STN	Zdravotnícka informatika. Komunikácia s osobným zdravotným prístrojom. Časť 20601: Aplikačný profil. Optimalizovaný výmenný protokol (ISO/IEEE 11073-20601: 2016 vrátane Cor 1: 2016).	STN EN ISO 11073-20601	
		84 8107	

Health informatics - Personal health device communication - Part 20601: Application profile - Optimized exchange protocol (ISO/IEEE 11073-20601:2016, including Cor 1:2016)

Táto norma obsahuje anglickú verziu európskej normy. This standard includes the English version of the European Standard.

Táto norma bola oznámená vo Vestníku ÚNMS SR č. 01/17

Obsahuje: EN ISO 11073-20601:2016, ISO/IEEE 11073-20601:2016, ISO/IEEE 11073-20601:2016/Cor 1:2016

Oznámením tejto normy sa ruší STN EN ISO 11073-20601 (84 8107) z augusta 2011

123743

Úrad pre normalizáciu, metrológiu a skúšobníctvo SR, 2017 Podľa zákona č. 264/1999 Z. z. v znení neskorších predpisov sa môžu slovenské technické normy rozmnožovať a rozširovať iba so súhlasom Úradu pre normalizáciu, metrológiu a skúšobníctvo SR.

EUROPEAN STANDARD NORME EUROPÉENNE

EN ISO 11073-20601

EUROPÄISCHE NORM

August 2016

ICS 35.240.80

Supersedes EN ISO 11073-20601:2011

English Version

Health informatics - Personal health device communication - Part 20601: Application profile -Optimized exchange protocol (ISO/IEEE 11073-20601:2016, including Cor 1:2016)

Informatique de santé - Communication entre dispositifs de santé personnels - Partie 20601: Profil d'application - Protocole d'échange optimisé (ISO/IEEE 11073-20601:2016, y compris Cor 1:2016) Medizinische Informatik - Kommunikation von Geräten für die persönliche Gesundheit - Teil 20601: Anwendungsprofil - Optimiertes Datenübertragungsprotokoll (ISO/IEEE 11073-20601:2016, einschließlich Cor 1:2016)

This European Standard was approved by CEN on 21 February 2016.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN-CENELEC Management Centre or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and United Kingdom.



EUROPEAN COMMITTEE FOR STANDARDIZATION COMITÉ EUROPÉEN DE NORMALISATION EUROPÄISCHES KOMITEE FÜR NORMUNG

CEN-CENELEC Management Centre: Avenue Marnix 17, B-1000 Brussels

© 2016 CEN All rights of exploitation in any form and by any means reserved worldwide for CEN national Members.

Ref. No. EN ISO 11073-20601:2016 E

Contents	Page
European foreword	

European foreword

This document (EN ISO 11073-20601:2016 including Cor 1:2016) has been prepared by Technical Committee ISO/TC 215 "Health informatics" in collaboration with Technical Committee CEN/TC 251 "Health informatics" the secretariat of which is held by NEN.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by February 2017, and conflicting national standards shall be withdrawn at the latest by February 2017.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN [and/or CENELEC] shall not be held responsible for identifying any or all such patent rights.

This document supersedes EN ISO 11073-20601:2011.

According to the CEN-CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

Endorsement notice

The text of ISO/IEEE 11073-20601:2016 including Cor 1:2016 has been approved by CEN as EN ISO 11073-20601:2016 without any modification.

INTERNATIONAL ISO/IEEE STANDARD



Second edition 2016-06-15

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





© 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 address below.

ISO copyright office Case postale 401 • CH-1214 Vernier, Geneva Tel. + 41 22 749 01 11 Fax + 41 22 749 09 47 E-mail copyright@iso.org Web www.iso.org Published in Switzerland 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

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 Comittee 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 Partner 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

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.

