This document has been accepted by the Academy Standards Board (ASB) for development as an American National Standard (ANS). For information about ASB and their process please refer to asb.aafs.org. This document is being made available at this stage of the process so that the forensic science community and interested stakeholders can be more fully aware of the efforts and work products of the Organization of Scientific Area Committees for Forensic Science (OSAC). The documents were prepared with input from OSAC Legal Resource Committee, Quality Infrastructure Committee, and Human Factors Committees, as well as the relevant Scientific Area Committee. The content of the documents listed below is subject to change during the standards development process within ASB, and may not represent the contents of the final published standard. All stakeholder groups or individuals, are strongly encouraged to submit technical comments on this draft document during the ASB's open comment period. Technical comments will not be accepted if submitted to the OSAC Scientific Area Committee or Subcommittees.

# Wildlife Forensics- Report Writing Standards



**DRAFT DOCUMENT** 

# Wildlife Forensics- Report Writing Standards

Keywords: wildlife forensics, taxonomic identification, reference collections, reporting

# Abstract

This document provides minimum standards and recommendations for report writing and report content for practicing wildlife forensic analysts. These minimum standards and recommendations are not intended to replace standards in ISO 17025, but are designed to guide analysts in proper report writing and report content. Notes and examples throughout this document offer clarifications and examples of how a lab may meet a specific standard.



# Foreword

# Acknowledgements

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# Wildlife Forensics Report Writing Standards

# 1 Scope

This document describes the information to be provided in reports of wildlife forensic examinations for use in legal proceedings. Requirements for both genetic and morphological examination reports are covered. Forensic reports serve a variety of audiences, and must provide a clear and concise summary of methods and results for the use of the investigator, the court, and the litigants.

# 2 Normative References

The following referenced documents are indispensable for the application of this document.

International Organization for Standardization. 2005. ISO/IEC 17025:2005 General Requirements for the Competence of Testing and Calibration Laboratories.

National Commission on Forensic Science. 2015. Views Document on Documentation, Case Record and Report Contents. (<u>https://www.justice.gov/ncfs/file/818191/download</u>).

Organization for Scientific Area Committees, Wildlife Forensics Subcommittee, Wildlife Forensics General Standards. Currently under review by the AAFS Standards Board (ASB).

# 3 Report Content

Reports shall include information on administrative detail, chain of custody, examination requested, methods, results, and conclusions. Suggested section headings are given below. Additional materials and case information may be included, as appropriate to the particular case.

# 3.1 Administrative Information

- 3.1.1 The report shall include:
  - a) Title of report, specifying type of analysis being reported (for example, Genetics Examination Report). This is distinct from the submitting investigator's case title, which may be noted separately.
  - b) Identity and location of the laboratory performing the analysis.
  - c) A unique case identifier assigned by the laboratory.
  - d) Pagination, including the total number of pages.
  - e) Date of report.
  - f) Name and signature of the author(s) of the report.

NOTE: Verified digital signatures are acceptable.

### 3.2 Chain of Custody Information

3.2.1 The report shall include:

- a) Investigator's Case number
- b) Name of submitter and (if appropriate) submitting agency.
- c) Date the evidence was received at laboratory.
- d) Name of laboratory functional unit (e.g. Evidence Unit) or staff member who initially received the evidence into the laboratory.
- e) The evidence item identifier(s) and submitted descriptions.
- f) Date the analyst received the evidence for the reported analysis.

### 3.3 Examination Requested

3.3.1 The report shall include a section describing the investigator's request(s) for analysis.

EXAMPLE: Species identification, source evaluation, minimum number of individuals.

### 3.4 Case Information

3.4.1 Information provided by the investigator regarding the evidence that was used to formulate the analytical approach and subsequent interpretation shall be noted in the report.

### 3.5 Examinations Conducted

- 3.5.1 The report shall include sufficient detail for another expert to be able to ascertain how the analyses were accomplished and conclusions drawn.
- 3.5.2 The report shall note that complete documentation of the analyses conducted and data collected is maintained in the case record, which is available from the laboratory.
- 3.5.3 The report shall state the technical methods used to reach the reported conclusion.

EXAMPLE: Mitochondrial DNA sequencing or morphological examination.

