
**Health informatics — Personal health
device communication —**

Part 20601:
**Application profile — Optimized
exchange protocol**

*Informatique de santé — Communication entre dispositifs de santé
personnels —*

Partie 20601: Profil d'application — Protocole d'échange optimisé





COPYRIGHT PROTECTED DOCUMENT

© ISO 2016
© IEEE 2014

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO or IEEE at the respective addresses below.

ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. +41 22 749 01 11
Fax +41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Institute of Electrical and Electronics Engineers, Inc.
3 Park Avenue, New York • NY 10016-5997, USA
E-mail stds.ipr@ieee.org
Web www.ieee.org

Published in Switzerland

Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

IEEE Standards documents are developed within the IEEE Societies and the Standards Coordinating Committees of the IEEE Standards Association (IEEE-SA) Standards Board. The IEEE develops its standards through a consensus development process, approved by the American National Standards Institute, which brings together volunteers representing varied viewpoints and interests to achieve the final product. Volunteers are not necessarily members of the Institute and serve without compensation. While the IEEE administers the process and establishes rules to promote fairness in the consensus development process, the IEEE does not independently evaluate, test, or verify the accuracy of any of the information contained in its standards.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is called to the possibility that implementation of this standard may require the use of subject matter covered by patent rights. By publication of this standard, no position is taken with respect to the existence or validity of any patent rights in connection therewith. ISO/IEEE is not responsible for identifying essential patents or patent claims for which a license may be required, for conducting inquiries into the legal validity or scope of patents or patent claims or determining whether any licensing terms or conditions provided in connection with submission of a Letter of Assurance or a Patent Statement and Licensing Declaration Form, if any, or in any licensing agreements are reasonable or non-discriminatory. Users of this standard are expressly advised that determination of the validity of any patent rights, and the risk of infringement of such rights, is entirely their own responsibility. Further information may be obtained from ISO or the IEEE Standards Association.

ISO/IEEE 11073-20601 was prepared by the IEEE 11073 Standards Committee of the IEEE Engineering in Medicine and Biology Society (as IEEE Std 11073-20601-2014). It was adopted by Technical Committee ISO/TC 215, *Health informatics*, in parallel with its approval by the ISO member bodies, under the “fast-track procedure” defined in the Patent Standards Development Organization cooperation agreement between ISO and IEEE. IEEE is responsible for the maintenance of this document with participation and input from ISO member bodies.

Abstract: Within the context of the ISO/IEEE 11073 family of standards for device communication, this standard defines a common framework for making an abstract model of personal health data available in transport-independent transfer syntax required to establish logical connections between systems and to provide presentation capabilities and services needed to perform communication tasks. The protocol is optimized to personal health usage requirements and leverages commonly used methods and tools wherever possible.

Keywords: IEEE 11073-20601™, medical device communication, personal health devices

The Institute of Electrical and Electronics Engineers, Inc.
3 Park Avenue, New York, NY 10016-5997, USA

Copyright © 2014 by The Institute of Electrical and Electronics Engineers, Inc.
All rights reserved. Published 10 October 2014. Printed in the United States of America.

IEEE and IEEE 802 are registered trademarks in the U.S. Patent & Trademark Office, owned by The Institute of Electrical and Electronics Engineers, Incorporated.

PDF: ISBN 978-0-7381-9314-4 STD98793
Print: ISBN 978-0-7381-9315-1 STDPD98793

IEEE prohibits discrimination, harassment, and bullying.

For more information, visit <http://www.ieee.org/web/aboutus/whatis/policies/p9-26.html>.

No part of this publication may be reproduced in any form, in an electronic retrieval system or otherwise, without the prior written permission of the publisher.

Important Notices and Disclaimers Concerning IEEE Standards Documents

IEEE documents are made available for use subject to important notices and legal disclaimers. These notices and disclaimers, or a reference to this page, appear in all standards and may be found under the heading “Important Notice” or “Important Notices and Disclaimers Concerning IEEE Standards Documents.”

