

# Archived Document

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December 2000

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## AUTO01-A

### Laboratory Automation: Specimen Container/ Specimen Carrier; Approved Standard

This document provides standards for the design and manufacture of specimen containers and carriers used for collecting and processing liquid samples, such as blood and urine, for clinical testing in laboratory automation systems.

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A standard for global application developed through the Clinical and Laboratory Standards Institute consensus process.

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Clinical and Laboratory Standards Institute  
950 West Valley Road, Suite 2500  
Wayne, PA 19087 USA  
P: +1.610.688.0100  
F: +1.610.688.0700  
[www.clsi.org](http://www.clsi.org)  
[standard@clsi.org](mailto:standard@clsi.org)

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## Laboratory Automation: Specimen Container/Specimen Carrier; Approved Standard

Volume 20 Number 29

Paul J. Orsulak, Ph.D., Chairholder  
Keiichi Naka, Ph.D., Vice-Chairholder  
Nelson S. Everhart, III  
Mary Haven  
L. Omar Henderson, Ph.D.  
Dennis A. Lamb  
Stewart Marsden

### Abstract

*Laboratory Automation: Specimen Container/Specimen Carrier; Approved Standard* (CLSI document AUTO01-A) was developed for those engaged in the design and manufacture of specimen collection containers used for specimen handling in the healthcare and clinical laboratory environments, and for those engaged in the design and manufacture of clinical laboratory instrumentation and clinical laboratory automation systems. This document is intended to *lead* design and manufacturing toward standardized products for a wider variety of instruments and automated laboratory systems.

CLSI. *Laboratory Automation: Specimen Container/Specimen Carrier; Approved Standard*. CLSI document AUTO01-A. Clinical and Laboratory Standards Institute, 950 West Valley Road, Suite 2500, Wayne, Pennsylvania 19087 USA, 2000.

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## Matrix of NCCLS Laboratory Automation Standards

The laboratory automation standards documents, AUTO01, AUTO2, AUTO3, AUTO4, and AUTO5 are interdependent with respect to their implementation in automated laboratory systems. The matrix describes the engineering relationships between the standards elements in each of the five documents. This matrix is provided so that designers and engineers, as well as users and customers, understand the relationships between the different standards' components in an automated system. The matrix format allows the users of one document to easily identify other standard elements, which relate to the standard elements in the document or documents from which they may be working, to design a system correctly.

**How to Read the Matrix** (See matrix on the next page.)

The numbers listed on the horizontal (X) and vertical (Y) axes contain multiple-digit numbers (e.g., (1)5.4, (5)5.4.1.3).

The 'first digit' (in parentheses) represents one of the five automation documents (e.g., (1)5.4 is from AUTO01; (5)5.4.1.3 is from AUTO5).

The 'remaining digits' represent the specific section of that document.

The symbol XX represents the direct 'engineering relationship' between two sections.

The symbol ## represents the section's 'self'; when it has been lined up with itself on the other axis.

SAMPLE

**Related NCCLS Publications\***

- AUTO2**     **Laboratory Automation: Bar Codes for Specimen Container Identification.** This document provides specifications for use of linear bar codes on specimen tubes in the clinical laboratory and for use on laboratory automation systems.
- AUTO3**     **Laboratory Automation: Communications with Automated Clinical Laboratory Systems, Instruments, Devices, and Information Systems.** The goal of this document is to facilitate accurate and timely electronic exchange of data and information between the automated laboratory elements.
- AUTO4**     **Laboratory Automation: Systems Operational Requirements, Characteristics, and Information Elements.** This document provides standards of interest to operators for display of system status information such as specimen location, reagent supply, and warnings and alerts to support laboratory automation operations.
- AUTO5**     **Laboratory Automation: Electromechanical Interfaces.** This document provides guidance for the standardization of electromechanical interfaces between instruments and/or specimen processing and handling devices and automation systems in the automated laboratory.
- GP2-A2-C**   **NCCLS Procedure Manual Template.** This computer template enables laboratorians to prepare consistent technical procedures in the NCCLS format. The template and its user manual, used along with the GP2-A3 guideline, provide a procedure format that is as easy to use as a word processing program. Procedures can be stored as individual files for easy retrieval and updating, or they can be networked through the local computer system for electronic distribution throughout the laboratory. The template format consists of tables for recording essential information for all procedures and an outline of key headings for incorporating procedure-specific details.
- GP2-A3**     **Clinical Laboratory Technical Procedure Manuals – Third Edition; Approved Guideline (1996).** This document provides guidance for the patient-testing community by addressing the design, preparation, maintenance, and use of paper or electronic technical procedure manuals.
- GP18-A**     **Laboratory Design; Approved Guideline (1998).** This guideline provides a foundation of information about laboratory design elements that can be used to help define the issues being considered when designing a laboratory.
- GP19-A**     **Laboratory Instruments and Data Management Systems: Design of Software User Interfaces and End-User Software Systems Validation, Operation, and Monitoring; Approved Guideline (1995).** The document identifies important factors that designers and laboratory managers should consider when developing new software-driven systems and selecting software user interfaces. Also included are simple rules to help prepare validation protocols for assessing the functionality and dependability of software.

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\* Proposed- and tentative-level documents are being advanced through the NCCLS consensus process; therefore, readers should refer to the most recent editions.

**Related NCCLS Publications (Continued)**

- H1-A4**      **Evacuated Tubes and Additives for Blood Specimen Collection - Fourth Edition; Approved Standard (1996).** *American National Standard.* This standard discusses requirements for blood collection tubes and additives, including heparin, EDTA, and sodium citrate.
- H18-A2**     **Procedures for the Handling and Processing of Blood Specimens; Approved Guideline—Second Edition (1999).** This guideline addresses multiple factors associated with handling and processing of specimens, and factors that can introduce imprecision or systematic bias into results.
- H38-P**      **Calibration and Quality Control of Automated Hematology Analyzers; Proposed Standard (1999).** This document addresses calibration and quality control strategies for multichannel hematology analyzers; assignment of values to calibrator materials; calibration using stabilized blood controls; internal quality control; pair difference analysis; and use of the weighted moving average ( $\bar{x}_B$ ) method.
- M29-A**      **Protection of Laboratory Workers from Instrument Biohazards and Infectious Disease Transmitted by Blood, Body Fluids, and Tissue; Approved Guideline (1997).** A consolidation of M29-T2 and I17-P, this document provides guidance on the risk of transmission of hepatitis viruses and human immunodeficiency viruses in any laboratory setting; specific precautions for preventing the laboratory transmission of blood-borne infection from laboratory instruments and materials; and recommendations for the management of blood-borne exposure.



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