- 3.5.4 Examinations conducted by any other person or persons shall be noted, including the identity of the person(s) and the nature of the examination.
- 3.5.5 The report shall include information on the reference material on which examination conclusions are based.

EXAMPLE: Such reference material includes databases, specimen collections, and published literature.

3.5.6 In the case of mtDNA analysis, the report shall include the name(s) of the loci on which the genetic conclusions are based.

#### 3.6 Examination Results

NOTE: This section refers to results only, prior to their interpretation.

3.6.1 The report shall include a statement of the results of the examination.

- 3.6.2 If no results were obtained, a statement to that effect shall be included.
- 3.6.3 Results that include a taxonomic category shall use currently accepted scientific names. Common names may be included as well.
- 3.6.4 When DNA sequencing produces meaningful results, the following shall be reported:
  - a) Total length of sequence used in comparison (base pairs).
  - b) The unique identifier of the reference sequence used for comparison, along with the organism's scientific name.
  - c) Percent identity or number of matching base pairs between the evidence sequence and the most similar reference sequence.
- 3.6.5 When DNA sequencing is used for haplotyping comparisons, the following shall be reported:
  - a) The locus/loci with reported results, specifying which items have the same haplotype and which items have different haplotypes.
  - b) For inclusions, indicate statistical support via confidence interval.
- 3.6.6 Source evaluation for individuals using Short Tandem Repeats (STRs)
  - a) When multiple evidence items are to be compared to one another, indicate which evidence items have the same genotype.
  - b) For inclusions, indicate the Likelihood Ratio.

EXAMPLE Indicate the propositions and present the likelihood ratio for these propositions.

- 3.6.7 When conducting population assignment analysis, include a confidence interval.
- 3.6.8 When evaluating parentage, include combined parenting index (CPI) support for inclusions.

NOTE: Statistical support is not necessary when determining minimum number of individuals using either morphology or sequencing or when determining exclusions. When conducting species identification using either morphology or sequencing, statistical support is not appropriate.

#### 3.7 Examination Conclusions

NOTE: Conclusions are an opinion statement requiring an expert's interpretation and evaluation of the results.

3.7.1 The report shall include a statement of the conclusions based on the examinations conducted.

3.7.2 Conclusions that include a taxonomic category shall use currently accepted scientific names. Common names may be included as well.

## 3.8 Optional Additional Information

Depending on the circumstances of particular cases, additional report sections may be appropriate. These may include References Cited (when providing the published literature, databases, or other sources consulted would be helpful to the Court) or a Glossary/Definitions section, when terms used need to be defined to avoid misunderstanding by non-technical recipients of the report.

# Annex A (informative)

# **Example Reports**

The following are examples of a variety of different analytical reports incorporating the standards. They have been annotated with the standards appropriate for each case. Laboratories may use these for guidance in developing their own reporting format.

## A.1 Source Evaluation for Individuals Using STRs

LAB HEADER with address here<sup>3.1.1.b</sup>

August 10, 2016<sup>3.1.1e</sup>

#### GENERIC GENETICS REPORT<sup>3.1.1.a</sup>

Agency: Submitting Agency<sup>3.2.1b</sup> Street Address City, State, ZIPCODE Lab Case #:10-000999Examiner:W. E. KiyoteAgency Case #:INV 2016123456Agency Case #:Investigator:Investigator:TroutSuspects:Leghorn, FCase Title:Unlawful Take of salmon

#### **EVIDENCE RECEIVED:**

The following evidence was received in the Evidence Unit<sup>3,2,1d</sup> of the Laboratory on March 17, 2010<sup>3,2,1c</sup>, and was transferred to the undersigned examiner on March 18, 2010<sup>3,2,1,f</sup>

LAB-1: "One of 2 salmon fillets from freezer search, Leghorn residence, 11-15-09." [Item# 1, ST######]<sup>3.2.1.e</sup>

LAB-2: "One of 2 salmon fillets from freezer search, Leghorn residence, 11-15-09." [Item# 2, ST######]

LAB-3: "Bloody hook-removal tool from campsite." [Item# 3, ST######]

#### EXAMINATION REQUESTED:<sup>3.3.1</sup>

The submitting investigator, Special Agent Bull Trout, requested analyses to determine the species origin of the evidence represented by LAB-1, LAB-2 and LAB-3, and whether the evidence represented by LAB-3 originated from the same individual as LAB-1 or LAB-2.<sup>3,3,1; 3,4,1</sup> Material was swabbed from LAB-3 and assigned item LAB-3A.