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

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 revisions 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 standard are responsible for observing or referring to the applicable regulatory regulatory requirements. IEEE does not, by the 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.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.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.jsp or contact IEEE at the address listed previously.

Errata

Errata, if any, for all IEEE standards can be accessed on the IEEE-SA Website at the following URL: <u>http://standards.ieee.org/findstds/errata/index.html</u>. 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:

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 Björsne Thomas Blackadar Marc Blanchet Thomas Bluethner Xavier Boniface Shannon Boucousis Julius Broma Lyle G. Bullock Bernard Burg Chris Burns Anthony Butt Jeremy Byford-Rew Satya Calloji Carole C. Carey Santiago Carot-Nemesio Randy W. Carroll Simon Carter Seungchul Chae Rahul Chauhan James Cheng Peggy Chien Chia-Chin Chong Saeed A. Choudhary

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

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-Quiros Jim DelloStritto Matthew d'Entremont Lane Desborough Kent Dicks Hyoungho Do Xiaolian Duan Brian Dubreuil Jakob Ehrensvard Fredrik Einberg Roger M. Ellingson Michihiro Enokida Javier Escavola 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 Nathaniel Hamming Rickey L. Hampton Sten 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 Kashihara Kohichi Kashiwagi

STN EN ISO 11073-20601: 2017

Ralph Kent Laurie M. Kermes Ikuo Keshi Junhyung Kim Min-Joon Kim Minho Kim Taekon Kim Tetsuya Kimura Alfred Kloos Jeongmee Koh Jean-Marc Koller John Koon Patty Krantz Alexander Kraus Ramesh Krishna Geoffrey Kruse Falko Kuester Rafael Lajara Pierre Landau Jaechul Lee JongMuk Lee Kyong Ho Lee Rami Lee Sungkee Lee Woojae Lee Yonghee Lee Joe Lenart Kathryn A. Lesh Qiong Li Ying Li Patrick Lichter Jisoon Lim Joon-Ho Lim John Lin Jiajia Liu Wei-Jung Lo Charles Lowe Don Ludolph Christian Luszick Bob MacWilliams Srikkanth Madhurbootheswaran Romain Marmot Sandra Martinez Miguel Martínez de Espronceda Cámara Peter Mayhew Jim McCain László Meleg Alexander Mense Ethan Metsger Yu Miao Jinsei Miyazaki Erik Moll Darr Moore Piotr Murawski Soundharya Nagasubramanian Jae-Wook Nah Alex Neefus Trong-Nghia Nguyen-Dobinsky Michael E. Nidd Tetsu Nishimura

Jim Niswander Hiroaki Niwamoto Thomas Norgall Anand Noubade Yoshiteru Nozoe Abraham Ofek Brett Olive Begonya Otal **Charles Palmer** Bud Panjwani Carl Pantiskas Harry P. Pappas Mikey Paradis Hanna Park Jong-Tae Park Myungeun Park Soojun Park Phillip E. Pash TongBi Pei Soren Petersen James Petisce Peter Piction Michael Pliskin Jeff Price Harald Prinzhorn John Quinlan Arif Rahman Tanzilur Rahman Steve Ray Phillip Raymond Tim Reilly Barry Reinhold Brian Reinhold Melvin I. Reynolds John G. Rhoads Jeffrey S. Robbins Moskowitz Robert Timothy Robertson David Rosales **Bill Saltzstein** Benedikt Salzbrunn Giovanna Sannino Jose A. Santos-Cadenas Stefan Sauermann John Sawyer Guillaume Schatz Alois Schloegl Paul S. Schluter Lars Schmitt Mark G. Schnell Richard A. Schrenker Antonio Scorpiniti Kwang Seok Seo Riccardo Serafin Sid Shaw Frank Shen Liqun Shen Bozhi Shi Min Shih Mazen Shihabi Redmond Shouldice

