



SLOVENSKI STANDARD SIST EN ISO 21549-5:2016

01-september-2016

Nadomešča:
SIST EN ISO 21549-5:2008

Zdravstvena informatika - Podatki o pacientu na zdravstveni kartici - 5. del: Identifikacijski podatki (ISO 21549-5:2015)

Health informatics - Patient healthcard data - Part 5: Identification data (ISO 21549-5:2015)

Medizinische Informatik – Patientendaten auf Karten im Gesundheitswesen – Teil 5:
Identifikationsdaten (ISO 21549-5:2015)

Informatique de santé - Données relatives aux cartes de santé des patients - Partie 5:
Données d'identification (ISO 21549-5:2015)

Ta slovenski standard je istoveten z: **EN ISO 21549-5:2016**

ICS:

35.240.15	Identifikacijske kartice. Čipne kartice. Biometrija	Identification cards. Chip cards. Biometrics
35.240.80	Uporabniške rešitve IT v zdravstveni tehniki	IT applications in health care technology

SIST EN ISO 21549-5:2016 **en,fr,de**

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EUROPEAN STANDARD

EN ISO 21549-5

NORME EUROPÉENNE

EUROPÄISCHE NORM

April 2016

ICS 35.240.80

Supersedes EN ISO 21549-5:2008

English Version

Health informatics - Patient healthcard data - Part 5: Identification data (ISO 21549-5:2015)

Informatique de santé - Données relatives aux cartes
de santé des patients - Partie 5: Données
d'identification (ISO 21549-5:2015)

Medizinische Informatik - Patientendaten auf Karten
im Gesundheitswesen - Teil 5: Identifikationsdaten
(ISO 21549-5:2015)

This European Standard was approved by CEN on 2 April 2016.

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COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

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

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European foreword

The text of ISO 21549-5:2015 has been prepared by Technical Committee ISO/TC 215 “Health informatics” of the International Organization for Standardization (ISO) and has been taken over as EN ISO 21549-5:2016 by 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of an identical text or by endorsement, at the latest by October 2016, and conflicting national standards shall be withdrawn at the latest by October 2016.

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 21549-5:2008.

This document has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association.

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.

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Endorsement notice

The text of ISO 21549-5:2015 has been approved by CEN as EN ISO 21549-5:2016 without any modification.

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INTERNATIONAL
STANDARD

ISO
21549-5

Second edition
2015-09-15

**Health informatics — Patient
healthcard data —**

**Part 5:
Identification data**

*Informatique de santé — Données relatives aux cartes de santé des
patients —*

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Partie 5: Données d'identification
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ISO copyright office
Ch. de Blandonnet 8 • CP 401
CH-1214 Vernier, Geneva, Switzerland
Tel. +41 22 749 01 11
Fax +41 22 749 09 47
copyright@iso.org
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ISO 21549-5:2015(E)

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. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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 on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT) see the following URL: [Foreword - Supplementary information \(standards.iteh.ai\)](#)

The committee responsible for this document is ISO/TC 215, *Health Informatics*.

This second edition cancels and replaces the first edition (ISO 21549-5:2008), which has undergone a minor revision. The following changes have been made:

- [Subclause 5.2](#), Table 1: condition of Sex optionality is added.
- [Subclause 5.2](#), Table 1: optionality of National representation of the name is corrected to match ASN.1 definition and [Figure 1](#).

ISO 21549 consists of the following parts, under the general title *Health informatics — Patient healthcard data*:

- *Part 1: General structure*
- *Part 2: Common objects*
- *Part 3: Limited clinical data*
- *Part 4: Extended clinical data*
- *Part 5: Identification data*
- *Part 6: Administrative data*
- *Part 7: Medication data*
- *Part 8: Links*

Introduction

With a more mobile population, greater healthcare delivery in the community and at patients' homes, together with a growing demand for improved quality of ambulatory care, portable information systems and stores have increasingly been developed and used. Such devices are used for tasks ranging from identification, through portable medical record files, and on to patient-transportable monitoring systems.

The functions of such devices are to carry and to transmit person-identifiable information between themselves and other systems; therefore, during their operational lifetime they may share information with many technologically different systems which differ greatly in their functions and capabilities.

Healthcare administration increasingly relies upon similar automated identification systems. For instance prescriptions may be automated and data exchange carried out at a number of sites using patient transportable computer readable devices. Healthcare funding institutions and providers are increasingly involved in cross-region care, where reimbursement may require automated data exchange between dissimilar healthcare systems. Administrative data objects may require linkage to external parties responsible for their own domains which are not within the scope of this part of ISO 21549. For instance, cross-border reimbursement of healthcare services are usually regulated by law and intergovernmental agreements which are not subject to standardization.

The advent of remotely accessible databases and support systems has led to the development and use of "Healthcare Person" identification devices that are also able to perform security functions and transmit digital signatures to remote systems via networks.

With the growing use of data cards for practical everyday healthcare delivery, the need has arisen for a standardized data format for interchange.

The person-related data carried by a data card can be categorised in three broad types: identification (of the device itself and the individual to whom the data it carries relates), administrative and clinical. It is important to realize that a given healthcare data card "de facto" contains device data and identification data and may in addition contain administrative, clinical, medication and linkage data.

Device data are defined to include:

- identification of the device itself;
- identification of the functions and functioning capabilities of the device.

Identification data are defined to include:

- unique identification of the device holder (and not information of other persons).

Administrative data can include:

- complementary person(s) related data;
- identification of the funding of healthcare, whether public or private, and their relationships, i.e. insurer(s), contract(s) and policy(ies) or types of benefits;
- identification of other persons as a part of the insurance contract (e.g. a family contract);
- other data (distinguishable from clinical data) that are necessary for the purpose of healthcare delivery.

Clinical data may include:

- items that provide information about health and health events;
- their appraisal and labelling by a healthcare provider;
- related actions planned requested or performed.