#### **EXAMINATION CONDUCTED:**

#### Mitochondrial DNA Analysis<sup>3.5.1; 3.5.3</sup>

A segment of the mitochondrial DNA (mtDNA) containing a portion of the cytochrome *b* gene<sup>3.5.6.</sup> from LAB-1, LAB-2, and LAB-3A was amplified by PCR and subjected to DNA sequence analysis. The resulting sequences were compared to An Agency reference database and reference sequences from the GenBank database for Rainbow trout (*Oncorhynchus mykiss*), Cutthroat trout (*Oncorhynchus clarkii*), Chinook salmon (*Oncorhynchus tschawytscha*), and Coho salmon (*Oncorhynchus kisutch*).<sup>3.5.5</sup>

Examiner's Initials

1 of 2<sup>3.1.1.d</sup>

Examination Report 10-000999 - Continued

August 10, 2016

#### Nuclear DNA Analysis<sup>3.5.1, 3.5.3</sup>

The DNA isolated from LAB-1, LAB-2, and LAB-3A was also characterized by STR analysis at ten nuclear loci designated as Loc01, Loc02, Loc03, Loc04, Loc05, Loc06, Loc07, Loc08, Loc09, and Loc10 to determine if the evidence items were of wild or hatchery origin. The resulting genotypes were compared to An Agency database of reference genotypes for salmon from rivers in southern Oregon.<sup>3.5.5</sup>

### EXAMINATION RESULTS:<sup>3.6.1</sup>

Mitochondrial DNA Analysis

The cytochrome *b* sequence obtained from LAB-1 and LAB-3A was identical to that of *Oncorhynchus tschawytscha*<sup>3.6.3</sup> reference sequence DNA A12345 at 430/430 base pairs (bp).<sup>3.6.3.a; b; c</sup>

The cytochrome *b* sequence obtained from LAB-2 was identical to that of *Oncorhynchus mykiss*<sup>3.6.3</sup> reference sequence DNA B40125 at 427/427 bp.<sup>3.6.4c</sup>

Nuclear DNA Analysis

The STR genotypes of LAB-1 and LAB-3A were the same<sup>3,6.6a</sup> at all ten loci.

The probability that LAB-1 and the material from LAB-3A originated from two different Chinook salmon from the Rogue River given that they share the same genotype is 1 in 120 billion.<sup>3.6.6b</sup>

The STR genotype of LAB-2 is not the same as that of LAB-1 and LAB-3A.

### **EXAMINATION CONCLUSIONS:**

There is strong support that LAB-1 and the material in LAB-3A originated from the same Chinook salmon, *Oncorhyncus tschawyscha*. <sup>3,7,1;3,7,2</sup>

LAB-2 originated from Oncorhynchus mykiss.<sup>3.7.1; 3.7.2</sup>

### **DISPOSITION OF EVIDENCE:**

All evidence items were transferred to the Evidence Unit pending return to the submitting agency. A complete record of the analysis may be obtained from the Laboratory.<sup>3.5.2</sup>

Validated Digital Signature here<sup>3.1.1f</sup>

W.E. Kiyote, Ph.D. Senior Forensic Analyst

<mark>2 of 2</mark>3.1.1d

# A.2 Species Identification Using Morphology

## FORENSIC MORPHOLOGY ASSOCIATES 101 FIRST AVENUE BIRDVILLE, CA 95555<sup>31.1b</sup>

August 10, 2016<sup>3.1.1e</sup>

### **MORPHOLOGY EXAMINATION REPORT** 3.1.1a

Agency:<sup>3.2.1b</sup> LE, Anytown 101 Main Street Anytown, OR 97500 Lab Case #:16-0111Examiner:SmithAgency Case #:2015123456Investigator:GreenSuspects:John DoeCase Title:Ex-birds

#### **EVIDENCE RECEIVED:**

The following evidence was received via FedEx by Evidence Technician Dusty Rhodes<sup>3.2.1d</sup> of the Laboratory on August 02, 2016<sup>3.2.1c</sup> and was transferred to the undersigned examiner on August 05, 2016:<sup>3.2.1f</sup>

