 3) the presence of extraneous alleles in case samples. The confirmation of exogenous DNA may reflect a system failure or contamination of the samples by an outside source. The source may be equipment, reagents, the working environment, laboratory/law enforcement personnel, or an analytical error. It can either be a single isolated event (such as cross-contamination between two samples) or it can be persistent (such as dirty reagents or equipment). To remedy a single isolated event, the appropriate extraction, quantitation, amplification and/or STR analysis is repeated.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

EXOGENOUS DNA PREVENTION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 30 OF 57

To aid in the identification of exogenous DNA, the LAB TYPES DATABASE procedure is used.

The Quality Assurance Manager and/or the appropriate Technical Leader must be notified if exogenous DNA is detected. The source of this DNA should be identified, if possible, and eliminated. For persistent events, the QUALITY INCIDENT REVIEW procedure must be followed to prevent the recurrence of the problem.

## G. Interpretation and reporting

Samples containing exogenous DNA must be interpreted and reported carefully. This is further discussed in the GENERAL GUIDELINES FOR DNA CASEWORK procedure.

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### Revision History:

February 9, 2010 – Initial version of procedure.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

LAB TYPES DATABASE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 31 OF 57

## Lab Types Database

### GUIDING PRINCIPLES AND SCOPE

“Lab Types” is a DNA database that contains the DNA profiles of individuals who have access to laboratory space and/or may come into contact with an item of evidence prior to or during processing. It contains locally- and nationally-recognized exogenous DNA profiles. This database is a part of the Quality Assurance Program of the laboratory and must be searched in order to assure that no casework DNA profile was contributed by someone during or after the investigation.

The individuals included in Lab Types include past and present personnel of the OCME, members of housekeeping staff, equipment vendors, select members of NYPD, and various visitors to the laboratory. Any DNA profiles that link cases together but are found to be exogenous will be kept in Lab Types under a contaminant listing.

This procedure describes the collection, identification, processing, and disposition of samples used to create the DNA profiles stored in the database. It also describes the processes for the operation and maintenance of the database as well as how the database is used by casework analysts.

### PROCEDURE

#### A. Sample Collection

1. All samples collected internally for Lab Types processing must be collected by an authorized individual. The OCME Human Resource Department most often collects and records each swab taken. Swabs are then sent to Forensic Biology for Processing.
2. The proper consent form must be completed by the donor prior to the collection of the swabs. This form will be stored with the Missing Persons/Exemplar Group.
3. A five-digit sample ID number is generated for each donor. The five-digit ID number meets the following conditions:
  - i. It falls within the numerical range 00000 to 99999, inclusive
  - ii. It is generated randomly each time a new swab is collected.
  - iii. It is unique to all other assigned ID numbers, past or present.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

LAB TYPES DATABASE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 32 OF 57

4. This number is placed on a large coin envelope that is also labeled with the donor's name. The information is recorded in Lab Types. This number becomes the sample identifier.
5. Lab Types samples are classified as reference materials.

## B. Sample Processing

1. Lab Types samples can be processed along with casework exemplar samples.
2. After cutting, the swabs are returned to their envelopes. In most cases, these envelopes are placed in the appropriate container for long-term storage. For situations where samples are not to be stored by Forensic Biology, see the *Sample Disposition* section.
3. Extraction, quantitation, amplification and STR analysis are performed identically to casework exemplar samples. The results are sent to the Lab Types Custodian.

## C. Sample Disposition

1. Lab Types samples and extracts are stored like all other exemplar swabs. In certain circumstances, a swab and extract may need to be destroyed or returned to an individual.
  - i. NYPD swabs and extracts will be returned to the NYPD Integrity Control Officer.
2. To return a sample, the envelope is cut open so that the Eppendorf tube containing the sample extract can be inserted along with the swabs. The five-digit ID number written on the envelope is obscured or removed.
3. In circumstances where samples need to be destroyed, the swabs and extract can be disposed of appropriately.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

LAB TYPES DATABASE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 33 OF 57

## D. Database Maintenance

1. The Lab Types Custodian is in charge of keeping the main Lab Types Database up to date with all relevant information as results arrive.
2. The information is maintained as an Access database and must include, but is not limited to:
  - i. ID number
  - ii. department/agency/employer of donor
  - iii. Date of swab receipt
  - iv. Date and time of extraction, quantitation, and amplification
  - v. quantitation value
  - vi. STR run name
  - vii. DNA profile

## E. Lab Types Reference Databases

1. Due to the nature of the information kept in the main Lab Types Database, the full version is not suitable for general usage by analysts for comparison to evidence profiles. For this reason, copies of the main Lab Types Database are created with various data fields deleted or hidden from view.
2. Two versions of the main Lab Types Database are periodically created for routine use by analysts or managers.
  - i. One version contains only the ID numbers and the corresponding DNA profiles and is designed for use by analysts for comparison with casework DNA profiles.
  - ii. A second version is designed for use by management, and has ID numbers, DNA profiles, and names of sample donors.
3. Each version is spot-checked and write-protected prior to placement online.
  - i. To spot-check a truncated version of the database, an authorized analyst other than the Lab Types Custodian checks the database entries against electropherograms of the samples.
  - ii. After this has been completed, the copies are created and write-protected. These copies are then directed to the Lab Types Manager for approval and placement on the network for general usage.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

