OSAC 2021-S-0029
Standard for Familial DNA Searching

Human Forensic Biology Subcommittee
Biology Scientific Area Committee (SAC)
Organization of Scientific Area Committees (OSAC) for Forensic Science

OSAC Proposed Standard

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Prepared by
Human Forensic Biology Subcommittee
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The STRP panel will consist of an independent and diverse panel, including subject matter experts, human factors scientists, quality assurance personnel, and legal experts, which will be tasked with evaluating the proposed standard based on a comprehensive list of science-based criteria.

For more information about this important process, please visit our website at: https://www.nist.gov/topics/organization-scientific-area-committees-forensic-science/scientific-technical-review-panels.

Version No.	Issue Date	Section	Reason
2.0	January 5, 2023	--	Added to the OSAC Registry and publicly announced.
2.1	December 28, 2023	All	Format and font updates only. No content changes made.

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Standard for Familial DNA Searching

1 Scope

This standard requires laboratories that perform familial DNA searches to have a written policy specifying criteria for:

a) accepting a familial DNA search request;

b) administrative structure and responsibilities;

c) the search process;

d) reporting results; and

e) safeguarding individual privacy and confidentiality of the results.

This standard also defines validation requirements.

NOTE This standard does not apply to Investigative Genetic Genealogy or to the investigation of partial matches that may occur during the normal course of forensic database searches.

2 Normative References

The document contains no normative references. See Annex A, Bibliography for other references.

3 Terms and Definitions

3.1

familial DNA search

familial DNA searching

A deliberate search of a DNA database using software to detect and statistically rank a list of potential candidates in the database who could be close biological relatives (e.g. parent, child, sibling) of the unknown individual contributing the evidence DNA profile.

3.2

likelihood ratio (LR)

The probability of the evidence under one proposition (hypothesis), divided by the probability of the evidence under an alternative, mutually exclusive proposition (hypothesis). The magnitude of its value expresses the weight of the evidence.

3.3

likelihood ratio threshold

The likelihood ratio below which a database profile specific to the relationship(s) under consideration would not be further investigated. For example, a laboratory may decide to investigate only those candidates above a certain likelihood ratio.

3.4

lineage testing

Genetic testing, such as YSTR, XSTR, or mitochondrial DNA analysis, used to trace maternal or paternal inheritance.

3.5

ranking threshold

The rank below which a database profile specific to the relationship(s) under consideration would not be further investigated. For example, a laboratory may decide to investigate only a set number of the best ranked candidates regardless of the likelihood ratio.

3.6

sensitivity study

An assessment of the proportion of familial searches that detect true relatives of the contributor of the evidence profile when one or more relatives is present in the database.

3.7

specificity study

An assessment of the proportion of familial searches that exclude non-relatives of the contributor of the evidence profile. Practical limitations can restrict this study to the initial statistical comparisons of simulated forensic unknowns to DNA database profiles.

4   Requirements

4.1   Policy and Procedure: The laboratory shall have documented policies and/or procedures for conducting familial DNA searching that shall contain the information detailed in 4.1.1 through 4.1.11, as applicable to the laboratory's involvement in the elements described in each substandard. To aid the public and law enforcement in understanding the laboratory's familial search results, the familial search policy should be publicly available.

4.1.1   The document shall define the process for case submission and acceptance for conducting DNA familial searching. The following should be considered in the case acceptance policy:
   a) seriousness or seriality of the crime;

   b) commitment to proceed with investigation and prosecution;

   c) case metadata and laboratory notes provided as available;

d) investigative stage to initiate a familial search, such as when other apparently viable investigative strategies have been exhausted;

e) the availability of additional DNA evidence; and

f) out of state search requests.

4.1.2 The document shall define the roles of the individuals involved in the DNA familial search process and oversight. The familial DNA search process may involve individuals in the following roles (an individual may fill more than one role):

a) a review committee (e.g. to evaluate requests, prioritize testing, verify results, and determine the propriety of moving forward with an investigation or disclosure of a name);

b) an individual or committee who has the ultimate control for the search and directs the release of investigative lead(s) and any follow-up;

c) an administrative representative from the source testing laboratory;

d) an administrative representative from the searching laboratory;

e) a technical representative from the source testing laboratory;

f) a technical representative from the searching laboratory;

g) a representative with access to investigative databases (metadata) typically restricted to law enforcement personnel;

h) a representative from the requesting law enforcement agency;

i) a representative from the prosecuting agency; and

j) a representative of the defendant when the search is being performed at the request of the defense.

4.1.3 The document shall define how information obtained through the familial search process is shared and released and should include:

a) Safeguards for the disclosure of sensitive information to the proper individuals; and

b) Provisions protecting the privacy of possible family members in the search database who could not be the perpetrator.

4.1.4 The document shall ensure that all the individuals with roles established in 4.1.2 are informed of the process, likely outcomes, limitations, and the need for additional investigation. This could be achieved through the use of a documented memorandum of understanding or equivalent.

