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Standard Guide for Sampling Seized Drugs for Qualitative and Quantitative Analysis¹

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1. Scope

1.1 This guide covers minimum considerations for sampling of seized drugs for qualitative and quantitative analysis.

2. Referenced Documents

2.1 ASTM Standards:²

E105 Practice for Probability Sampling of Materials

E122 Practice for Calculating Sample Size to Estimate, With Specified Precision, the Average for a Characteristic of a Lot or Process

E141 Practice for Acceptance of Evidence Based on the Results of Probability Sampling

E1732 Terminology Relating to Forensic Science

E2329 Practice for Identification of Seized Drugs

E2334 Practice for Setting an Upper Confidence Bound For a Fraction or Number of Non-Conforming items, or a Rate of Occurrence for Non-conformities, Using Attribute Data, When There is a Zero Response in the Sample

2.2 ISO Standards:³

ISO 3534-1 Statistics – Vocabulary and symbols ~~Symbols~~ – Part 1: Probability and ~~general statistical terms~~ General Statistical Terms

ISO 3534-2 Statistics – Vocabulary and symbols ~~Symbols~~ – Part 2: Statistical ~~quality control~~ Quality Control

3. Significance and Use

3.1 This guide provides information for the sampling of seized-drug submissions.

3.2 The principal purpose of sampling in the context of this guide is to answer relevant questions about a population by examination of a portion of the population. For example:

What is the net weight of the population?

What portion of the units of a population can be said to contain a given drug at a given level of confidence?

3.3 By developing a sampling strategy and implementing appropriate sampling schemes, as illustrated in Fig. 1, a laboratory will minimize the total number of required analytical determinations, while ensuring that all relevant legal and scientific requirements are met.

4. Sampling Strategy

4.1 A sampling strategy is highly dependent on the purpose of the investigation, the original question, and the ultimate use of the results. Laws and legal practices form the foundation of most strategies and shall be taken into account when designing a sampling scheme. Therefore, specific sampling strategies are not defined in this guide.

4.2 The laboratory has the responsibility to develop its own strategies consistent with these recommendations. It is recommended that the following key points be addressed:

4.2.1 Sampling may be statistical or non-statistical.

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² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For Annual Book of ASTM Standards volume information, refer to the standard's Document Summary page on the ASTM website.

³ Available from International Organization for Standardization (ISO), 1 rue de Varembe, Case postale 56, CH-1211, Geneva 20, Switzerland, http://www.iso.ch.

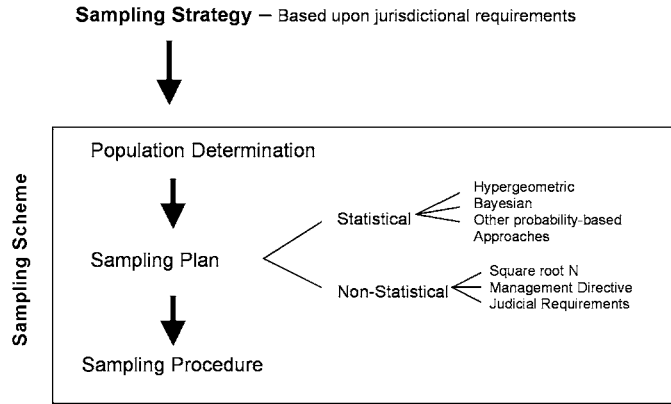


FIG. 1 Relationship of Various Levels Required in Sampling

NOTE 1—For the purpose of this guide, the use of the term *statistical* is meant to include the notion of an approach that is probability-based.

4.2.1.1 In many cases, a non-statistical approach may suffice. The sampling plan shall provide an adequate basis for answering questions of applicable law. For example,

Is there a drug present in the population?

Are statutory enhancement levels satisfied by the analysis of a specified number of units?

4.2.1.2 If an inference about the whole population is to be drawn from a sample, then the plan shall be statistically based and limits of the inference shall be documented.

4.2.2 Statistically selected units shall be analyzed to meet Practice E2329 if statistical inferences are to be made about the whole population.

5. Sampling Scheme

5.1 The sampling scheme is an overall approach that includes population determination, selection of the sampling plan and procedure and, when appropriate, sample reduction prior to analysis (Fig. 2).