LAB TYPES DATABASE		
DATE EFFECTIVE 08-14-2015	APPROVED BY Quality Assurance Manager	PAGE 34 OF 57

## F. Searching the Lab Types Database

1. The Lab Types databank in Access can be sorted by genotype at each locus.
2. Double click *LabTypes:Table* on the left side of the screen if the table does not automatically populate.
3. There are two ways to search: manual and filtered. In both tables, profiles are automatically sorted in numerical order from top to bottom across all columns.
4. **Manual Search.** To search manually, an analyst scrolls down until they find the genotype at the locus in the first column.
5. **Filtered Search.** To perform a filtered search, click on arrow next to the locus name at the top of the column. Check each allele or allele combination you wish to search

This process can be done with as many subsequent loci as necessary. To reset the filter and display the entire database again, click *Toggle Filter* on the Home toolbar

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### Revision History:

February 9, 2010 – Initial version of procedure.

April 30, 2012 – Revised the “Guiding Principles and Scope” section. Lab Types contains both locally- and nationally-recognized exogenous DNA profiles and are kept under a “contaminant” listing.

July 16, 2012 – Specific terminology was removed and replaced with generic terminology to accommodate LIMS.

October 1, 2012 – A Target Date of 60 days for Lab Types samples was added to “Section A – Sample Collection.”

August 14, 2015 – Updated section to reflect current practices.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PREVENTIVE ACTION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 35 OF 57

## Preventive Action

### GUIDING PRINCIPLES AND SCOPE

Preventive action is a pro-active process to identify opportunities for improvement and potential sources of non-conformities rather than a re-active process to the identification of problems or complaints. Aside from the review of the operational procedures, preventive action may involve analysis of data including trend and risk analyses and proficiency test results.

This document describes the Department's procedure to identify potential preventive actions, either technical or concerning the Management System, and the steps to be taken to deal with the issues identified.

### PROCEDURE

- Any staff member that becomes aware of potential sources of non-conformities in laboratory operations informs their immediate supervisor and/or Assistant Director as soon as practicable.
  - Immediate supervisors notify their Assistant Director if the AD was not part of the initial notification. The initial process to communicate potential preventive actions up the chain-of-command ensures that any follow-up action is implemented sooner, rather than later.
- The immediate supervisor and/or Assistant Director investigates the potential problem and conducts a preliminary review of the root cause(s) of any potential non-conformity to determine if action is necessary. The appropriate Technical Leader (if the potential problem is a technical problem), the Quality Assurance Manager, and/or other supervisors/managers may be consulted for assistance.
  - If the investigating supervisor/manager does not agree that a potential problem exists, no further action is necessary.
- If the investigating supervisor/manager agrees that a potential problem exists, and if a root cause of the potential non-conformity is determined, the immediate supervisor and/or Assistant Director develops a plan of action to deal with the issue. This may include a change in technical procedures and/or the initiation of new guiding principles. The plan of action shall include the initiation of controls to ensure that the preventive actions are effective. A description of the potential problem, root cause, and plan of action is documented on a **Preventive Action Form** and submitted to the Quality Assurance

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PREVENTIVE ACTION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 36 OF 57

Manager. If the preventive action is of a technical nature, the Quality Assurance Manager will forward the form to the appropriate Technical Leader for review.

4. If the preventive action is of a technical nature, the appropriate Technical Leader either approves the plan or decides on an alternate arrangement.

If the preventive action concerns a potential non-conformity in the Management System, the Director or his/her designee either approves the plan or decides on an alternate arrangement.

5. The Preventive Action Form and any associated documentation (such as Manual Change Forms, copies of emails, etc.) are filed with the Quality Assurance Unit.
6. The Quality Assurance Manager reviews the Preventive Action Form within six months to determine if the preventive action plan that was put into place has been effective.
  - a. The Quality Assurance Manager records their evaluation of effectiveness on the Preventive Action form, e.g. a notation that none of the anticipated non-conformities had occurred.
  - b. If the action plan is determined to have been effective, the preventive action is considered to be complete.
  - c. If the action plan is determined not to have been effective, the Quality Manager will determine whether the changes made as a result of the action plan need to be discontinued or revised.

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## Revision History:

February 9, 2010 – Initial version of procedure.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PROFICIENCY TESTING PROGRAM		
EFFECTIVE DATE 08-14-2015	APPROVING AUTHORITY DNA TECHNICAL LEADERS	PAGE 37 OF 57

## Proficiency Testing Program

### GUIDING PRINCIPLES AND SCOPE

Proficiency tests are given to qualified analysts to evaluate both their individual competence and the quality performance of the laboratory. Proficiency tests must be analyzed using only approved methods and/or procedures. While there are several types of proficiency tests, the Department of Forensic Biology utilizes open-external proficiency testing and blind reanalysis proficiency testing.