Notice and Disclaimer of Liability Concerning the Use of IEEE Standards Documents

IEEE Standards documents (standards, recommended practices, and guides), both full-use and trial-use, are developed within IEEE Societies and the Standards Coordinating Committees of the IEEE Standards Association (“IEEE-SA”) Standards Board. IEEE (“the Institute”) develops its standards through a consensus development process, approved by the American National Standards Institute (“ANSI”), which brings together volunteers representing varied viewpoints and interests to achieve the final product. Volunteers are not necessarily members of the Institute and participate without compensation from IEEE. While IEEE administers the process and establishes rules to promote fairness in the consensus development process, IEEE does not independently evaluate, test, or verify the accuracy of any of the information or the soundness of any judgments contained in its standards.

IEEE does not warrant or represent the accuracy or content of the material contained in its standards, and expressly disclaims all warranties (express, implied and statutory) not included in this or any other document relating to the standard, including, but not limited to, the warranties of: merchantability; fitness for a particular purpose; non-infringement; and quality, accuracy, effectiveness, currency, or completeness of material. In addition, IEEE disclaims any and all conditions relating to: results; and workmanlike effort. IEEE standards documents are supplied “AS IS” and “WITH ALL FAULTS.”

Use of an IEEE standard is wholly voluntary. The existence of an IEEE standard does not imply that there are no other ways to produce, test, measure, purchase, market, or provide other goods and services related to the scope of the IEEE standard. Furthermore, the viewpoint expressed at the time a standard is approved and issued is subject to change brought about through developments in the state of the art and comments received from users of the standard.

In publishing and making its standards available, IEEE is not suggesting or rendering professional or other services for, or on behalf of, any person or entity nor is IEEE undertaking to perform any duty owed by any other person or entity to another. Any person utilizing any IEEE Standards document, should rely upon his or her own independent judgment in the exercise of reasonable care in any given circumstances or, as appropriate, seek the advice of a competent professional in determining the appropriateness of a given IEEE standard.

IN NO EVENT SHALL IEEE BE LIABLE FOR ANY DIRECT, INDIRECT, INCIDENTAL, SPECIAL, EXEMPLARY, OR CONSEQUENTIAL DAMAGES (INCLUDING, BUT NOT LIMITED TO: PROCUREMENT OF SUBSTITUTE GOODS OR SERVICES; LOSS OF USE, DATA, OR PROFITS; OR BUSINESS INTERRUPTION) HOWEVER CAUSED AND ON ANY THEORY OF LIABILITY, WHETHER IN CONTRACT, STRICT LIABILITY, OR TORT (INCLUDING NEGLIGENCE OR OTHERWISE) ARISING IN ANY WAY OUT OF THE PUBLICATION, USE OF, OR RELIANCE UPON ANY STANDARD, EVEN IF ADVISED OF THE POSSIBILITY OF SUCH DAMAGE AND REGARDLESS OF WHETHER SUCH DAMAGE WAS FORESEEABLE.

Translations

The IEEE consensus development process involves the review of documents in English only. In the event that an IEEE standard is translated, only the English version published by IEEE should be considered the approved IEEE standard.

Official statements

A statement, written or oral, that is not processed in accordance with the IEEE-SA Standards Board Operations Manual shall not be considered or inferred to be the official position of IEEE or any of its committees and shall not be considered to be, or be relied on as, a formal position of IEEE. At lectures, symposia, seminars, or educational courses, an individual presenting information on IEEE standards shall make it clear that his or her views should be considered the personal views of that individual rather than the formal position of IEEE.

Comments on standards

Comments for revision of IEEE Standards documents are welcome from any interested party, regardless of membership affiliation with IEEE. However, IEEE does not provide consulting information or advice pertaining to IEEE Standards documents. Suggestions for changes in documents should be in the form of a proposed change of text, together with appropriate supporting comments. Since IEEE standards represent a consensus of concerned interests, it is important that any responses to comments and questions also receive the concurrence of a balance of interests. For this reason, IEEE and the members of its societies and Standards Coordinating Committees are not able to provide an instant response to comments or questions except in those cases where the matter has previously been addressed. For the same reason, IEEE does not respond to interpretation requests. Any person who would like to participate in discussions to an IEEE standard is welcome to join the relevant IEEE working group.

Comments on standards should be submitted to the following address:

Secretary, IEEE-SA Standards Board
445 Hoes Lane
Piscataway, NJ 08854 USA

Laws and regulations

Users of IEEE Standards documents should consult all applicable laws and regulations. Compliance with the provisions of any IEEE Standards document does not imply compliance to any applicable regulatory requirements. Implementers of the standards are responsible for observing or referring to the applicable regulatory requirements. IEEE does not, with publication of its standards, intend to urge action that is not in compliance with applicable laws, and these documents may not be construed as doing so.