Sternly K. Simon Marjorie Skubic Robert Smith Ivan Soh Motoki Sone Emily Sopensky Rajagopalan Srinivasan Andreas Staubert Nicholas Steblay Beth Stephen Lars Steubesand John (Ivo) Stivoric Raymond A. Strickland Hermanni Suominen Lee Surprenant Ravi Swami Ray Sweidan Jin Tan Haruyuyki Tatsumi John W. Thomas Brad Tipler Jonas Tirén James Tomcik Janet Traub Gary Tschautscher Masato Tsuchid Ken Tubman Yoshihiro Uchida Sunil Unadkat Fabio Urbani Philipp Urbauer Laura Vanzago Alpo Värri Ciro de la Vega Dalimar Velez Naveen Verma Rudi Voon Isobel Walker David Wang Jerry P. Wang Yao Wang Yi Wang Steve Warren Fujio Watanabe Toru Watsuji Mike Weng Kathleen Wible Paul Williamson Jan Wittenber Jia-Rong Wu Will Wykeham Ariton Xhafa Junjie Yang Ricky Yang Melanie Yeung Done-Sik Yoo Jason Zhang Zhiqiang Zhang Thomas Zhao Miha Zoubek Szymon Zysko

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 nonvoting 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	
1.3 Context	
	2
2. Normative references	5
3. Definitions, acronyms, and abbreviations	5
3.1 Definitions	5
3.2 Acronyms and abbreviations	
	0
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	
5.4 Communication model	
5.5 Compliance with other standards	
5.6 Security	9
6. Personal health device DIM	. 10
6.1 General	. 10
6.2 Nomenclature usage	
6.3 Personal health object class definitions	
6.3.1 General	
6.3.2 MDS class	. 14
6.3.3 Metric class	. 22
6.3.4 Numeric class	
6.3.5 RT-SA class	
6.3.6 Enumeration class	
6.3.7 PM-store class	
6.3.8 PM-segment class	
6.3.9 Scanner classes.	
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	
7.4 Specific application of object access EVENT REPORT services for personal health devices	
7.4.1 General	
7.4.2 Confirmed and unconfirmed event reports	
7.4.3 Configuration event report	
7.4.4 Agent- and manager-initiated measurement data transmission	
7.4.5 Variable, fixed, and grouped format event reports	
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	
8.3 Communications characteristics	
8.3.1 General	
8.3.2 Common communications characteristics	
8.3.3 Reliable communications characteristics	
8.3.4 Best-effort communications characteristics	
8.4 State machines	
8.4.1 Agent state machine	
8.4.2 Manager state machine	
8.4.3 Timeout variables	
8.5 Connected procedure	
8.5.1 General	
8.5.2 Entry conditions	
8.5.3 Normal procedures	
8.5.4 Exit conditions	
8.5.5 Error conditions	
8.6 Unassociated procedure	
8.6.1 General.	
8.6.2 Entry conditions	
8.6.3 Normal procedures	
8.6.4 Exit conditions	
8.6.5 Error conditions	
8.7 Associating procedure	
8.7.1 General	
8.7.2 Entry conditions	78
8.7.3 Normal procedures	
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	
8.8.4 Exit conditions	87
8.8.5 Error conditions	88
8.9 Operating procedure	
8.9.1 General	88
8.9.2 Entry conditions	
8.9.3 Normal procedures	
8.9.4 Exit conditions	
8.9.5 Error conditions	
8.10 Disassociating procedure	
8.10.1 General	
8.10.2 Entry conditions	
8.10.3 Normal procedures	
8.10.4 Exit conditions	
8.10.5 Error conditions	
8.11 Message encoding	
8.12 Time coordination	
8.12.1 General	
8.12.2 Absolute time	104

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

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.

1.3 Context

Figure 1 shows categories and typical devices supporting the personal health space. Agents (e.g., blood pressure monitors, weighing scales, and pedometers) collect information about a person (or persons) and transfer the information to a manager (e.g., cell phone, health appliance, or personal computer) for collection, display, and possible later transmission. The manager may also forward the data to remote support services for further analysis. The information is available from a range of domains including disease management, health and fitness, or aging independently applications.

