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**Standard for Processing Evidence for the Detection of  
Friction Ridge Impressions**

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# Standard for Processing Evidence for the Detection of Friction Ridge Impressions

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## Foreword

This document has been developed to improve the quality and consistency of friction ridge examination practices.

This document specifies (or establishes) broad class processing techniques for the detection of friction ridge impressions. The specific processing techniques applied are determined by the FSP based on the specific processes that are appropriate for each particular substrate and matrix combination.

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This document was revised, prepared, and finalized as a standard by the Friction Ridge Consensus Body of the AAFS Standards Board. The draft of this standard was developed by the Friction Ridge Subcommittee of the Organization of Scientific Area Committees (OSAC) for Forensic Science.

Questions, comments, and suggestions for the improvement of this document can be sent to ASB Secretariat, [asb@aafs.org](mailto:asb@aafs.org) or 410 N 21<sup>st</sup> Street, Colorado Springs, CO 80904.

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# Standard for Processing Evidence for the Detection of Friction Ridge Impressions

## 1 Scope

This document provides requirements for the processing of evidence, within a laboratory setting, for the detection of friction ridge impressions. The standard specifies the broad class of processing techniques and sequences to be applied when processing such evidence. This document does not address the processing of evidence at a crime scene, the photography or digital processing of friction ridge impressions or the validation of the various processing techniques, necessary equipment, or storage requirements.

## 2 Normative References

There are no normative reference documents. Annex A, Bibliography, contains informative references.

## 3 Terms and Definitions

For purposes of this document, the following terms and definitions apply.

### 3.1

#### **forensic light source**

A light source that may be fixed, filtered or tunable to a variety of spectral ranges.

### 3.2

#### **Forensic Service Provider**

#### **FSP**

Organization or individual that conducts and/or supplies forensic services.

### 3.3

#### **matrix**

Transfer medium that is deposited or removed by the friction ridge skin when making an impression (e.g., grease/oil, sweat, blood).

### 3.4

#### **semi-porous**

Partially but not freely or wholly permeable (e.g., glossy paper).

### 3.5

#### **sequential processing**

The application of chemical and/or physical friction ridge development techniques in a specific order to target specific constituents of friction ridge impressions which may be visualized for examination, and to maximize the preservation of the friction ridge detail during each process.

34 **3.6**35 **substrate**

36 Surface or material upon which an item of interest is deposited (e.g., porous, non-porous).

37 *ISO 21043-1*<sup>1</sup>38 **4 Processing Considerations**39 **4.1** The processes applied by each FSP shall be based on the efficacy and limitations of the  
40 process, availability of resources and processing techniques, and the type and condition of the  
41 evidence.42 **4.2** The FSP shall apply processing techniques in a sequence (i.e., sequential processing) from  
43 least destructive to most destructive for the detection of friction ridge impressions.44 **4.2.1** The FSP shall document deviations from the processing sequences. The FSP may  
45 supplement and/or deviate from the sequences for the detection of friction ridge impressions in  
46 certain situations. Some examples of when the FSP may supplement and/or deviate from the  
47 sequences are as follows.48 a) The item does not react to a processing technique as expected or reacts adversely to a chemical  
49 (e.g., thermal paper).

50 b) The item of evidence has an obvious known contaminant such as blood or grease.

51 c) The processing technique has not been validated to perform sufficiently in certain  
52 environmental conditions.53 d) The size of the item does not allow for a specific processing technique that aligns to the  
54 required sequence.55 **4.3** Prior to applying specific processing techniques to evidence, the FSP shall assess the potential  
56 for negative implications to other forensic discipline examinations, communicate those concerns to  
57 the stakeholder, and document the communication. (e.g., DNA, trace evidence, questioned  
58 documents, firearms, drug chemistry, etc.).59 **4.3.1** The FSP shall have a written policy about communication with the stakeholder regarding  
60 negative implications to future examinations. If FSP personnel communicate with a stakeholder, it  
61 shall be documented. Some potential negative implications to consider are as follows.62 a) Forensic light source(s), such as short-wave ultraviolet (UV) light source, and the potential  
63 negative impact on DNA examinations.64 b) Cyanoacrylate dye stains and the potential negative impact on adhesive side processing,  
65 questioned documents, drug chemistry, and trace evidence examinations.66 c) Porous chemical processing and the potential negative impact on thermal paper and  
67 Questioned Documents examinations.

