INTERNATIONAL STANDARD

ISO/IEEE 11073-10419

Second edition 2019-03

Health informatics — Personal health device communication —

Part 10419:

Device specialization — Insulin pump

Informatique de santé — Communication entre dispositifs de santé personnels —

Partie 10419: Spécialisation des dispositifs — Pompe à insuline





© IEEE 2018

*entation, no part of this cal, including photor ted from IEEE at * All rights reserved. Unless otherwise specified, or required in the context of its implementation, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested from IEEE at the address below.

3 Park Avenue, New York NY 10016-5997, USA

Email: stds.ipr@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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted (see www.iso.org/directives).

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.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see www.iso.org/iso/foreword.html.

ISO/IEEE 11073-10419 was prepared by the IEEE 11073 Standards Committee of the IEEE Engineering in Medicine and Biology Society (as IEEE Std 11073-10419-2017) and drafted in accordance with its editorial rules. It was adopted, under the "fast-track procedure" defined in the Partner Standards Development Organization cooperation agreement between ISO and IEEE, by Technical Committee ISO/TC 215, *Health informatics*.

This second edition cancels and replaces the first edition (ISO 11073-10419:2016), which has been technically revised.

A list of all parts in the ISO 11073 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

Abstract: Within the context of the ISO/IEEE 11073 family of standards for device communication, a normative definition of communication between personal telehealth insulin pump devices and compute engines (e.g., cell phones, personal computers, personal health appliances, set top boxes), in a manner that enables plug-and-play interoperability, is established in this standard. It leverages appropriate portions of existing standards including ISO/IEEE 11073 terminology, information models, application profile standards, and transport standards. It specifies the use of specific term codes, formats, and behaviors in telehealth environments ba, unicatik

J73-10419™,

**Electrica* restricting optionality in base frameworks in favor of interoperability. The standard defines a common core of communication functionality for personal telehealth insulin pump devices.

Keywords: IEEE 11073-10419™, insulin pump, medical device communication, personal health devices

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

Copyright @ 2018 by The Institute of Electrical and Electronics Engineers, Inc. All rights reserved. Published 25 January 2018. Printed in the United States of America.

IEEE is a registered trademark in the U.S. Patent & Trademark Office, owned by The Institute of Electrical and Electronics Engineers, Incorporated.

ISBN 978-1-5044-4291-6 STD22758 ISBN 978-1-5044-4292-3 STDPD22758 Print:

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 Notices and Disclaimers Concerning IEEE Standards Documents." They can also be obtained on request from IEEE or viewed at http://standards.ieee.org/IPR/disclaimers.html.

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. IEEE Standards are documents developed through scientific, academic, and industry-based technical working groups. Volunteers in IEEE working groups 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 Standards do not guarantee or ensure safety, security, health, or environmental protection, or ensure against interference with or from other devices or networks. Implementers and users 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.

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 upon 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 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 Xplore at http://ieeexplore.ieee.org/ 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, Chair Michael J. Kirwan, Chair Melanie S. Yeung, Vice Chair Akib Uddin, Vice Chair

Karsten Aalders Charles R. Abbruscato Nabil Abujbara Maher Abuzaid James Agnew Haidar Ahmad Manfred Aigner Jorge Alberola Murtaza Ali Rolf Ambuehl David Aparisi Paolo Ariano 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

Rudy Belliardi Kathryn M. Bennett Daniel Bernstein George A. Bertos Chris Biernacki Ola Björsne Thomas Blackadar Marc Blanchet Thomas Bluethner Douglas P. Bogia Xavier Boniface Shannon Boucousis Julius Broma Lyle G. Bullock Bernard Burg

Chris Burns

Anthony Butt

Satya Calloji

Xiaoying Cao

Carole C. Carey

Jeremy Byford-Rew

David Bean

John Bell

Craig Carlson Santiago Carot-Nemesio Randy W. Carroll Simon Carter

Seungchul Chae Rahul Chauhan James Cheng Peggy Chien David Chiu Jinyong Choi 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

Ndifor Cyril Fru

Xiaolian Duan

Brian Dubreuil

Allen Curtis

Jesús Daniel Trigo
Eyal Dassau
David Davenport
Russell Davis
Sushil K. Deka
Ciro de la Vega
Pedro de-las-Heras-Quiros
Jim DelloStritto
Matthew d'Entremont
Kent Dicks
Hyoungho Do
Alistair Donaldson