The communication path between agent and manager is assumed to be a logical point-to-point connection. Generally, an agent communicates with a single manager at any point in time. A manager may communicate with multiple agents simultaneously using separate point-to-point connections.

The overlay shows the focus area of the IEEE 11073[™] Personal Health Devices Working Group. The primary concentration is the interface and data exchange between the agents and manager. However, this interface cannot be created in isolation by ignoring the remainder of the solution space. Remaining cognizant of the entire system helps to move data reasonably from the agents all the way to the remote support services when necessary. This path may include converting the data format, exchange protocols, and transport protocols across different interfaces. Much of the standardization effort is outside of the scope of the Personal Health Devices Working Group; however, aligning all standardization efforts allows data to flow seamlessly through the overall set of systems.

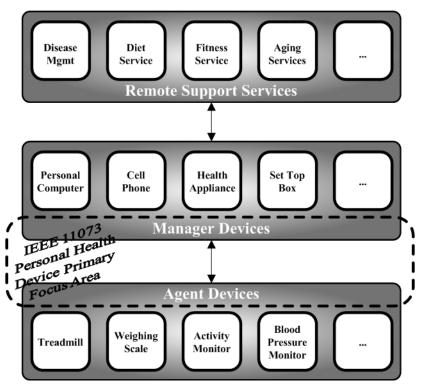


Figure 1—Overall context of work

Figure 2 shows a hierarchical view of the architecture of an agent or manager superimposed with a view of the related standards. The application layers are, for the most part, not specific to any particular transport. Where necessary, this standard identifies assumptions that require direct support by a transport or a "shim" layer above the transport. This approach allows support for various transports. The definition of the transports is outside the scope of this standard and the working group.

Above the transport layer is the Optimized Exchange Protocol (described in this standard). This protocol consists of two aspects: the application layer services and the definition of the data exchange protocol between agents and managers. The application layer services provide the protocol for connection management and reliable transfer of actions and data between agent and manager. The data exchange protocol defines the commands, agent configuration information, data format, and overall protocol. The Optimized Exchange Protocol provides the basis to support any type of agent. For a specific device type, the reader is directed to the device specialization for that agent to understand the capabilities of the device and its implementation according to this standard. The device specialization indicates which aspects of this standard to comprehend and where further information to implement the device is found.

Above the exchange protocol are device specializations that describe specific details relative to the particular agent (e.g., blood pressure monitor, weighing scale, or pedometer). The specializations describe the details of how these agents work and act as a detailed description for creating a specific type of agent. Additionally, they provide reference to a related standard for further details. The standard numbers reserved for device specializations range from IEEE Std 11073-10401 through IEEE Std 11073-10499, inclusive. When the collection of standards is being referenced, the term *IEEE 11073-104zz* is used where zz could be any number in the range from 01 to 99, inclusive.

Some device specializations describe broad categories of device types (e.g., the IEEE 11073-10441TM model device types that promote cardiovascular activity such as step counters or exercise cycles). Other device specializations have a narrow focus on a single device type (e.g., IEEE 11073-10408TM model thermometers). Specializations that address one or more device types may also define *profiles*. A profile further constrains the model defined in a specialization to increase interoperability (e.g., the step counter profile utilizes a limited portion of IEEE 11073-10441 modeling).

The IEEE Std 11073-00103-2012 [B5]¹ technical report describes the overall personal health space with further definition of the underlying use cases and usage models.

	-00103 Technical Report - Overview					7	
-10404 Pulse Oximeter	-10407 Blood Pressure	-10408	e Spe -10415 Weighing Scale	-10417	tions -10441/- 10442 H&F Cardio/ Strength	-10471 Activity Hub	Phase II
-20601 Optimized Exchange Protocol							
	Transports						

Figure 2—Document map

¹ The numbers in brackets correspond to the numbers of the bibliography in Annex K.