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68 d) Powder and the potential negative impact on electronic evidence examinations.

69 **4.4** The FSP shall preserve potentially suitable friction ridge impressions prior to applying the  
70 next processing techniques within the processing sequence. The FSP shall have a policy for the  
71 appropriate methods for digital imaging (e.g., scanning and photography) of the friction ridge  
72 impressions.

73 **4.5** The FSP shall establish appropriate health and safety practices, along with universal  
74 precautions to ensure the safety of personnel while maintaining the integrity of the evidence.

## 75 **5 Processing Sequences**

### 76 **5.1 General**

77 **5.1.1** The processing sequences below are based on the appropriate techniques that are specific  
78 for a particular substrate and matrix combination and shall be used for laboratory-based  
79 processing.

80 **5.1.2** Many items of evidence consist of more than one physical property (e.g., a porous envelope  
81 with a glassine window). In these situations, the FSP shall apply the processing techniques using  
82 sequences appropriate for the relevant areas in a manner that does not negatively impact other  
83 areas of the evidence.

84 **5.1.3** Wet items should be allowed to dry prior to processing. Once dried, processing of the items  
85 should proceed under one of the sequences listed below.

86 NOTE 1 The processing sequences below are meant to describe the most universal chemical processing  
87 sequences for routinely encountered substrates. It is not meant to be an exhaustive list of all available  
88 techniques.

89 NOTE 2 Guidance related to application, formulation, and optimization of specific processing techniques can  
90 be found in the publications listed in Annex A.

91 **5.1.4** Every processing sequence shall start with a visual examination and then also be completed  
92 following every processing technique.

### 93 **5.2 Non-porous**

94 The following is the minimum processing sequence an FSP shall perform for non-porous items.

95 a) Visual.

96 b) Forensic light source(s).

97 c) Cyanoacrylate fuming.

98 d) Contrast, such as dye stain, forensic light source(s), and/or powder.

### 99 5.3 Porous

100 The following is the minimum processing sequence an FSP shall perform for porous items.

- 101 a) Visual.
- 102 b) Forensic light source(s).
- 103 c) Amino acid reagent(s) (e.g., 1,2 Indanedione, 1,8 Diazafuoren 9-one, Ninhydrin)

104 If a fluorescent reagent is used, it shall be used prior to Ninhydrin.

105 NOTE If the condition of the evidence supports the use of a sebaceous reagent, Oil Red O and/or physical  
106 developer may be used.

### 107 5.4 Semi-porous

108 The following is the minimum processing sequence an FSP shall perform for semi-porous items.

- 109 a) Visual.
- 110 b) Forensic light source(s).
- 111 c) Cyanoacrylate fuming.

112 d) Magnetic Powder.

113 If a fluorescent amino acid reagent is not going to be used, regular powder would be an  
114 acceptable alternative to magnetic powder.

115 e) Amino acid reagent(s) (e.g., 1,2 Indanedione, 1,8 Diazafuoren 9-one, Ninhydrin).

116 If a fluorescent reagent is used, it shall be used prior to Ninhydrin

117 f) Contrast, such as dye stain, forensic light source(s), and/or powder.

### 118 5.5 Adhesive

119 The following is the minimum processing sequence an FSP shall perform for adhesive surfaces.

- 120 a) Visual.
- 121 b) Forensic light source(s).
- 122 c) Adhesive side powder suspension/Gentian Violet/fluorescent reagent.

### 123 5.6 Blood

124 **5.6.1** The bloody surface/item shall be dried prior to processing.

125 **5.6.2** Depending on the blood process used, a blood fixative may be needed prior to processing.

126 NOTE Examination by other forensic disciplines, such as DNA or trace evidence, may need to be preserved or  
127 completed prior to blood processing.

128 **5.6.3** The following is the minimum processing sequence an FSP shall perform for bloody  
129 surfaces/items.

- 130 a) Visual.
- 131 b) Forensic light source(s).
- 132 c) Protein reagent/Heme reagent/Acid reagent.
- 133 d) Forensic light source(s) (if applicable).

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## Annex A (informative)

### Bibliography

The following bibliography is not intended to be an all-inclusive list, review, or endorsement of literature on this topic. The goal of the bibliography is to provide publications cited informationally, and publications relevant to the standard.

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