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<text> **Electronic Health Record-System Functional Model, Release 1.1** 



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# **EUROPEAN STANDARD** NORME EUROPÉENNE **EUROPÄISCHE NORM**

# **EN ISO 10781**

November 2009

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**English Version** 

## Electronic Health Record-System Functional Model, Release 1.1 (ISO 10781:2009)

Modèle fonctionnel d'un système d'enregistrement électronique de la santé HL7, version 1.1 (ISO 10781:2009)

Medizinische Informatik - HL7 funktionales Modell für ein elektronisches Gesundheitsaktensystem (ISO 10781:2009)

This European Standard was approved by CEN on 7 November 2009.

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

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Ref. No. EN ISO 10781:2009: E

## Foreword

This document (EN ISO 10781:2009) 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of an identical text or by endorsement, at the latest by May 2010, and conflicting national standards shall be withdrawn at the latest by May 2010.

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.

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### **Endorsement notice**

em. , y CEN as. The text of ISO 10781:2009 has been approved by CEN as a EN ISO 10781:2009 without any modification.

# **HL7 EHR Work Group**

# Electronic Health Record-System **Functional Model, Release 1.1 June 2009**

# Chapter One: Overview

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HL7 EHR System Functional Model

## Preface

## i. Notes to Readers

The HL7 EHR-S Functional Model, which was approved in July, 2004 as a Draft Standard for Trial Use (DSTU), has undergone a series of enhancements in the last year as it made its way to a fully approved American National Standards Institute (ANSI) standard. A broad constituency - including intensive outreach to industry, care providers, and healthcare organizations - has worked to refine the initial EHR-S Functional Model. This version reflects the changes made as part of the reconciliation process in the successful membership level balloting that took place at the January 2007 HL7 Workgroup Meeting.

## ii. Acknowledgements

The committee is indebted to the following past co-chairs and facilitators for their contributions towards the Public Health domain and the material presented here. Direct and indirect participants in the development of the model, including workgroup contributors and other participants, can be found in the "Contributor Listing" found at <u>www.HL7.org/EHR</u> in the "documents" section.

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Chapter 1: Overview

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## iii. Changes from Previous Release

The HL7 EHR-S Functional Model was promoted to an ISO International Standard after passing HL7 consensus ballot as a normative standard and achieving ANSI approval. This promotion follows the process outlined in the ISO/HL7 Pilot Agreement. The June, 2009 release of this document includes updates from the joint ISO/HL7 ballot. The dates of the comment for the 60 day Draft International Standard (DIS) ballot were February through April, 2009. The HL7 comment period was a 30 day window that coordinted with the last 30 days of the ISO-DIS ballot period. Reconciliation was accomplished in May and early June of 2009.

HL7 EHR System Functional Model

# **Chapter 1 Introduction and Overview**

The HL7 Electronic Health Records Special Interest Group (EHR SIG) was established in the spring of 2002. In the spring of 2003 the HL7 group began efforts to develop a standardized functional specification for Electronic Health Records Systems (EHR-S). In May 2004 the SIG was promoted to a full HL7 Technical Committee, becoming the EHR TC. The EHR TC is intended primarily to serve as a body which promotes the uptake of Electronic Health Record (EHR) implementation by standardizing the functions that may be present, based on user selection, in an EHR-S.

The Department of Health and Human Services, the Veterans Health Administration, the Health Information Management Systems Society and the Robert Wood Johnson Foundation, in a public-private partnership, approached HL7 to accelerate their existing work to develop a consensus standard to define the functions of an EHR-S. HL7, through its EHR SIG, responded by developing an EHR-S Functional Model that passed ballot as a Draft Standard for Trial Use (DSTU) in April 2004. The Functional Model DSTU was published and formally registered with the American National Standards Institute (ANSI) in July 2004. The Functional Model was then balloted and passed as a normative standard as part of the January 2007 HL7 Workgroup Meeting and is now registered as a normative standard with ANSI

Learning important lessons from the ballot process, a Functional Model with a clearer, more simplified list of functions, has been created. The HL7 EHR System Functional Model provides a reference list of functions that may be present in an Electronic Health Record System (EHR-S). The function list is described from a user perspective with the intent to enable consistent expression of system functionality. This EHR-S Model, through the creation of Functional Profiles, enables a standardized description and common understanding of functions sought or available in a given setting (e.g. intensive care, cardiology, office practice in one country or primary care in another country).

## 1 Background

## 1.1 What is HL7?

Established in 1987, Health Level Seven (HL7) is an American National Standards Institute (ANSI) accredited, not-for-profit standards-development organization, whose mission is to provide standards for the exchange, integration, sharing, and retrieval of electronic health information; support clinical practice; and support the management, delivery and evaluation of health services. ANSI accreditation, coupled with HL7's own procedures, dictates that any standard published by HL7 and submitted to ANSI for approval, be developed and ratified by a process that adheres to ANSI's procedures for open consensus and meets a balance of interest requirement by attaining near equal participation in the voting process by the various constituencies that are materially affected by the standard (e.g., vendors, providers, government agencies, consultants, non-profit organizations). This balance of interest goal ensures that a particular constituency is neither refused participation nor is it allowed to dominate the development and ratification of a proposed standard. More information and background on ANSI is available on their website at: http://www.ANSI.org