Souray Dutta Jakob Ehrensvard Fredrik Einberg Michihiro Enokida Javier Escayola Calvo Mark Estes Leonardo Estevez Roger Feeley Hailing Feng Bosco T. Fernandes Christoph Fischer Morten Flintrup Joseph W. Forler Russell Foster Eric Freudenthal Matthias Frohner

Ken Fuchs
Jing Gao
Xuemei 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 Aki Harma Jordan Hartmann Kai Hassing Marc Daniel Haunschild

Wolfgang Heck Nathaniel Heintzman Charles Henderson Jun-Ho Her Helen B. Hernandez Takashi Hibino Timothy L. Hirou Allen Hobbs Alex Holland Arto Holopainen Kris Holtzclaw Robert Hoy Frank Hsu Anne Huang Sen-Der Huang Zhiqiang Huang Ron Huby David Hughes Robert D. Hughes Jiyoung Huh Hugh Hunter Hitoshi Ikeda Yutaka Ikeda

Mark G. Schnell

Philip O. Isaacson Jinsei Miyazaki Atsushi Ito Erik Moll Michael Jaffe Darr Moore Praduman Jain Carsten Mueglitz Wei Jin Piotr Murawski Danny Jochelson Soundharya Nagasubramanian Phaneeth Junga Jae-Wook Nah Akiyoshi Kabe Alex Neefus Steve Kahle Trong-Nghia Nguyen-Dobinsky Tomio Kamioka Michael E. Nidd Tetsu Nishimura Kei Kariya Andy Kaschl Jim Niswander Junzo Kashihara Hiroaki Niwamoto Kohichi Kashiwagi Thomas Norgall Ralph Kent Anand Noubade Laurie M. Kermes Yoshiteru Nozoe Ikuo Keshi Abraham Ofek Junhyung Kim Brett Olive Minho Kim Begonya Otal Min-Joon Kim Marco Paleari Taekon Kim Charles Palmer Tetsuya Kimura Bud Panjwani Alfred Kloos Carl Pantiskas Jeongmee Koh Harry P. Pappas Jean-Marc Koller Hanna Park John Koon Jong-Tae Park Patty Krantz Myungeun Park Raymond Krasinski Soojun Park Phillip E. Pash Alexander Kraus TongBi Pei Ramesh Krishna Geoffrey Kruse Lucian Pestritu Falko Kuester Soren Petersen Rafael Lajara James Petisce Pierre Landau Peter Piction Michael Pliskin Jaechul Lee JongMuk Lee Varshney Prabodh Jeff Price Kyong Ho Lee Rami Lee Harald Prinzhorn Sungkee Lee Harry Qiu Arif Rahman Woojae Lee Yonghee Lee Tanzilur Rahman Joe Lenart Steve Ray Phillip Raymond Kathryn A. Lesh Catherine Li Tim Reilly Barry Reinhold Qiong Li Brian Reinhold Patrick Lichter Jisoon Lim Melvin I. Reynolds Joon-Ho Lim John G. Rhoads Jeffrey S. Robbins John Lin Chris Roberts Wei-Jung Lo Charles Lowe Moskowitz Robert

Bob MacWilliams Fatemeh Saki
Srikkanth Madhurbootheswaran Bill Saltzstein
Miriam L. Makhlouf Benedikt Salzbrunn
Romain Marmot Giovanna Sannino
Sandra Martinez Jose A. Santos-Cadenas

Don Ludolph

Christian Luszick

Sandra Martinez

Miguel Martínez de Espronceda
Cámara

Peter Mayhew
Jim McCain
László Meleg
Alexander Mense

Jose A. Santos-Cac
Stefan Sauermann
John Sawyer
Guillaume Schatz
Alois Schloegl
Paul S. Schluter
Lars Schmitt

Richard A. Schrenker Antonio Scorpiniti Kwang Seok Seo Riccardo Serafin Sid Shaw Frank 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 Lars Steubesand John (Ivo) Stivoric Raymond A. Strickland Chandrasekaran Subramaniam

Hermanni Suominen Lee Surprenant Ravi Swami Ray Sweidan Jin Tan Yi Tang Haruyuyki Tatsumi John W. Thomas

Jonas Tirén Alexandra Todiruta Janet Traub Gary Tschautscher Masato Tsuchid Ken Tubman Yoshihiro Uchida Sunil Unadkat Fabio Urbani Philipp Urbauer Laura Vanzago Alpo Värri Dalimar Velez Rudi Voon Barry Vornbrock 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

