Specification for Low-Level Protocol to Transfer Messages Between Clinical Laboratory Instruments and Computer Systems; Approved Standard—Second Edition

This document describes the electronic transmission of digital information between clinical laboratory instruments and computer systems.

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



Clinical and Laboratory Standards Institute

Advancing Quality in Health Care Testing

Clinical and Laboratory Standards Institute (CLSI, formerly NCCLS) is an international, interdisciplinary, nonprofit, standards-developing, and educational organization that promotes the development and use of voluntary consensus standards and guidelines within the health care community. It is recognized worldwide for the application of its unique consensus process in the development of standards and guidelines for patient testing and related health care issues. Our process is based on the principle that consensus is an effective and cost-effective way to improve patient testing and health care services.

In addition to developing and promoting the use of voluntary consensus standards and guidelines, we provide an open and unbiased forum to address critical issues affecting the quality of patient testing and health care.

PUBLICATIONS

A document is published as a standard, guideline, or committee report.

Standard A document developed through the consensus process that clearly identifies specific, essential requirements for materials, methods, or practices for use in an unmodified form. A standard may, in addition, contain discretionary elements, which are clearly identified.

Guideline A document developed through the consensus process describing criteria for a general operating practice, procedure, or material for voluntary use. A guideline may be used as written or modified by the user to fit specific needs.

Report A document that has not been subjected to consensus review and is released by the Board of Directors.

CONSENSUS PROCESS

The CLSI voluntary consensus process is a protocol establishing formal criteria for:

- the authorization of a project
- the development and open review of documents
- the revision of documents in response to comments by users
- the acceptance of a document as a consensus standard or guideline.

Most documents are subject to two levels of consensus— "proposed" and "approved." Depending on the need for field evaluation or data collection, documents may also be made available for review at an intermediate consensus level.

Proposed A consensus document undergoes the first stage of review by the health care community as a proposed standard or guideline. The document should receive a wide and thorough technical review, including an overall review of its scope, approach, and utility, and a line-by-line review of its technical and editorial content.

Approved An approved standard or guideline has achieved consensus within the health care community. It should be reviewed to assess the utility of the final document, to ensure attainment of consensus (ie, that comments on earlier versions have been satisfactorily addressed), and to identify the need for additional consensus documents.

Our standards and guidelines represent a consensus opinion on good practices and reflect the substantial agreement by materially affected, competent, and interested parties obtained by following CLSI's established consensus procedures. Provisions in CLSI standards and guidelines may be more or less stringent than applicable regulations. Consequently, conformance to this voluntary consensus document does not relieve the user of responsibility for compliance with applicable regulations.

COMMENTS

The comments of users are essential to the consensus process. Anyone may submit a comment, and all comments are addressed, according to the consensus process, by the committee that wrote the document. All comments, including those that result in a change to the document when published at the next consensus level and those that do not result in a change, are responded to by the committee in an appendix to the document. Readers are strongly encouraged to comment in any form and at any time on any document. Address comments to Clinical and Laboratory Standards Institute, 940 West Valley Road, Suite 1400, Wayne, PA 19087, USA.

VOLUNTEER PARTICIPATION

Health care professionals in all specialties are urged to volunteer for participation in CLSI projects. Please contact us at customerservice@clsi.org or +610.688.0100 for additional information on committee participation.

LIS01-A2 ISBN 1-56238-665-4 ISSN 0273-3099

Volume 28 Number 13ISSN 0273-3Specification for Low-Level Protocol to Transfer Messages Between
Clinical Laboratory Instruments and Computer Systems; Approved
Standard—Second Edition

David Chou, MD Andrzej J. Knafel, PhD Charles D. Hawker, PhD, MBA, FACB Ed Heierman, PhD David A. Lacher, MD, MEd William Neeley, MD, FACP Eugene T. Reilly Richard S. Seaberg, MT(ASCP)

Abstract

CLSI document LIS01-A2—Specification for Low-Level Protocol to Transfer Messages Between Clinical Laboratory Instruments and Computer Systems; Approved Standard—Second Edition describes the electronic transmission of digital information between clinical laboratory instruments (those that measure one or more parameters from one or multiple samples) and computer systems (those that are configured to accept instrument results for further processing, storage, reporting, or manipulation).

Clinical and Laboratory Standards Institute (CLSI). Specification for Low-Level Protocol to Transfer Messages Between Clinical Laboratory Instruments and Computer Systems; Approved Standard—Second Edition. CLSI document LIS01-A2 (ISBN 1-56238-665-4). Clinical and Laboratory Standards Institute, 940 West Valley Road, Suite 1400, Wayne, Pennsylvania 19087-1898 USA, 2008.