Copyrights

IEEE draft and approved standards are copyrighted by IEEE under U.S. and international copyright laws. They are made available by IEEE and are adopted for a wide variety of both public and private uses. These include both use, by reference, in laws and regulations, and use in private self-regulation, standardization, and the promotion of engineering practices and methods. By making these documents available for use and adoption by public authorities and private users, IEEE does not waive any rights in copyright to the documents.

Photocopies

Subject to payment of the appropriate fee, IEEE will grant users a limited, non-exclusive license to photocopy portions of any individual standard for company or organizational internal use or individual, non-commercial use only. To arrange for payment of licensing fees, please contact Copyright Clearance Center, Customer Service, 222 Rosewood Drive, Danvers, MA 01923 USA; +1 978 750 8400. Permission to photocopy portions of any individual standard for educational classroom use can also be obtained through the Copyright Clearance Center.

Updating of IEEE Standards documents

Users of IEEE Standards documents should be aware that these documents may be superseded at any time by the issuance of new editions or may be amended from time to time through the issuance of amendments, corrigenda, or errata. An official IEEE document at any point in time consists of the current edition of the document together with any amendments, corrigenda, or errata then in effect.

Every IEEE standard is subjected to review at least every ten years. When a document is more than ten years old and has not undergone a revision process, it is reasonable to conclude that its contents, although still of some value, do not wholly reflect the present state of the art. Users are cautioned to check to determine that they have the latest edition of any IEEE standard.

In order to determine whether a given document is the current edition and whether it has been amended through the issuance of amendments, corrigenda, or errata, visit the IEEE-SA Website at <http://ieeexplore.ieee.org/xpl/standards.jsp> or contact IEEE at the address listed previously. For more information about the IEEE-SA or IEEE's standards development process, visit the IEEE-SA Website at <http://standards.ieee.org>.

Errata

Errata, if any, for all IEEE standards can be accessed on the IEEE-SA Website at the following URL: <http://standards.ieee.org/findstds/errata/index.html>. Users are encouraged to check this URL for errata periodically.

Patents

Attention is called to the possibility that implementation of this standard may require use of subject matter covered by patent rights. By publication of this standard, no position is taken by the IEEE with respect to the existence or validity of any patent rights in connection therewith. If a patent holder or patent applicant has filed a statement of assurance via an Accepted Letter of Assurance, then the statement is listed on the IEEE-SA Website at <http://standards.ieee.org/about/sasb/patcom/patents.html>. Letters of Assurance may indicate whether the Submitter is willing or unwilling to grant licenses under patent rights without compensation or under reasonable rates, with reasonable terms and conditions that are demonstrably free of any unfair discrimination to applicants desiring to obtain such licenses.

Essential Patent Claims may exist for which a Letter of Assurance has not been received. The IEEE is not responsible for identifying Essential Patent Claims for which a license may be required, for conducting inquiries into the legal validity or scope of Patents Claims, or determining whether any licensing terms or conditions provided in connection with submission of a Letter of Assurance, if any, or in any licensing agreements are reasonable or non-discriminatory. Users of this standard are expressly advised that determination of the validity of any patent rights, and the risk of infringement of such rights, is entirely their own responsibility. Further information may be obtained from the IEEE Standards Association.

Participants

At the time this standard was submitted to the IEEE-SA Standards Board for approval, the Personal Health Devices Working Group had the following membership:

Daidi Zhong, *Co-Chair*
Michael J. Kirwan, *Co-Chair*
Douglas P. Bogia, *Co-Chair*

Charles R. Abbruscato
Nabil Abujbara
Maher Abuzaid
Manfred Aigner
Jorge Alberola
Karsten Alders
Murtaza Ali
Rolf Ambuehl
David Aparisi
Lawrence Arne
Diego B. Arquillo
Serafin Arroyo
Muhammad Asim
Merat Bagha
Doug Baird
David Baker
Anindya Bakshi
Ananth Balasubramanian
Sunlee Bang
M. Jonathan Barkley
Gilberto Barrón
David Bean
John Bell
Rudy Belliardi
Kathryn M. Bennett
Daniel Bernstein
George A. Bertos
Chris Biernacki
Ola Björnsne
Thomas Blackadar
Marc Blanchet
Thomas Bluethner
Xavier Boniface
Shannon Boucousis
Julius Broma
Lyle G. Bullock
Bernard Burg
Chris Burns
Anthony Burns
Jeremy Byford
Satya C. Challoji
John C. Carey
Santiago Carot-Nemesio
Gandy W. Carroll
Simon Carter
Seungchul Chae
Rahul Chauhan
James Cheng
Peggy Chien
Chia-Chin Chong
Saeed A. Choudhary