The proficiency testing program is designed to meet the requirements of ASCLD/LAB and the Quality Assurance Standards for Forensic DNA Testing Laboratories. The external proficiency testing program is not just a requirement; it is also a quality assurance measure used to monitor performance and identify areas in which improvement may be needed.

External DNA proficiency tests are obtained from New York State and ASCLD/LAB approved proficiency test providers, for example, Collaborative Testing Service (CTS), Orchid Cellmark (IQAS), and the College of American Pathologists (CAP).

Serology results are reported on DNA tests obtained from CTS.

### PROCEDURE

#### A. DNA Open-External Proficiency Testing Program

1. All analysts, technical reviewers, and technicians undergo semiannual external proficiency testing to the full extent in which they perform each technology in casework. Technology refers to the type of forensic DNA analysis performed (i.e. STR, Y-STR, mtDNA.) The program is administered in an open proficiency-testing format and in accordance with the FBI Quality Assurance Standards for Forensic DNA Testing Laboratories.
2. One test is assigned to each participant in the first six months of the calendar year and the second test is assigned in the last six months of the calendar year.
  - a. The interval between consecutive tests must be at least four months and cannot exceed eight months.
  - b. **The laboratory uses the assigned date/start date to calculate the interval between tests.**

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PROFICIENCY TESTING PROGRAM		
EFFECTIVE DATE 08-14-2015	APPROVING AUTHORITY DNA TECHNICAL LEADERS	PAGE 38 OF 57

- c. Newly qualified individuals enter the external proficiency testing program within six months of the date of their qualification.
3. The scheduling of external proficiency tests is completed by a member of the Quality Assurance Unit prior to the start of each calendar year. While minor changes may be made during the year (test vendor, paired analyst, addition/removal of personnel, etc.), the schedule of each analyst/technician is not changed unless a change is necessary due to an extended leave of absence.
4. All specimens of an external proficiency test are analyzed according to current standard operating procedures. However, some exceptions are made in order to comply with the FBI Quality Assurance Standards for Forensic DNA Testing Laboratories. For example, the following sample types, which during normal casework analysis might only be tested in one or two multiplex reactions, must be amplified at all CODIS core loci or CODIS core sequences and tested in all applicable technologies (Autosomal STR, Y-STR, and/or Mitochondrial DNA) to the full extent that the analyst participated in casework:
  - 1) Excluded suspects
  - 2) Mixtures, even if there are other clean profiles
  - 3) Epithelial cell fractions from an unknown stain or from a body orifice swab, even if the results match the victim type.
5. The laboratory utilizes a team approach for casework testing. Therefore, proficiency tests are conducted in the same manner. However, each individual is proficiency tested at least once per year in each methodology to the full extent of his or her participation in casework.

Methodology refers to analytical procedures used to support a DNA-typing technology [i.e. extraction methods (manual v. automated,) quantification methods, typing test kits and instrument platforms]. The extent in which each individual participates in casework may be team dependent.

Individuals who perform STR, YSTR, and/or mtDNA amplification, analysis and/or review must perform these skills twice per year per technology.

- a. Unlike other titles, Criminalist Level I's are competent only in selected areas of the analytical process and their competency differs between the different teams within the laboratory. Criminalist I's cannot interpret the final DNA typing data or prepare an associated written scientific report.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PROFICIENCY TESTING PROGRAM		
EFFECTIVE DATE 08-14-2015	APPROVING AUTHORITY DNA TECHNICAL LEADERS	PAGE 39 OF 57

Thus, their participation in proficiency tests is limited to the methodologies that they are competent in and they are paired with a DNA Analyst on proficiency tests.

- b. Individuals using both manual and automated methods are proficiency-tested in each at least once per year.
6. A laboratory report to summarize the results of the Proficiency Test is written by the DNA interpreting analyst. The DNA interpreting analyst is also responsible for completing any vendor paperwork to document the results. The DNA interpreting analyst must ensure that the data transcribed to the vendor's paperwork is accurate. The proficiency test file is then forwarded to the analyst's supervisor, manager, and/or designee for a full technical review.
7. In addition to conducting a full technical review of the proficiency test file, the reviewer(s) must also review the completed vendor's paperwork to ensure that data has been transcribed correctly.
8. After the proficiency test has been completed (including a full administrative review), the DNA interpreting analyst assigned to the proficiency test is responsible for delivering the test results to the test vendor. The delivery method may vary from vendor to vendor, but is typically either by fax or e-mail.
9. After official results have been received by the proficiency test provider, a designated Quality Assurance Unit member grades the tests.
  - a. Non-administrative discrepancies on proficiency tests that affect typing results and/or conclusions should be reported to the appropriate Technical Leader at the time of discovery. If confirmed, the Technical Leader must inform the CODIS Custodian/Supervisor so that appropriate follow-up action can be initiated. A formal QUALITY INCIDENT REVIEW may be required.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PROFICIENCY TESTING PROGRAM		
EFFECTIVE DATE 08-14-2015	APPROVING AUTHORITY DNA TECHNICAL LEADERS	PAGE 40 OF 57

10. All proficiency-test participants are informed of their final test results. Participants are required to sign the appropriate area on the Proficiency Test Evaluation Form to document that they have received and have been informed of the final test results.
11. After the grading of all proficiency tests within the series, the designee informs the appropriate Technical Leader of the results of all participants.