The Clinical and Laboratory Standards Institute consensus process, which is the mechanism for moving a document through two or more levels of review by the health care community, is an ongoing process. Users should expect revised editions of any given document. Because rapid changes in technology may affect the procedures, methods, and protocols in a standard or guideline, users should replace outdated editions with the current editions of CLSI/NCCLS documents. Current editions are listed in the CLSI catalog and posted on our website at www.clsi.org. If your organization is not a member and would like to become one, and to request a copy of the catalog, contact us at: Telephone: 610.688.0100; Fax: 610.688.0700; E-Mail: customerservice@clsi.org; Website: www.clsi.org



Number 13

LIS01-A2

Copyright [©]2008 Clinical and Laboratory Standards Institute. Except as stated below, neither this publication nor any portion thereof may be adapted, copied, or otherwise reproduced, by any means (electronic, mechanical, photocopying, recording, or otherwise) without prior written permission from Clinical and Laboratory Standards Institute ("CLSI").

CLSI hereby grants permission to each individual member or purchaser to make a single reproduction of this publication for use in its laboratory procedure manual at a single site. To request permission to use this publication in any other manner, contact the Executive Vice President, Clinical and Laboratory Standards Institute, 940 West Valley Road, Suite 1400, Wayne, Pennsylvania 19087-1898, USA.

Suggested Citation

(CLSI. Specification for Low-Level Protocol to Transfer Messages Between Clinical Laboratory Instruments and Computer Systems; Approved Standard—Second Edition. CLSI document LIS01-A2. Wayne, PA: Clinical and Laboratory Standards Institute; 2008.)

Approved Standard April 2003

Approved Standard—Second Edition April 2008

ISBN 1-56238-665-4 ISSN 0273-3099

Volume 28

Committee Membership

Area Committee on Automation and Informatics

David Chou, MD Chairholder University of Washington Medical Center Seattle, Washington

Andrzej J. Knafel, PhD Vice-Chairholder Roche Instrument Center AG Rotkreuz, Switzerland

Charles D. Hawker, PhD, MBA, FACB ARUP Laboratories, Inc. Salt Lake City. Utah

Ed Heierman, PhD Abbott Diagnostics Irving, Texas

David A. Lacher, MD, MEd National Center for Health Statistics Hyattsville, Maryland

William Neeley, MD, FACP, DABCC Detroit Medical Center University Laboratories Detroit, MI

Eugene T. Reilly FDA Ctr. For Devices/Rad Health Rockville, MD

Richard S. Seaberg, MT(ASCP) North Shore University Hospital Manhasset, New York

Advisors

Michael G. Bissell, MD, PhD, MPH Ohio State University Columbus, Ohio Suzanne H. Butch, MA, MT(ASCP), SB The University of Michigan Ann Arbor, Michigan

Randy R. Davis Siemens Medical Solutions Diagnostics Bear, Delaware

Al DeStefano Sysmex America, Inc. Mundelein, Illinois

Jeffrey A. DuBois, PhD NOVA Biomedical Corp. Waltham, Massachusetts

Arden W. Forrey, Jr., PhD, FACB University of Washington Seattle, Washington

David A. Herold, MD, PhD VA (San Diego) Medical Center San Diego, California

Georg E. Hoffmann, MD Trillium GmbH Grafrath, Germany

Stephen Howlett Beckman Coulter, Inc. Miami, Florida

Brian Richard Jackson, MD ARUP Laboratories Salt Lake City, Utah

Gary W. Kramer, PhD National Institute of Standards and Technology Gaithersburg, Maryland Rodney S. Markin, MD, PhD Univ. of Nebraska Medical Center Omaha, Nebraska

Richard A. McPherson, MD Medical College of Virginia Hospital Richmond, Virginia

Paul J. Mountain, MSc, MT(ASCP) Flamborough, Ontario, Canada

David O'Bryan, PhD Hibernia Consulting Portsmouth, New Hampshire

Jeff Quint, PhD Beckman Coulter, Inc. Brea, California

Hiroshi Sekiya Olympus America Inc. Irving, Texas

Russell H. Tomar, MD Chicago, Illinois

Staff

Clinical and Laboratory Standards Institute Wayne, Pennsylvania

Lois M. Schmidt, DA Vice President, Standards Development and Marketing

David E. Sterry, MT(ASCP) Staff Liaison

Patrice E. Polgar Project Manager

Melissa A. Lewis Editor

Acknowledgment

CLSI and the Area Committee on Automation and Informatics gratefully acknowledge the contributions of Ed Heierman, PhD, Abbott Diagnostics; Randy Davis, Siemens Medical Solutions Diagnostics; and Bill Coughlin, BS, MBA, Data Innovations, for their contributions during the revision of the LIS01-A2 consensus standard.

