SLOVENSKI STANDARD
kSIST FprEN ISO 22870:2016

01-september-2016

Preskušanje ob preiskovancu (POCT) - Zahteve za kakovost in kompetentnost (ISO/FDIS 22870:2016)

Point-of-care testing (POCT) - Requirements for quality and competence (ISO/FDIS 22870:2016)

Patientennahe Untersuchungen (point-of-care testing, POCT) - Anforderungen an Qualität und Kompetenz (ISO/FDIS 22870:2016)

Analyse de biologie délocalisée (ADBD) - Exigences en matière de qualité et de compétences (ISO/FDIS 22870:2016)

Ta slovenski standard je istoveten z: **FprEN ISO 22870**

ICS:

03.120.10	Vodenje in zagotavljanje kakovosti	Quality management and quality assurance
11.100.01	Laboratorijska medicina na splošno	Laboratory medicine in general

kSIST FprEN ISO 22870:2016 **en**

FINAL
DRAFT

INTERNATIONAL
STANDARD

**ISO/FDIS
22870**

ISO/TC **212**

Secretariat: ANSI

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Point-of-care testing (POCT) — Requirements for quality and competence

Analyses de biologie délocalisées (ADBD) — Exigences concernant la qualité et la compétence

Please see the administrative notes on page ii

Reference number
ISO/FDIS 22870:2016(E)

ISO/CEN PARALLEL PROCESSING

This final draft has been developed within the International Organization for Standardization (ISO), and processed under the **ISO-lead** mode of collaboration as defined in the Vienna Agreement. The final draft was established on the basis of comments received during a parallel enquiry on the draft.

This final draft is hereby submitted to the ISO member bodies and to the CEN member bodies for a parallel two-month approval vote in ISO and formal vote in CEN.

Positive votes shall not be accompanied by comments.

Negative votes shall be accompanied by the relevant technical reasons.

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Foreword

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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).

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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 World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see the following URL: www.iso.org/iso/foreword.html.

The committee responsible for this document is ISO/TC 212, *Clinical laboratory testing and in vitro diagnostic test systems*.

This second edition cancels and replaces the first edition (ISO 22870:2006), which has been technically revised that includes cross-references to the applicable clauses in ISO 15189:2012.