Jinhan Chung
Malcolm Clarke
John A. Cogan
John T. Collins
Cory Condek
Todd H. Cooper
David Cornejo
Douglas Coup
Nigel Cox
Hans Crommenacker
Tomio Crosley
David Culp
Allen Curtis
Ndifor Cyril Fru
Jesús Daniel Trigo
Eyal Dassau
David Davenport
Russell Davis
Ed Day
Sushil K. Deka
Pedro de-las-Heras-Guiro
Jim DelloStitto
Matthew d'Entremont
Lane Desborough
Kent Dickson
Hyoungho Do
Xioliang Duan
Bruno Dabreuil
Jacob Ehrensvard
Fredrik Einberg
Roger M. Ellingson
Michihiro Enokida
Javier Escayola Calvo
Leonardo Estevez
Roger Feeley
Bosco T. Fernandes
Christoph Fischer
Morten Flintrup
Joseph W. Forler
Russell Foster
Eric Freudenthal
Matthias Frohner
Ken Fuchs
Jing Gao
Marcus Garbe
John Garguilo
Rick Geimer
Igor Gejdos
Ferenc Gerbovics
Nicolae Goga
Julian Goldman

Raul Gonzalez Gomez
Chris Gough
Channa Gowda
Charles M. Gropper
Amit Gupta
Jeff Guttmacher
Rasmus Haahr
Christian Habermann
Michael Hagerty
Jerry Hahn
Robert Hall
Nancy J. Hamming
Keye L. Hampton
Steve Hanke
Jordan Hartmann
Kai Hassing
Marc Daniel Haunschild
Wolfgang Heck
Charles Henderson
Jun-Ho Her
Takashi Hibino
Timothy L. Hirou
Allen Hobbs
Alex Holland
Arto Holopainen
Robert Hoy
Frank Hsu
Anne Huang
Sen-Der Huang
Zhiqiang Huang
Ron Huby
Robert D. Hughes
David Hughes
Jiyoung Huh
Hugh Hunter
Hitoshi Ikeda
Yutaka Ikeda
Philip O. Isaacson
Atsushi Ito
Michael Jaffe
Praduman Jain
Danny Jochelson
Chris Johnson
Phaneeth Junga
Akiyoshi Kabe
Steve Kahle
Tomio Kamioka
Kei Kariya
Andy Kaschl
Junzo Kashiara
Kohichi Kashiwagi

The following members of the balloting committee voted on this standard. Balloters may have voted for approval, disapproval, or abstention.

Hector Barron Gonzalez
 Pieter Botman
 Lyle G. Bullock
 Juan Carreon
 Randy W. Carroll
 Lawrence Catchpole
 Jianwen Chen
 Keith Chow
 Donald Cragun
 Paul Croll
 Russell Davis
 Douglas Dorr
 Sourav Dutta
 Christoph Fischer

David Friscia
 David Fuschi
 Randall Groves
 Kai Hassing
 Werner Hoelzl
 Ruimin Hu
 Noriyuki Ikeuchi
 Akio Iso
 Atsushi Ito
 Raj Jain
 Junghoon Jee
 Piotr Karocki
 Stuart Kerry
 Geoff Ladwig
 Richard Lancaster

Charles Ngethe
 Melvin I. Reynolds
 Terence Rout
 Bartien Sayogo
 Lars Schmitt
 Carl Singer
 Kapil Sood
 Raymond A. Strickland
 Walter Struppler
 Jiande Sun
 Hung-Yu Wei
 Jan Wittenber
 Oren Yuen
 Daidi Zhong

When the IEEE-SA Standards Board approved this standard on 21 August 2014, it had the following membership:

John Kulick, Chair
Jon Walter Rosdahl, Vice-chair
Richard H. Hulett, Past Chair
Konstantinos Karachalios, Secretary