LAB-1:	One of "Three (3) individual bags of feathers" [ST#xxxxx;Item#1]
LAB-2:	One of "Three (3) individual bags of feathers" [ST#xxxxxx;Item#1]
LAB-3:	One of "Three (3) individual bags of feathers" [ST#xxxxx;Item#1]
LAB-4:	One of "Two (2) birds" [ST#xxxxxx;Item#2]
LAB-5:	One of "Two (2) birds" [ST#xxxxxx;Item#2] <sup>3.2.1e</sup>

## EXAMINATION REQUESTED:<sup>3.3.1</sup>

The submitting investigator, Special Agent Rhett Green, requested analysis to determine the species origin of the evidence and the minimum number of individuals represented.

#### **EXAMINATION CONDUCTED:** 3.5.1

The evidence was examined visually, and identification was made by macroscopic comparison with known reference specimens in the collection of the Forensic Morphology Associates Laboratory. In all cases, similar species were considered and excluded, based on the external morphological characters exhibited by the evidence. <sup>3.5.3</sup> Complete documentation of the analyses conducted and data collected is maintained in the case record, which is available from the laboratory. <sup>3.5.2</sup>



#### EXAMINATION RESULTS:<sup>3.6.1</sup>

- LAB-1: The evidence consisted of tail feathers exhibiting diagnostic morphological characters of Northern Mockingbird, *Minus polyglottos*, <sup>3.6.3</sup> as verified by comparison with reference specimens in the collection of the Forensic Morphology Associates Laboratory. <sup>3.5.5</sup>
- LAB-2: The evidence consisted of tail feathers exhibiting diagnostic morphological characters of Northern Mockingbird, *Mimus polyglottos*, as verified by comparison with reference specimens in the collection of the Forensic Morphology Associates Laboratory.
- LAB-3: The evidence consisted of tail feathers exhibiting diagnostic morphological characters of Red-tailed Hawk, *Buteo jamaicensis*, as verified by comparison with reference specimens in the collection of the Forensic Morphology Associates Laboratory.
- LAB-4: The evidence consisted of a carcass exhibiting diagnostic morphological characters of Hairy Woodpecker, *Picoides villosus*, as verified by comparison with reference specimens in the collection of the Forensic Morphology Associates Laboratory.
- LAB-5: The evidence consisted of a carcass exhibiting diagnostic morphological characters of Killdeer, *Charadrius vociferus*, as verified by comparison with reference specimens in the collection of the Forensic Morphology Associates Laboratory.

### **EXAMINATION CONCLUSIONS:**<sup>3.7.1</sup>

- LAB-1: Tail feathers of NORTHERN MOCKINGBIRD (*Mimus polyglottos*<sup>3.7.2</sup>)
- LAB-2: Tail feathers of NORTHERN MOCKINGBIRD (*Mimus polyglottos*)
- LAB-3: Tail feathers of RED-TAILED HAWK (*Buteo jamaicensis*)
- LAB-4: Carcass of HAIRY WOODPECKER (*Picoides villosus*)
- LAB-5: Carcass of KILLDEER (*Charadrius vociferus*)

### SUMMARY OF MINIMUM NUMBER OF INDIVIDUALS 37.1

The evidence consisted of a minimum of two Northern Mockingbirds (*Mimus polyglottos*), based on the presence of twenty tail feathers. All other species in this evidence were represented by a minimum of one individual each.

### **DISPOSITION OF EVIDENCE:**

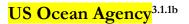
All evidence item(s) were transferred to the Evidence Unit pending return to the submitting agency.