The personal health device specializations are not being created independently of all other standards. There are a number of existing standards generated for clinical environments upon which these standards draw. Figure 3 shows the relationship to the remainder of the IEEE 11073 documents. There are two types of relationships:

- Drawing ideas and/or content from the other documents (dashed lines)
- Leveraging information from the other document and introducing new content into that document to support this standard (solid lines)

This standard imports information from ISO/IEEE 11073-10201:2004 [B17] and ISO/IEEE 11073-20101:2004 [B21] as normative annexes. If there is a discrepancy between these standards, this standard takes priority. Because of the reuse of constructs from these standards, some of the names appear to be more clinically focused [e.g., medical device system (MDS) instead of personal health device system]; however, to maintain consistency, the traditional names have been preserved.

This standard replicates relevant portions of ISO/IEEE 11073-10101 [B16] and incorporates new nomenclature codes.

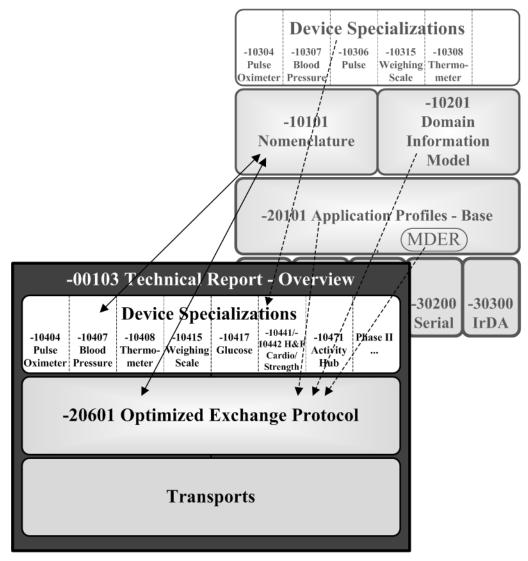


Figure 3—Relationship to other IEEE 11073 documents

2. Normative references

The following referenced documents are indispensable for the application of this standard (i.e., they must be understood and used; therefore, each referenced document is cited in the text and its relationship to this standard is explained). For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments or corrigenda) applies.

IEEE Std 802[®]-2014, IEEE Standard for Local and Metropolitan Area Networks: Overview and Architecture.^{2,3}

IEEE Std 1541TM-2002 (Reaff 2008), IEEE Standard for Prefixes for Binary Multiples.

ISO/IEC 80000-13:2008, Quantities and units — Part 13: Information science and technology.^{4, 5}

ITU-T Rec. X.667 (Sept. 2004), Information technology — Open Systems Interconnection — Procedures for the operation of OSI Registration Authorities: Generation and registration of universally unique identifiers (UUIDs) and their use as ASN.1 object identifier components.⁶

koniec náhľadu – text ďalej pokračuje v platenej verzii STN

² IEEE publications are available from The Institute of Electrical and Electronics Engineers (<u>http://standards.ieee.org</u>).

³ The IEEE standards or products referred to in this clause are trademarks of The Institute of Electrical and Electronics Engineers, Inc. ⁴ This standard cancels and replaces sections 3.8 and 3.9 of IEC 60027-2 (2005).

⁵ ISO/IEC publications are available from the International Organization for Standardization (http://www.iso.ch/). ISO/IEC publications are also available in the United States from Global Engineering Documents (http://global.ihs.com/). Electronic copies are available in the United States from the American National Standards Institute (http://www.ansi.org/).

⁶ ITU-T publications are available from the International Telecommunications Union (http://www.itu.int/).

⁷ The IEEE Standards Dictionary Online subscriptions are available at

 $[\]underline{http://www.ieee.org/portal/innovate/products/standard/standards_dictionary.html}$