Peter Balma
 Farooq Bari
 Ted Burse
 Clint Chaplain
 Stephen Dukes
 Jean-Phillippe Faure
 Gary Hoffman

Michael Janezic
 Jeffrey Katz
 Joseph L. Koepfinger *
 David Law
 Hung Ling
 Oleg Logvinov
 T. W. Olsen
 Glenn Parsons

Ron Peterson
 Adrian Stephens
 Peter Sutherland
 Yatin Trivedi
 Phil Winston
 Don Wright
 Yu Yuan

*Member Emeritus

Also included are the following non-voting IEEE-SA Standards Board liaisons:

Richard DeBlasio, *DOE Representative*
 Michael Janezic, *NIST Representative*

Don Messina
IEEE-SA Content Publishing

Kathryn Bennett
IEEE-SA Technical Community Programs

Introduction

This introduction is not part of IEEE Std 11073-20601-2014, Health informatics—Personal health device communication—Part 20601: Application profile—Optimized Exchange Protocol.

ISO and IEEE 11073 standards enable communication between medical devices and external computer systems. This standard and corresponding IEEE 11073-104zz standards address a need for a simplified and optimized communication approach for personal health devices, which may or may not be regulated devices. These standards align with, and draw upon, the existing clinically focused standards to provide easy management of data from either a clinical or personal health device.

This document addresses a need for an openly defined, independent standard for converting the collected information into an interoperable transmission format so the information can be exchanged between agents and managers.

Other closely related standards include the following:

- IEEE Std 11073-00103-2012 [B5]^a provides an overview of the personal health space and defines the underlying use cases and usage models.
- ISO/IEEE 11073-10101 [B16] documents the nomenclature terms that can be used.
- ISO/IEEE 11073-10201:2004 [B17] documents the extensive domain information model (DIM) leveraged by this standard.
- ISO/IEEE 11073-104zz standards define specific device specializations. For example, ISO/IEEE 11073-10404 [B18] defines how interoperable pulse oximeters work.
- ISO/IEEE 11073-20101:2004 [B21] defines the medical device encoding rules (MDER) used in this standard.

^a The numbers in brackets correspond to the numbers of the bibliography in Annex K.

Contents

1. Overview	1
1.1 Scope	1
1.2 Purpose	1
1.3 Context	2
2. Normative references.....	5
3. Definitions, acronyms, and abbreviations	5
3.1 Definitions	5
3.2 Acronyms and abbreviations	6
4. Guiding principles	7
5. Introduction to IEEE 11073 personal health devices.....	8
5.1 General	8
5.2 Domain information model (DIM)	9
5.3 Service model	9
5.4 Communication model	9
5.5 Compliance with other standards.....	9
5.6 Security.....	9
6. Personal health device DIM	10
6.1 General	10
6.2 Nomenclature usage	11
6.3 Personal health object class definitions	12
6.3.1 General.....	12
6.3.2 MDS class	14
6.3.3 Metric class.....	22
6.3.4 Numeric class.....	28
6.3.5 RT-SA class	31
6.3.6 Enumeration class	33
6.3.7 PM-store class.....	35
6.3.8 PM-segment class	41
6.3.9 Scanner classes.....	46
6.4 Information model extensibility rules.....	57
7. Personal health device service model.....	58
7.1 General	58
7.2 Association service	58
7.3 Object access services.....	58
7.4 Specific application of object access EVENT REPORT services for personal health devices.....	59
7.4.1 General.....	59
7.4.2 Confirmed and unconfirmed event reports.....	59
7.4.3 Configuration event report.....	59
7.4.4 Agent- and manager-initiated measurement data transmission.....	63
7.4.5 Variable, fixed, and grouped format event reports.....	64
7.4.6 Single-person and multiple-person event reports.....	65