# John J. Smith 3.1.1f

John J. Smith Senior Forensic Scientist

#### $2 \text{ of } 2^{3.1.1d}$

# A.3 Species Identification Using Sequencing



Agency Address<sup>3.1.1b</sup>

# **Genetics Examination Report**<sup>3.1.1a</sup>

<mark>SA Lotsa Fish</mark><sup>3.2.1b</sup>

**LAB CASE #:** 12335<sup>3.1.1c</sup>

**AGENCY:** 

<mark>US Ocean</mark> Agency<sup>3.2.1b</sup>

**AGENCY CASE #: 6789**<sup>3.2.1A</sup>

### Administrative Information

Evidence received	
Date:	DD/MM/YYYY <sup>3.2.1c, 3.2.1F</sup>
From:	SA Lotsa Fish <sup>3.2.1B</sup>
	US Ocean Agency <sup>3.2.1B</sup>
	Street Address
	City, State Zip Code
Via:	UPS Next Day Air (tracking #123 45 6789)
By:	Receiver's name <sup>3.2.1d</sup>
<b>Brief Description:</b>	One sealed package of frozen suspected whale meat.

Analysis Performed by: Analyst name

**Case summary and examination requested:** SA Lotsa Fish requested "DNA analysis to confirm that the imported meat, identified by passenger as whale meat (species unknown) is a marine mammal product."<sup>3.3.1, 3.4.1, 3.2.1e</sup>

**Disposition of Evidence:** Evidence will be held at the laboratory pending further instruction from SA Fish.

### Conclusions

I identified the submitted evidence item as originating from *Balaenoptera acutorostrata<sup>3.7.2</sup>*, Minke whale.<sup>3.7.1</sup>

**Conclusions Table.** Laboratory item number, sample information, and identification conclusion for the submitted evidence.

1	Laboratory Item #	Item #	Seized Prop. #	Laboratory Description	Identification
	Smp01	1	<mark>1234<sup>3.2.1</sup> e</mark>	suspected whale meat	Balaenoptera acutorostrata <sup>3.7.1, 3.7.2</sup>

MM/DD/YYYY<sup>3.1.1e</sup> | PAGE <u>X</u> OF <u>Y</u><sup>3.1.1d</sup>

# **Genetics Examination Report**

<b>INVESTIGATOR:</b>	SA Lotsa Fish	LAB CASE #:	12345
AGENCY:	US OCEAN AGENCY	AGENCY CASE #:	6789

#### **Details of Examination**

**Methods**<sup>3.5.1</sup>: A small tissue subsample was taken from the submitted item. Standard laboratory protocols were used for molecular genetic analysis<sup>3.5.3</sup>. DNA was extracted from the evidence. The mtDNA control region <sup>3.5.6</sup> was amplified from the extracted DNA and controls and sequenced. Resulting sequences were evaluated for quality, edited, aligned, and compared to appropriate reference sequences, following standard procedures.

The sequence data from the unknown sample was compared with reference sequences and identifications were made based on sequence similarity and phylogenetic reconstruction. A full record of the work is available from the laboratory.<sup>3.5.2</sup>

#### Results:<sup>3.6.1</sup>

Evidence sequences produced a 445<sup>3.6.3a</sup> bp contig, which shared 99.8% identity <sup>3.6.4c</sup> with an Ocean Agency Forensic Laboratory reference sequence for *Balaenoptera acutorostrata*.<sup>3.6.3</sup>

**Results Table.** Laboratory item and evidence bag numbers, most similar reference sequence and source species, and the number of base pairs (bp) in the evidence sequence identical to those in the most similar reference sequence.

Laborato ry Item #	Item #	Most similar species	Most similar reference sequence(s)	# Identical bp/ Total # bp (%)
Smp01		Balaenoptera acutorostrata <sup>3.6.4b</sup>	Bacu001 <sup>3.6.4b</sup>	<mark>444/445<sup>3.6.4c</sup> (99.8%)</mark>

**Reference Material:** Forensics Laboratory internal database "Cetacean standards dlp V1" and DNA Surveillance databases "All cetaceans Vs4.3" and "Mysticetes Vs4.3"<sup>3.5.5</sup>.

# Analyst name<sup>3.1.1f</sup>

**Analyst's signature** Certified Wildlife Forensic Scientist

### -END-

MM/DD/YYYY<sup>3.1.1E</sup> | PAGE <u>X</u> OF <u>Y</u><sup>3.1.1D</sup>

# Annex B

(informative)

# **Bibliography**

This is not meant to be an all-inclusive list as the group recognizes other publications on this subject may exist. At the time these standards were drafted, these were the publications available to the working group members for reference. Additionally, any mention of a particular software tool or vendor as part of this bibliography is purely incidental, and any inclusion does not imply endorsement by the authors of this document.

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