## B. Serology Open-External Proficiency Testing Program

Serology is a sub-discipline of the Biology discipline (as per ASCLD/LAB). The laboratory will endeavor to arrange for each employee to annually complete a serology proficiency test, but it is not required to do so.

Forensic Biology proficiency tests purchased from CTS allows the participant to report results for serology tests as well as for DNA testing. Therefore, serology proficiency testing is satisfied in this manner. The management of this test is identical to the management of DNA external proficiency tests – tests are reported, reviewed, and participants are evaluated in the same manner.

## C. Blind Re-analysis Proficiency Testing Program

1. The Blind Re-analysis Proficiency Testing Program is a quality assurance program where a previously examined sample is re-examined by a different analyst to check for correctness of the initial examination and results.
2. **DNA Blind Reanalysis Program.** The Quality Assurance Unit is responsible for reanalyzing DNA samples, reviewing the results, and comparing them to the original analyses.
  - a. Each month, a minimum of two (2) exemplar samples are selected from cases completed within the previous year.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

PROFICIENCY TESTING PROGRAM		
EFFECTIVE DATE 08-14-2015	APPROVING AUTHORITY DNA TECHNICAL LEADERS	PAGE 41 OF 57

- b. Each sample is submitted for extraction, quantitation, amplification (in at least one casework multiplex system), analyzed for STR results, and the results compared to the original results. Re-examined results are documented separate from the case file and maintained as a record by the Quality Assurance Unit.
  - c. A second reanalysis must be performed if the results are not concordant. All follow-up actions must be documented and maintained.
- 3. **Serology Blind Reanalysis Program.** The laboratory has a blind serology re-analysis program for negative cases. The purpose of this program is to ensure that negative serology results are accurate.
  - a. Approximately 25% of negative sexual assault kits are selected by a Quality Assurance member for re-analysis. The re-analysis must occur prior to the release of any report.
  - b. Re-analysis of negative serology cases is conducted by casework analysts that are not involved in the original analysis. Each sample within the case that was previously analyzed is re-analyzed to ensure consistent results and checked for correct itemization. Examination notes and confirmatory-test results are compared. Original and re-examined results are retained in the case file.
  - c. If discrepancies between results occur, the Quality Assurance Manager and/or the Quality Assurance Supervisors must be contacted to determine what follow-up action is necessary. All follow-up actions must be documented and maintained.

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## Revision History:

- February 9, 2010 – Initial version of procedure.
- March 30, 2012 – Removed the requirement to use the Proficiency Test Review Form to document the review of proficiency tests (consistent with current practice).
- July 16, 2012 – Specific forms and worksheets were removed and replaced with generic terminology to accommodate LIMS.
- August 14, 2015 – Updated percentage of cases chosen for Blind Reanalysis per year, as well as changing “QA supervisor” to “designated QA member”.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

QUALITY INCIDENT REVIEW		
DATE EFFECTIVE 04-01-2014	APPROVED BY QUALITY ASSURANCE MANAGER	PAGE 42 OF 57

## Quality Incident Review

### GUIDING PRINCIPLES AND SCOPE

Action must be taken when major departures from the policies and procedures in the Management System have been identified. These quality incidents shall be identified and reported so that appropriate follow-up action can be implemented. The identification of problem areas and subsequent preventive actions will improve the quality of our Department.

This document describes the Department's process for dealing with quality incidents when major departures from the policies and procedures in the Management System have been identified. Problems or difficulties can arise in all phases of laboratory operations, and these must be evaluated and dealt with appropriately. Listing each potential problem is impractical, and this topic is considered in general terms.

Technical errors or problems related to casework testing are dealt with as per the CONTROL OF NON-CONFORMING TESTING procedure. The procedure provides direction with respect to when such problems must be dealt with via a Root-Cause Analysis.

This procedure ensures that, when required, our accrediting bodies and/or appropriate entities are notified in a timely manner.

### PROCEDURE

A problem with the Management System of the laboratory may be identified through a variety of activities such as internal or external audits, management reviews, feedback from customers, and from staff observations.

Not every quality incident or departure from Management System policies and procedures is serious enough to require a Quality Incident Review.