4.1.5 the document shall address the collection, testing, and retention/destruction of the following comparison samples:

4.1.5.1 surreptitiously collected samples associated with the person of interest.

4.1.5.2 samples known not to be from the perpetrator but that could provide additional confirmatory information (e.g. possible relatives of the perpetrator), whether collected surreptitiously or voluntarily.

4.1.5.3 confirmatory samples, which are direct reference samples from a person of interest that are collected with a chain of custody.

4.1.6 The document shall define the acceptance requirements of the evidence profile to be searched to include the information in 4.1.6.1 through 4.1.6.5.

4.1.6.1 That the DNA sample was recovered directly from the crime scene (or associated to the crime scene) and is attributed to the putative perpetrator.

4.1.6.2 The minimum threshold for the number of searchable loci and/or profile rarity.

4.1.6.3 The acceptable level of potential allelic dropout in the profile submitted for the familial DNA search.

4.1.6.4 The number of possible contributors, e.g. single source, deduced single source from a mixture, or mixtures up to a defined maximum number of contributors.

4.1.6.5 A requirement that a direct comparison search of the profile has occurred in relevant databases (e.g., Combined DNA Index System (CODIS)) prior to the familial DNA search being conducted.

4.1.7 The document shall define which database categories will be searched (e.g. convicted offenders, arrestees, suspects).

4.1.8 The document shall define which relationships (e.g., 1st degree relatives only) will be considered.

4.1.9 When possible, the laboratory shall conduct expanded autosomal STR (beyond the original 13 CODIS core loci) and/or lineage testing (e.g. YSTR, XSTRs and/or mitochondrial DNA) of potential relatives. The laboratory shall document situations where and justify conditions when conducting additional testing is not required.

4.1.10 The document shall define the thresholds (e.g. likelihood ratio and/or ranking) for proceeding with the release of information regarding potential relatives.

4.1.11 The document shall define criteria for the release of the name(s) of the potential relatives(s). The following criteria may be considered prior to the release of the name(s) of the potential relative(s):
a) The use of metadata for the evaluation of a potential relative.

b) Assessing kinship using non-invasive investigation, i.e., one that does not require direct contact with a person of interest.

4.2   Technical

4.2.1   The laboratory shall use validated analytical procedures/systems and software.

4.2.2   Internal validation shall include sensitivity and specificity studies to include the range of search profiles defined in 4.1.6.2, 4.1.6.3, and 4.1.6.4.

It is impractical to perform both sensitivity and specificity studies on the entire process with a meaningfully sized dataset when the process includes extensive additional testing and investigative work.

A reasonable sensitivity test would seed a database of unrelated individuals with the profiles of true relatives (at a minimum, those defined in section 4.1.9) of the experimental search profile. Ranking the LRs for all comparisons from highest to lowest, sensitivity is defined as the proportion of true relatives (real or in silico) included at each rank. Alternatively, the ranked list of unrelated individuals' LRs could be seeded with the LRs for each of the true family members, thus avoiding the insertion of experimental profiles into the database.

A reasonable specificity test would examine how many individuals remain as candidates after the statistical process (e.g., the initial LR rankings based solely upon autosomal STR loci). Coupled with how many of those individuals would meet the requirements to be tested with additional lab work, this will give an estimate of how likely it would be to see a false positive. Knowing whether the statistical process the lab has put in place will lead to tens of candidates, hundreds of candidates, or thousands of candidates will give a sense of whether or not the subsequent lineage testing (or additional autosomal markers) might adventitiously include a non-relative.

Search criteria developed from sensitivity and specificity studies should be established to err on the side of minimizing false positives. This approach will enhance the probability of releasing reliable leads.

Search criteria developed from sensitivity and specificity studies can be impacted by the capacity and resources of the laboratory in addition to the size of the database being searched.

4.2.3   The laboratory shall determine if the process will incorporate the following:

4.2.3.1 population substructure (Fst or theta)

4.2.3.2 mutation model

4.2.3.3 silent (null) allele model

4.2.3.4 correction of linkage

4.2.3.5 probabilistic genotyping

4.2.3.6 prior odds

4.2.4　The laboratory shall have a documented training program specific to the familial DNA searching including training in kinship analysis.

Training of an individual evaluating familial search results should include the following concepts.

    a) The expected levels of DNA sharing for various relationships, including:
       i. The flow of DNA through pedigrees; and
       ii. lineage vs. autosomal patterns of inheritance

    b) The calculations performed in familial searching, including:
       i. likelihood ratios; and
       ii. conditional probabilities given an assumed level of relatedness

    c) Likelihood ratio distributions for various relationship categories.

    d) Evaluating pedigrees larger than two-person comparisons.

    e) Addressing the issues of linked loci and meiotic mutations.

    f) The effect of population database selection on the likelihood ratio.

    g) The effect of database size on the ranking of true relatives.

    h) The benefits and limitations of lineage vs. autosomal loci.

Annex A
(informative)
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