Number 13

I-A2
ŀ

Contents

Abstra	ct		i	
Comm	ittee Me	embership	iii	
Forewo	ord		vii	
1	Scope1			
2	Introduction1			
3	Terminology1			
4	Significance and Use			
5	Physical Layer for Serial Binary Data Exchange			
	5.1 5.2 5.3	Overview Electrical Characteristics Mechanical Characteristics	3 3 5	
6	Data L	ink Layer for Serial Binary Data Exchange	5	
	6.1 6.2 6.3 6.4 6.5 6.6	Overview Establishment Phase (Link Connection) Transfer Phase Termination Phase (Link Release) Error Recovery Restricted Message Characters	5 6 7 9 9 10	
7	Physic	Physical Layer for TCP/IP Data Exchange		
	7.1 7.2 7.3	Overview Electrical Characteristics Mechanical Characteristics	10 10 12	
8	Data Link Layer for TCP/IP Data Exchange			
	8.1 8.2 8.3 8.4 8.5 8.6	Overview Establishment Phase (Link Connection) Transfer Phase Termination Phase (Link Release) Error Recovery Restricted Message Characters	12 13 14 16 16 17	
Refere	nces		18	
Appendix A. Mandatory Information19				
Appendix B. Nonmandatory Information				
Summary of Delegate Comments and Area Committee Responses				
The Quality Management System Approach				
Related	1 CLSI I	Reference Materials	25	

Number 13

Volume 28

LIS01-A2

Foreword

In 2001, ASTM Committee E31 decided to restructure its operations, with the intent of focusing on standards-development issues such as security, privacy, and the electronic health record. Part of the reorganization plan was to transfer responsibility for E31.13 standards to CLSI, then known as NCCLS.

Following this transfer, nine standards (formerly ASTM E792; E1029; E1238; E1246; E1381; E1394; E1466; E1639; and E2118) were redesignated as CLSI/NCCLS standards LIS1 through LIS9.¹⁻⁸ This collection of standards provides a wide variety of information relating to clinical laboratory computer systems. Some included documents are of general interest as reference sources; others represent specifications of primary importance to instrument manufacturers. LIS2 is a revision of the former ASTM E1381-02.

The Area Committee on Automation and Informatics has assumed responsibility for maintaining the documents and will revise or update each document in accord with the CLSI Administrative Procedures. The area committee prioritized LIS1-A as the second standard from this collection to be updated, incorporated into the CLSI document template, and advanced through the CLSI consensus process. The area committee will revise other documents in the series in a similar manner.

With the transfer of the former ASTM standards, the Area Committee on Automation and Informatics has expanded its mission statement to include laboratory information systems. In the future, the area committee will develop additional standards addressing informatics issues, as well as issues related to the integration of patient clinical data.

This document replaces the first edition of the approved guideline, LIS1-A, which was published in 2003. Several changes were made in this edition; among them, TCP/IP communication is now included (Sections 4.4 and 4.5) and the state diagram was replaced (see Appendix A) so it is consistent with the text of the document.

The revisions in this edition of the LIS01 standard are also intended to delineate this document from its former ASTM edition. The title and text have been revised throughout to indicate that this standard applies to clinical laboratory instruments. The term *computer* has been replaced with the term *information* to better reflect the current terminology (ie, LIS) and the headings of Sections 6 and 8 have been changed to make them more specific.

Key Words

data link layer, physical layer, serial binary data exchange, TCP/IP data exchange

Number 13

Volume 28

LIS01-A2

Specification for Low-Level Protocol to Transfer Messages Between Clinical Laboratory Instruments and Computer Systems; Approved Standard— Second Edition

1 Scope

This specification describes the electronic transmission of digital information between clinical laboratory instruments and computer systems.

This specification addresses the low-level protocol used for both serial binary data exchange and TCP/IP data exchange. For message content in the interface between clinical instruments and computer systems, reference CLSI/NCCLS document LIS2.¹

2 Introduction

The clinical laboratory instruments under consideration are those that measure one or more parameters from one or more patient samples. Often they will be automated instruments that measure many parameters from many patient samples. The computer systems considered here are those that are configured to accept instrument results for further processing, storage, reporting, or manipulation. This instrument output may include patient results, quality control results, and other related information. Typically, the computer system will be a clinical laboratory information management system (CLIMS).

The terminology of the International Organization for Standardization (ISO) Reference Model for Open Systems Interconnection (OSI) is generally followed in describing the communications protocol and services.⁹ The electrical and mechanical connection between instrument and computer is described in the Physical Layer sections (see Sections 5 and 7). The methods for establishing communication, error detection, error recovery, and sending and receiving of messages are described in the Data Link Layer sections (see Sections 6 and 8). The data link layer interacts with higher layers in terms of sending and receiving "messages," handles data link connection and release requests, and reports the data link status.

3 Terminology

- **3.1** receiver the device that responds to the sender and accepts the message.
- 3.2 sender the device that has a message to send and initiates the transmission process.
- **3.3** The parts of a communication between instrument and computer are identified by the following terms. The parts are hierarchical and are listed in order of most encompassing first.
- **3.3.1 session** a total unit of communication activity, used in this standard to indicate the events starting with the establishment phase and ending with the termination phase, as described in subsequent sections.
- **3.3.2 message** a collection of related information on a single topic, used here to mean all the identity, tests, and comments sent at one time; **NOTE:** When used with CLSI/NCCLS document LIS2,¹ this term means a record as defined by CLSI/NCCLS document LIS2.¹
- **3.3.3 frame** a subdivision of a message, used to allow periodic communication housekeeping, such as error checks and acknowledgments.