It is impossible to anticipate all situations in which a Quality Incident Review must be conducted; therefore, sound judgment is required in determining the extent and level of reporting and documentation required.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

QUALITY INCIDENT REVIEW		
DATE EFFECTIVE 04-01-2014	APPROVED BY QUALITY ASSURANCE MANAGER	PAGE 43 OF 57

Issues which require a Quality Incident Review include, but are not limited to:

- Systemic non-conformance with the policies and procedures in the Management System (e.g., failure to properly document staff qualifications and training)

The Quality Assurance Manager should be consulted if there is any question as to whether a Quality Incident Review is required.

## A. Quality Incident Reporting

1. All staff members are responsible for reporting apparent quality incidents that come to their attention.
  - i. All quality incidents are reported to the staff member's immediate supervisor.
  - ii. Supervisors or managers who become aware of any non-technical quality incidents, either directly or through notification from other staff members, must proceed to Step 2.
  - iii. Any member of staff who believes that the potential quality incident is of major concern, but is concerned about confidentiality, may inform the Quality Assurance Manager immediately and directly
2. The supervisor or manager investigates the issue to determine the details of the potential problem.
3. If the initial investigation indicates that a quality incident occurred, but that a formal Quality Incident Review is not needed, the investigating supervisor/manager shall inform the Quality Assurance Manager of the incident via email.
4. If the initial investigation indicates that a formal Quality Incident Review is needed, the investigating supervisor/manager consults with the Quality Assurance Manager.
5. The Quality Assurance Manager determines whether to proceed with a formal Quality Incident Review.
6. The Quality Assurance Manager informs the investigating supervisor/manager of the decision.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

QUALITY INCIDENT REVIEW		
DATE EFFECTIVE 04-01-2014	APPROVED BY QUALITY ASSURANCE MANAGER	PAGE 44 OF 57

7. The decision not to proceed with a formal Quality Incident Review does not prevent a supervisor or manager from conducting other follow-up action, e.g., counseling of an individual.

## B. Quality Incident Review

1. The Quality Assurance Manager assigns a supervisor or manager to conduct the Quality Incident Review.
2. The Quality Incident Review (QIR) form guides the steps in the process.
3. The incident is described in detail on the QIR, including the effect(s) of the discrepancy.
4. The assigned supervisor/manager conducts an investigation to determine the **root cause** of the incident. The root cause(s) may not be obvious and thus a careful analysis of all potential causes of the problem is required. Potential causes could include, but are not limited to, problems with:
  - customer requirements
  - the samples
  - sample specifications
  - methods and procedures
  - staff skills and training
  - consumables, or
  - equipment and its calibration.

The assigned supervisor/manager conducting the QIR shall use the OCME Root Cause Analysis Procedure as a guide to determine the root cause of the incident.

5. Follow-up actions are proposed to correct the immediate problem and minimize the potential for recurrence of the problem. Corrective actions may include, for example, personnel counseling, modifying procedures or forms, etc. The follow-up action plans should also include:
  - The parties responsible for implementing the corrective actions
  - The monitoring that will be conducted to ensure that the proposed actions have been effective. Very serious and/or systemic issues may require follow-up audits of the affected areas of activity.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

QUALITY INCIDENT REVIEW		
DATE EFFECTIVE 04-01-2014	APPROVED BY QUALITY ASSURANCE MANAGER	PAGE 45 OF 57

6. The QIR is forwarded to the Quality Assurance Manager for review and approval of the proposed corrective actions.

## C. Close-out of Quality Incident Reviews

1. The completion of corrective actions is documented on the QIR.
2. If the monitoring activities indicate that the initial follow-up actions were insufficient to address the quality incident, the Quality Assurance Manager may initiate additional follow-up actions.
3. The Quality Assurance Manager determines which incidents must be disclosed to accrediting bodies and/or appropriate entities.
4. The QIR is “closed” when monitoring activities are completed and all individuals agree that corrective action(s) have been satisfactorily implemented and effectively addressed the quality issue.
5. The QIR and any supporting records are filed with the Quality Assurance Unit.

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### Revision History:

February 9, 2010 – Initial version of procedure.

September 24, 2010 – Added step in procedure to document quality incidences that do not rise to the level of a Quality Incident Review on the Non-conformity Reporting Form.

April 1, 2014- Any references to technical non-conformities have been removed from the QIR and will now solely be addressed via the Non-Conforming Work Procedure.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 46 OF 57

## Reagents

### GUIDING PRINCIPLES AND SCOPE

A **reagent** is any substance used because of its chemical or biological activity. Reagents are used directly, or at a dilution, in a given analytical procedure. Reagents are different than *chemicals*, which are used in the preparation of in-house reagents.

Only reagents suitable for the methods employed may be used in the Department of Forensic Biology. This procedure describes in general terms the requirements for the documentation and quality control of commercial reagents and for the formulation, documentation, and quality control of in-house reagents. The last section in this document is a list of the reagents used by the Department.

### PROCEDURE

Reagents are classified into two general categories:

A **critical reagent** is determined by empirical studies or routine practice to require testing on established samples before use on evidentiary or casework reference samples in order to prevent unnecessary or irreparable loss of sample. "Critical reagents" includes a variety of test kits or systems used in DNA testing.