Dan Xiao

Yaxi Yan

Timothy Robertson

David Rosales

Done-Sik Yoo Zhiqiang Zhang Qifeng Yan Junjie Yang Jianchao Zeng Thomas Zhao Ricky Yang Jason Zhang Miha Zoubek Szymon Zyskoter Qiang Yin

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

John Ballingall Noriyuki Ikeuchi Melvin I. Reynolds Hector Barron Gonzalez Bartien Sayogo Atsushi Ito Lyle G. Bullock Piotr Karocki Lars Schmitt Keith Chow Patrick Keith-Hynes Raymond Strickland Joseph El Youssef Walter Struppler Patrick Kinney Randall Groves Robert Kircher Jan Wittenber Michael J. Kirwan Oren Yuen Kai Hassing Nick S. A. Nikjoo Werner Hoelzl Daidi Zhong Henry Pinto

When the IEEE-SA Standards Board approved this standard on 28 September 2017, it had the following membership:

> Jean-Philippe Faure, Chair Gary Hoffman, Vice Chair John D. Kulick, Past Chair Konstantinos Karachalios, Secretary

Chuck Adams Thomas Koshy Robby Robson Masayuki Ariyoshi Joseph L. Koepfinger* Dorothy Stanley Ted Burse Kevin Lu Adrian Stephens Stephen Dukes Daleep Mohla Mehmet Ulema Doug Edwards Damir Novosel Phil Wennblom J. Travis Griffith Ronald C. Petersen Howard Wolfman Annette D. Reilly Michael Janezic

^{*}Member Emeritus

Introduction

This introduction is not part of IEEE Std 11073-10419-2017, Health informatics-Personal health device communication—Part 10419: Device Specialization—Insulin Pump.

ds en e optim.
communic.
ally focused sa
ces (PHDs). ISO/IEEE 11073 standards enable communication between medical devices and external computer systems. This document uses the optimized framework created in ISO/IEEE 11073-20601:2016 and describes a specific, interoperable communication approach for insulin pumps. These standards align with, and draw on, the existing clinically focused standards to provide support for communication of data from clinical or personal health devices (PHDs).

¹Information on references can be found in Clause 2.

Contents

1. Overview	12
1.1 Scope	
1.2 Purpose	
1.3 Context	
1.5 Context	12
2. Normative references	13
3. Definitions, acronyms, and abbreviations	12
3.1 Definitions	13
3.2 Acronyms and abbreviations	
3.2 Actoryths and aboreviations	13
4. Introduction to ISO/IEEE 11073 personal health devices (PHDs)	16
4.1 General	
4.2 Introduction to ISO/IEEE 11073-20601 modeling constructs	
4.3 Compliance with other standards	
5. Insulin pump device concepts and modalities	
5.1 General	
5.2 Device types	
5.3 Collected data	
5.4 Stored data	
5.5 Scheduled data	24
6 Insulin nume domain information model (DIM)	24
6. Insulin pump domain information model (DIM)	24
6.2 Class extensions	24
6.3 Object instance diagram	24
6.4 Types of configuration	25
6.5 Profiles	
6.6 MDS object	
6.7 Numeric objects	
6.8 Real-time sample array objects	
6.9 Enumeration objects	43
6.10 PM-store objects	
6.11 Schedule-store objects	
6.12 Scanner objects	
6.13 Class extension objects	
6.14 Insulin pump information model extensibility rules	
0.14 Insum pump information model extensionity rules	
7. Insulin pump service model	65
7.1 General	
7.2 Object access services	
7.3 Object access event report services	
·	
8. Insulin pump communication model	69
8.1 Overview	
8.2 Communications characteristics	
8.3 Association procedure	
8.4 Configuring procedure	71
8.5 Operating procedure	
8.6 Time synchronization	

9. Test associations	
9.1 Behavior with standard configuration	
9.2 Behavior with extended configurations	 74
10. Conformance	
10.1 Applicability	
10.2 Conformance specification	 75
10.3 Levels of conformance	
10.4 Implementation conformance statements (ICSs)	76
Annex A (informative) Bibliography	 81
Annex B (normative) Any additional ASN.1 definitions	 82
B.1 Device status and insulin pump status bit mapping	
B.2 Capability-mask	
B.3 State-flag	
Annex C (normative) Allocation of identifiers	86
10.5 General	
10.6 Definitions of terms and codes	
10.7 Systematic derivations of terms and codes	
Annex D (informative) Message sequence examples	97
· militi 2 (militimus v) militiago soquente manipale	, ,
Annex E (normative) Schedule-store class	90
E.1 Schedule-store class	
E.2 Schedule-segment class	
212 20110010 216110110 11122	
Annex F (normative) Schedule class ASN.1 definitions	107
F.1 ACTION-method-related data types	
F.2 Data types for new object attributes and object services	
F.3 Data protocol definitions	
Annex G (informative) The schedule-store concept	
G.1 General	
G.2 Schedule-store object hierarchy	
Annex H (informative) Scedule communication model	115
H.1 Operating procedure	115
Annex I (informative) Protocol data unit (PDU) examples	119
I.1 General	
I.2 Association information exchange	
I.3 Configuration information exchange	
I.4 GET MDS attributes service	
I.5 Data reporting	
I.6 Disassociation.	
2.0 2.2.2.2.000	120
Annex I (informative) Revision history	129

