



Technical Report

ISO/TR 4419

Health informatics — Pathways for human-computer interaction in electronic health information record systems to reduce clinician burden

Informatique de santé — Voies d'interaction homme-machine dans les systèmes électroniques d'enregistrement des informations de santé afin de réduire la charge de travail des cliniciens

**First edition
2026-01**

ITEH Standards
(<https://standards.iteh.ai>)

Table of Content Preview

[ISO/TR 4419:2026](#)

<https://standards.iteh.ai/catalog/standards/iso/54376627-b1dd-4b5f-bfce-2e048366c476/iso-tr-4419-2026>

iTeh Standards
(<https://standards.iteh.ai>)
Document Preview

[ISO/TR 4419:2026](https://standards.iteh.ai/catalog/standards/iso/54376627-b1dd-4b5f-bfce-2e048366c476/iso-tr-4419-2026)

<https://standards.iteh.ai/catalog/standards/iso/54376627-b1dd-4b5f-bfce-2e048366c476/iso-tr-4419-2026>



COPYRIGHT PROTECTED DOCUMENT

© ISO 2026

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 either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
CP 401 • Ch. de Blandonnet 8
CH-1214 Vernier, Geneva
Phone: +41 22 749 01 11
Email: copyright@iso.org
Website: www.iso.org

Published in Switzerland

Contents

	Page
Foreword	iv
Introduction	v
1 Scope	1
2 Normative references	1
3 Terms and definitions	1
4 Abbreviated terms	4
5 Clinician burden	5
5.1 Health information record systems' unfulfilled promises	5
5.2 Health information record systems	6
5.3 Health modules and decision support systems	6
5.4 Patient portals	8
5.5 Remote patient monitoring (RPM) and wearable devices	8
5.6 Ontology-based systems	8
5.7 Knowledge repositories	9
6 Burden mitigation	9
6.1 Approaches to reducing clinician burden	9
6.2 Human-computer interaction	10
6.3 HCI benefits	12
7 Summary	13
Bibliography	15

iTeh Standards

(<https://standards.iteh.ai>)
Document Preview

[ISO/TR 4419:2026](https://standards.iteh.ai)

<https://standards.iteh.ai/catalog/standards/iso/54376627-b1dd-4b5f-bfce-2e048366c476/iso-tr-4419-2026>

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).

ISO draws attention to the possibility that the implementation of this document may involve the use of (a) patent(s). ISO takes no position concerning the evidence, validity or applicability of any claimed patent rights in respect thereof. As of the date of publication of this document, ISO had not received notice of (a) patent(s) which may be required to implement this document. However, implementers are cautioned that this may not represent the latest information, which may be obtained from the patent database available at www.iso.org/patents. ISO shall not be held responsible for identifying any or all such patent rights.

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.

This document was prepared by Technical Committee ISO/TC 215, *Health informatics*, in collaboration with HL7.

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.

ISO/TR 4419:2026

<https://standards.iteh.ai/catalog/standards/iso/54376627-b1dd-4b5f-bfce-2e048366c476/iso-tr-4419-2026>

Introduction

The implementation of electronic health information record systems (HIRS), defined in [Clause 3](#), is one of the primary factors contributing both positive and negative impacts on clinical practice and patient experience. HIRS can be used to create complex hierarchical structures from basic primitive types. HIRS has substantially altered the norms of clinician-patient interaction, often unintentionally diminishing the most meaningful aspects of healthcare practice for the clinician and the patient.[\[1\]](#),[\[2\]](#),[\[3\]](#),[\[4\]](#),[\[5\]](#) Among the various segments of the healthcare industry, there is widespread agreement that the structure, capabilities and operations within HIRS are not working for clinicians, patients, health organizations, health technology vendors or the businesses who enable the provision of health insurance, equipment, biotechnology services, research, and pharmaceuticals. The healthcare industry faces a diverse range of challenges causing numerous serious issues for each of these groups. Challenges include rising numbers of preventable mistakes,[\[6\]](#) declining quality of care, sub-optimal health outcomes, out-of-control costs, financial pressure, resource constraints, clinician burnout, and problems integrating technology with artificial intelligence (AI) and telehealth. These ever-changing technical, regulatory, and environmental challenges, along with cybersecurity threats to patient data requiring robust security measures and constant vigilance, demand significant additional effort, planning and resources. These challenges impact operational efficiency and can lead to workforce shortages. There is a growing gap in the availability of skilled healthcare professionals and healthcare services relative to growing demand for healthcare services as populations age and the prevalence of chronic diseases rises.

The root causes of many of these challenges relate to deeper issues with HIRS, including:

- inadequate standards for data quality and insufficient auditing and enforcement of existing data quality standards;
- complex nonintuitive data structures and functionalities;
- lack of functionalities to mitigate cognitive overload;
- fragmented care, poor workflow integration, and lack of context and specialty specific support.[\[7\]](#)

While the problems are daunting, they appear to have motivated many healthcare and non-healthcare actors to start to identify solutions. This document reviews a large body of clinician opinion and research findings suggesting that the HIRS status quo cannot continue and that comprehensive HIRS reform is necessary, possibly leading to a completely new HIRS which would be termed the digital healthcare system (DHS). During the transition to a new HIRS, clinicians need to continue to treat patients with the available HIRS; the idea of moving to a more perfect DHS is an evolutionary process over a period of years. The goal now is to use the knowledge, principles, standards and experience gained from the first 15 years of wide scale HIRS implementation, and the advances made in several domains to support continuing efforts to improve HIRS and move toward that more perfect DHS.

While the burden of disease is a known epidemiologic concept, clinician burden is a less known and more recent term.[\[8\]](#) Clinician burden occurs when the clinician's environment and workload impose physical, cognitive, psychological, and time burdens on clinicians without sufficiently improving quality of care and clinician functioning.[\[2\]](#) Multiple factors can contribute to clinician burden, including

- time and productivity pressures,
- excessive bureaucratic tasks of low clinical value,
- limited capacities of human cognition versus high demands of information-intensive clinical practice,[\[9\]](#),[\[10\]](#), and
- limited clinician-centred design in health information record systems and clinical decision support tools, which contributes to ineffective functionality and increased burden on clinical workflows.

While contributing to improved healthcare, the volume and velocity of new patient-related healthcare information, patient-specific data, and underlying biomedical knowledge also unintentionally increase clinician burden.[\[11\]](#) A large and growing body of research suggests that poor HIRS usability and poor integration within clinician workflows are important factors preventing electronic health information