A **non-critical reagent** is a reagent whose failure to work properly will not cause irreparable loss of sample. Therefore, the use of a QC test procedure to check the reliability of the reagent prior to its use in casework is not an absolute requirement, but will be performed by the Department on a reagent-by-reagent basis.

Reagents are prepared **in-house** or are obtained **commercially**.

Personnel preparing reagents, and those who use reagents, are to exercise care at all times to ensure that no exogenous DNA will be introduced to a stock reagent.

#### A. Reagents Prepared In-House

- 1) Reagents are prepared in-house according to an approved formula or procedure. Reagent preparation is usually performed by a member of the Quality Assurance Unit.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 47 OF 57

- 2) A **reagent sheet** form exists for every reagent prepared in the laboratory and is used as a guide for the preparation of the reagent.
- 3) Each reagent record contains the following information:
  - i. the identity of the reagent
  - ii. date of preparation
  - iii. identity of individual preparing the reagent
  - iv. standard batch size
  - v. ingredients of the reagent
  - vi. data entry section
4. Some reagent records (such as critical reagents) may also include:
  - i. lot numbers
  - ii. expiration dates (see step 6)
  - iii. quality control procedures (aka, “reliability checks”) to be performed and passed before the reagent is released for use in the laboratory.
5. Reagents prepared in the laboratory are labeled with, at a minimum:
  - i. the identity of the reagent
  - ii. the lot number
  - iii. the expiration date (see step 6)

When a reagent is aliquotted into tubes that are too small to be labeled with all of the required information, each tube is marked with the identity of the reagent and its lot number and stored in a “cryobox” that is labeled with the required identifying information listed above.

6. The expiration date given is usually one year from date of make/aliquot or the earliest expiration date of the reagents being used, whichever comes first. This may also be stated in each reagent forms.
7. Staff is notified via email by the Quality Assurance Unit regarding reagents that are expiring.

## B. Commercial Reagents

2. Commercial reagents include, but are not limited to, kits for DNA quantitation and genetic typing.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 48 OF 57

3. A **Raw Materials** form exists for each commercial reagent that requires quality testing prior to use in casework. The applicable quality control procedure is contained on the form.
4. Commercial reagents are labeled with, at a minimum:
  - i. The identity of the reagent
  - ii. The expiration date as provided by the manufacturer or as determined by the laboratory.
    - 1) If identical reagents with the same lot number are assigned different expiration dates by the manufacturer, then the expiration date will be extended to the latest date provided that it passes quality control testing.

For example, Lot #1234 of a reagent was received on June 1, 2011 (Bottle A) and has a manufacturer assigned expiration date of June 1, 2012. A second bottle of Lot #1234 was received on December 1, 2011 (Bottle B) and has a manufacturer-assigned expiration date of December 1, 2012. Since the manufacturer supports the use of this particular lot of reagents until December 1, 2012, the expiration date of Bottle A will be extended to December 1, 2012 provided that Bottle B passes quality control testing.
    - 2) Commercial reagents without an expiration date provided by the manufacturer shall expire two years *after receipt* unless otherwise indicated.

## C. Reagent Quality Control Testing

Quality control (QC) tests are reliability checks and may be used by the Department to ensure that reagents are performing as expected. If needed, these tests must be completed prior to the reagent being used in actual casework. A reliability check may be a combination of several quality control tests and, for ease of classification, are assigned QC testing procedure numbers. If a reagent sheet lists a "procedure" for its quality control test, then the reagent must pass all the quality control tests listed below. If it only lists a specific "QC" number, then the reagent must pass that quality control test only.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 49 OF 57

	QC Tests Included	Analysis
<b>Procedure 1</b>	QC615, QC616 or QC620	Real Time Quantitative PCR
<b>Procedure 2</b>	QC240, QC350	PCR Amplification and STRs
<b>Procedure 3</b>	QC145A, QC615/620, QC350	Organic Extraction, Real Time Quantitative PCR, PCR Amplification, and STRs
<b>Procedure 4</b>	QC145/165, QC160, QC615/620, QC350	Chelex/M48 Extraction, Real Time Quantitative PCR, PCR Amplification, and STRs
<b>Procedure 5</b>	QC350	3130xl STRs

## D. Reagent Records

Reagent records, such as reagent sheets and Raw Materials Forms are a form of Quality Record, and shall be stored in accordance to the guiding principles and procedures that govern such records. See CONTROL OF RECORDS in the Administrative Manual for further information.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 50 OF 57

## E. REAGENTS USED BY THE DEPARTMENT

This section shows a list of reagents used in the Department of Forensic Biology. The list includes reagents prepared in-house as well as commercial reagents. Each reagent is classified as “Critical” or “Non-Critical”.