Health informatics—Personal health device communication

Part 10419: Device Specialization—Insulin Pump

1. Overview

1.1 Scope

This standard establishes a normative definition of communication between personal telehealth insulin pump devices (agents) and managers (e.g., cell phones, personal computers, personal health appliances, set top boxes) in a manner that enables plug-and-play interoperability. It leverages work done in other ISO/IEEE 11073 standards including existing terminology, information profiles, application profile standards, and transport standards. It specifies the use of specific term codes, formats, and behaviors in telehealth environments, restricting optionality in base frameworks in favor of interoperability. This standard defines a common core functionality of personal telehealth insulin pump devices.

In the context of personal health devices (PHDs), an insulin pump is a medical device used for the administration of insulin in the treatment of diabetes mellitus, also known as continuous subcutaneous insulin infusion (CSII) therapy.

This standard provides the data modeling according to ISO/IEEE 11073-20601 and does not specify the measurement method.

1.2 Purpose

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

1.3 Context

See ISO/IEEE 11073-20601:2016 for an overview of the environment within which this standard is written.

² Information on references can be found in Clause 2.

IEEE Std 11073-10419-2017

Health informatics—Personal health device communication—Part 10419: Device Specialization—Insulin Pump

This standard defines the device specialization for the insulin pump, being a specific agent type, and provides a description of the device concepts, its capabilities, and its implementation according to this standard.

This standard is based on ISO/IEEE 11073-20601:2016, which in turn draw information from both ISO/IEEE 11073-10201:2004 [B8] and ISO/IEEE 11073-20101:2004 [B9].³ The medical device encoding rules (MDER) used within this standard are fully described in ISO/IEEE 11073-20601:2016.

This standard reproduces relevant portions of the nomenclature found in ISO/IEEE 11073-10101:2004 [B6] and ISO/IEEE 11073-10101a:2015 [B7] and adds new nomenclature codes for the purposes of this standard. Among these standards and ISO/IEEE 11073-20601:2016, all required nomenclature codes for implementation are documented.

NOTE—In this standard, ISO/IEEE 11073-104zz is used to refer to the collection of device specialization standards that utilize ISO/IEEE 11073-20601, where zz can be any number from 01 to 99, inclusive.⁴

2. Normative references

The following referenced documents are indispensable for the application of this document (i.e., they must be understood and used; therefore, each referenced document is cited in text and its relationship to this document 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.

ISO/IEEE 11073-20601:2016, Health informatics—Personal health device communication—Part 20601: Application Profile—Optimized Exchange Protocol.⁵

See Annex A for all informative material referenced by this standard.

3. Definitions, acronyms, and abbreviations

3.1 Definitions

For the purposes of this document, the following terms and definitions apply. The *IEEE Standards Dictionary Online* should be consulted for terms not defined in this clause.⁶

agent: A node that collects and transmits personal health data to an associated manager.

artificial pancreas: A system combining diabetes devices to provide similar functionality as a pancreas. Examples include linking a continuous glucose monitor to an insulin pump to automatically reduce or increase insulin infusion based upon specified thresholds of measured interstitial glucose.

basal insulin: Insulin required to cover the basic insulin needs of the body.

basal rate: Rate of continuously delivered insulin to cover the basic insulin needs of the body

³ The numbers in brackets correspond to the numbers of the bibliography in Annex A.

⁴ Notes in text, tables, and figures of a standard are given for information only and do not contain requirements needed to implement the standard.

⁵ ISO/IEEE publications are available from the ISO Central Secretariat (http://www.iso.ch/). ISO/IEEE publications are also available in the United States from The Institute of Electrical and Electronics Engineers (http://standards.ieee.org/).

⁶ IEEE Standards Dictionary Online is available at http://dictionary.ieee.org.