7.4.7 Temporarily stored measurements	66
8. Communication model	66
8.1 General	66
8.2 System context.....	67
8.3 Communications characteristics	68
8.3.1 General.....	68
8.3.2 Common communications characteristics.....	69
8.3.3 Reliable communications characteristics	70
8.3.4 Best-effort communications characteristics	70
8.4 State machines	71
8.4.1 Agent state machine.....	71
8.4.2 Manager state machine.....	74
8.4.3 Timeout variables.....	75
8.5 Connected procedure	76
8.5.1 General.....	76
8.5.2 Entry conditions.....	76
8.5.3 Normal procedures.....	76
8.5.4 Exit conditions	77
8.5.5 Error conditions	77
8.6 Unassociated procedure	77
8.6.1 General.....	77
8.6.2 Entry conditions	77
8.6.3 Normal procedures.....	77
8.6.4 Exit conditions	77
8.6.5 Error conditions	77
8.7 Associating procedure	78
8.7.1 General.....	78
8.7.2 Entry conditions	78
8.7.3 Normal procedures.....	78
8.7.4 Exit conditions	82
8.7.5 Error conditions	82
8.7.6 Test association	83
8.8 Configuring procedure.....	84
8.8.1 General.....	84
8.8.2 Entry conditions	84
8.8.3 Normal procedures.....	84
8.8.4 Exit conditions	87
8.8.5 Error conditions	88
8.9 Operating procedure	88
8.9.1 General.....	88
8.9.2 Entry conditions	88
8.9.3 Normal procedures.....	88
8.9.4 Exit conditions	100
8.9.5 Error conditions	101
8.10 Disassociating procedure	102
8.10.1 General.....	102
8.10.2 Entry conditions	102
8.10.3 Normal procedures.....	103
8.10.4 Exit conditions	103
8.10.5 Error conditions	103
8.11 Message encoding.....	103
8.12 Time coordination.....	104
8.12.1 General.....	104
8.12.2 Absolute time	104

8.12.3 Base time with offset.....	106
8.12.4 Relative time	106
8.12.5 High-resolution relative time	107
9. Conformance model	108
9.1 Applicability	108
9.2 Conformance specification	108
9.3 Implementation conformance statements (ICSs)	109
9.4 General conformance.....	109
9.4.1 General ICS.....	109
9.4.2 Minimum requirements ICS.....	111
9.4.3 Service support ICS	112
9.5 Device additions/extensions ICS	113
9.5.1 General additions/extensions ICS	113
9.5.2 Personal health device DIM object and class (POC) ICS	114
9.5.3 POC attribute ICS	114
9.5.4 POC behavior ICS.....	115
9.5.5 POC notification ICS	115
9.5.6 POC nomenclature ICS.....	116
Annex A (normative) ASN.1 definitions.....	117
Annex B (informative) Scale and range specification example.....	151
Annex C (informative) The PM-store concept	153
Annex D (informative) Transport profile types.....	158
Annex E (normative) State tables.....	161
Annex F (normative) Medical device encoding rules (MDER).....	181
Annex G (informative) Encoded data type definitions.....	193
Annex H (informative) Examples.....	213
Annex I (normative) Nomenclature codes.....	228
Annex J (informative) Derivation and modification history.....	233
Annex K (informative) Bibliography	236

Health informatics—Personal health device communication

Part 20601: Application profile— Optimized Exchange Protocol

IMPORTANT NOTICE: IEEE Standards documents are not intended to ensure safety, security, health, or environmental protection, or ensure against interference with or from other devices or networks. Implementers of IEEE Standards documents are responsible for determining and complying with all appropriate safety, security, environmental, health, and interference protection practices and all applicable laws and regulations.

This IEEE document is made available for use subject to important notices and legal disclaimers. These notices and disclaimers appear in all publications containing this document and may be found under the heading “Important Notice” or “Important Notices and Disclaimers Concerning IEEE Documents.” They can also be obtained on request from IEEE or viewed at <http://standards.ieee.org/IPR/disclaimers.html>.

1. Overview

1.1 Scope

Within the context of the ISO/IEEE 11073 personal health device standard family, this standard defines an optimized exchange protocol and modeling techniques to be used by implementers of personal health devices to create interoperability between device types and vendors. This standard establishes a common framework for an abstract model of personal health data available in transport-independent transfer syntax required to establish logical connections between systems and to provide presentation capabilities and services needed to perform communication tasks. The protocol is optimized to personal health usage requirements and leverages commonly used methods and tools wherever possible.

1.2 Purpose

This standard addresses a need for an openly defined, independent standard for controlling information exchange to and from personal health devices and managers (e.g., cell phones, personal computers, personal health appliances, and set top boxes). Interoperability is key to growing the potential market for these devices and enabling people to be better informed participants in the management of their health.