REAGENT	CRITICAL
a-Amylase powder from Human Saliva	N
Acid Phosphatase Test Reagent	Y
Alkaline Substrate Buffer	Y
Agilent DNA 1000 Kits	N
AmpFSTR Identifiler PCR Amplification Kit	Y
AmpFSTR MiniFiler PCR Amplification Kit	Y
AmpliTaq Gold DNA Polymerase Kit (all components)	Y
BigDye Terminator Cycle Sequencing Kit	Y
Centrisep columns, strips, and plates	N
Chelex, 20%	Y
Chelex, 5%	Y
Deoxynucleotide Triphosphates 2.5 mM (dNTPs)	Y
Digest Buffer	Y
Dithiothreitol (DTT) 1M	Y
EDTA, 0.5 M	N
EDTA, 0.5M for WTC	Y
ExoSAP-IT	Y
Fish Sperm DNA	Y
Genetic Analyzer Buffer (ABI)	N
HiDi Formamide	N
Human Leukemia 60 (HL60)	Y

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 51 OF 57

REAGENT	CRITICAL
Hydrogen Peroxide, 3%	N
Kastle-Meyer (KM) Reagent	Y
MagAttract DNA Mini M48 Kit (Qiagen)	Y
Magnesium Chloride (MgCl <sub>2</sub> )	N
Nuclear Fast Red	Y
Organic Extraction Buffer	Y
PBS for Chelex Extraction	Y
PBS for Nail Extraction, 25mM EDTA	Y
PBS Solution for Seratec (PBS tablets)	Y
PBS Solution, Irradiated (LCN DNA)	Y
Phase lock gel tubes	N
Phenol Chloroform Isoamyl Alcohol (PCIA)	Y
Picric Indigo Carmine (PIC)	Y
POP-4	N
POP-6	N
Poly A RNA	Y
Primer, FBI – A1, B1, C1, D1, C2, D2, A4, B4, HVIF, HVIR, HVIIF, HVIIR (mtDNA)	Y
Proteinase K solution	Y
Quantifiler Trio DNA Quantification Kit	Y
Roche Primer and Reaction Mix	Y
Saline (0.85% NaCl)	N
SDS, 2%	N
SDS, 20%	Y
SDS, 0.01%, 0.05%, and 1% (LCN DNA)	Y

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 52 OF 57

REAGENT	CRITICAL
Seratec PSA Semiquant Kits	Y
Seratec Amylase Forensic Test	Y
Sequencing Loading Buffer	Y
Sodium Acetate, 0.1 M	N
Sterile Deionized Water	Y
Terg-a-zyme	N
TAE, 1X	Y
TBE buffer	N
Tris-EDTA, 1X	Y
Tris-HCl, 1M (pH 8.0)	N
UltraPure Water	Y
Xylene	N
Yfiler™ PCR Amplification Kit	Y

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

REAGENTS		
DATE EFFECTIVE 12-24-2015	APPROVED BY Quality Assurance Manager	PAGE 53 OF 57

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## Revision History:

February 9, 2010 – Initial version of procedure.  
October 28, 2010 – Added the MagAttract DNA Mini M48 Kit and the MiniFiler PCR Amplification Kit to the list of reagents.  
December 29, 2011 – Revised Section B.3 to clarify how the laboratory determines the expiration dates of commercial reagents.  
July 16, 2012 – Portions revised to generalize terminology to accommodate LIMS.  
August 20, 2012 – Revised Section B.3 to clarify how the laboratory determines the expiration dates of commercial reagents.  
April 1, 2014 – Revised Section A to include clarification of expiration dates of Reagents made in-house. Replaced YM1 STR with Yfiler™ PCR in Critical Reagent List.  
November 24, 2014 – Updated wording for reagent Expiration dates. Added EDTS, 0.5M for WTC to the critical reagent list and replaced Irradiated Water with UltraPure Water.  
February 2, 2015- Updated Section E. Added Quantifiler and Seratec Reagents, Removed outdated reagents from the Reagent List.  
May 1, 2015 – Updated QC procedures to reflect use of QC615 (QC with Quantiflier Trio).  
August 14, 2015 – Updated procedure 1 with new QC616 protocol and corrected reference to location of the Control of Records procedure.  
December 24, 2015 – Removed Rotorgene reagents from Reagent List as this procedure is no longer on-line.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

VALIDATION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 54 OF 57

## Validation

### GUIDING PRINCIPLES AND SCOPE

Validation is the process by which a procedure is evaluated to determine its efficacy and reliability for forensic casework analysis. It is the accumulation of test data within the laboratory to demonstrate that established methods and procedures perform as expected. Only validated methods and procedures may be used with casework samples.

This is different from a performance check, which is a quality assurance measure to assess the functionality of laboratory instruments, equipment, and software that affect the accuracy and/or validity of forensic sample analysis.

The validation process identifies the critical aspects of a procedure which must be carefully controlled and monitored. Validation studies must have been conducted by the Department of Forensic Biology prior to the adoption of a procedure by our laboratory. This procedure describes the requirements of the validation process.

### PROCEDURE

All staff members are encouraged to propose new technologies, methodologies, or procedures to be used in casework. Proposals may be forwarded to the Forensic Biology Future Technologies Planning Team. The Director shall make a final determination on whether or not to validate any proposed new technology, methodology, or procedure.

Validations are a planned activity, and the exact tests of one validation may differ from another depending on the new technology, methodology, or procedure being tested. The appropriate Technical Leader shall be consulted to determine which studies must be conducted to ensure efficacy and reliability for forensic casework use. If the technology, methodology, or procedure concerns DNA testing, the Technical Leader must ensure that the appropriate tests, as listed in the FBI's Quality Assurance Standards for Forensic DNA Testing, are conducted.

Validation plans may differ from the initial assessment of the Technical Leader. They may be updated as development proceeds.

While not required, prior to starting any validation, a preliminary assessment may be done to ensure the time and effort that will be dedicated to the validation will be worthwhile.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

VALIDATION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 55 OF 57

## A. Developmental Validation

1. Developmental validation is the acquisition of test data and determination of conditions and limitations of a ***new or novel*** methodology for use on forensic samples.
2. If another laboratory's developmental validation studies are being used, appropriate documentation or citations for these studies must be available.
3. Developmental validation studies must include the following, where applicable:
  - i. Testing using case-type samples, including samples from adjudicated cases or mock samples that mimic casework samples
  - ii. Characterization of genetic marker
  - iii. Sensitivity, stability, and species specificity studies
  - iv. Reproducibility studies
  - v. Population studies such as allele frequency distributions and independence of the population databases
  - vi. Mixture studies
  - vii. Precision and accuracy studies
  - viii. PCR-based studies, including reaction conditions, assessment of differential and preferential amplification, effects of multiplexing, assessment of appropriate controls, and product detection studies.
4. All developmental validations conducted by the Department must include an executive summary, which summarizes all the studies conducted. The executive summary must include specific recommendations (such as settings, quality assurance parameters, interpretation guidelines, or mixture interpretation guidelines) and must include a statement as to whether the method is fit for the intended use. While not required, it is recommended that each study conducted have an individual summary of results.

## B. Internal Validation

1. Internal validation is an accumulation of test data within the laboratory to demonstrate that ***established*** methods and procedures (such as forensic DNA methods or procedures that are published in peer reviewed articles) perform as expected in the laboratory.

# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

VALIDATION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 56 OF 57

2. Prior to implementing a new or revised methodology or procedure, the Department must first demonstrate the reliability of the method or procedure internally. This includes changes in detection platform, changes in DNA test kits, or the implementation of new body-fluid identification procedures. Internal validation studies must be sufficient to support and document the reliability of the method or procedure and must include the following, where applicable:
  - i. Testing using known samples
  - ii. Testing using non-probative evidence samples or mock evidence samples
  - iii. Reproducibility and precision
  - iv. Sensitivity and stochastic studies
  - v. Mixture studies
  - vi. Contamination assessment
3. As a result of the internal validation studies, quality assurance parameters, interpretation guidelines, and mixture interpretation guidelines (where applicable) shall be defined.
4. The documentation of an internal validation includes an executive summary, which summarizes all the testing conducted. The executive summary must include specific recommendations (such as settings, quality assurance parameters, interpretation guidelines, or mixture interpretation guidelines) and a statement as to whether or not the method is fit for the intended use. While not required, it is recommended that each study conducted have an individual summary of results.

## C. Review and Approval of Validation

1. Completed validation project packages are submitted to the appropriate Technical Leader for review and approval. The package includes:
  - i. Test records and all required summaries
  - ii. Draft technical procedure
2. All validations must be reviewed and approved by the appropriate Technical Leader before the technology and/or procedure is used in casework.

**Note:** Approval of a validation does not necessarily denote that a technology or procedure is online for casework. Training needs, budgetary concerns, etc., must be taken into consideration before the technology or procedure is implemented.



# FORENSIC BIOLOGY QUALITY ASSURANCE/QUALITY CONTROL MANUAL

VALIDATION		
DATE EFFECTIVE 02-09-2010	APPROVED BY Quality Assurance Manager	PAGE 57 OF 57

3. At the Technical Leader's discretion, the technology or procedure may be used on select cases prior to lab-wide implementation. However, the technology or procedure are not be used on any casework until standard operating procedures are written and have been approved by the appropriate Technical Leader.

## D. Training

Training commences after approval of the validation by the appropriate Technical Leader. The initial training of analysts can be considered a "dry-run" of the procedure, and the technology, methodology, and/or procedure are not used in casework until all concerns that may be raised during the initial training have been addressed.

## E. Storage of Validation Records

Records of validation studies are stored by the Quality Assurance Unit indefinitely. In general, validations that have been reviewed by an external audit team will be stored on the fourth floor of the DNA Building (Records Storage), while validations that have not been reviewed by an external audit team will be stored within the operational areas of the Quality Assurance Unit. However, general convenience and spacing issues may alter the exact location of any validation study.

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### Revision History:

February 9, 2010 – Initial version of procedure.