

SLOVENSKI STANDARD
SIST-TS CEN ISO/TS 22367:2010
01-junij-2010

Medicinski laboratoriji - Zmanjšanje števila napak z obvladovanjem tveganja in stalnim izboljševanjem (ISO/TS 22367:2008, vključno s popravkom Cor 1:2009)

Medical laboratories - Reduction of error through risk management and continual improvement (ISO/TS 22367:2008, including Cor 1:2009)

Medizinische Laboratorien - Fehlerrückführung durch Risikomanagement und ständige Verbesserung (ISO/TS 22367:2008, einschließlich Cor 1:2009)

Laboratoires médicaux - Réduction d'erreurs par gestion du risque et amélioration continue (ISO/TS 22367:2008, Cor 1:2009 inclus)

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Ta slovenski standard je istoveten z: CEN ISO/TS 22367:2010

ICS:

03.100.01	Organizacija in vodenje podjetja na splošno	Company organization and management in general
11.100.01	Laboratorijska medicina na splošno	Laboratory medicine in general

SIST-TS CEN ISO/TS 22367:2010 **en**

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TECHNICAL SPECIFICATION
SPÉCIFICATION TECHNIQUE
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CEN ISO/TS 22367

January 2010

ICS 11.100.01

English Version

**Medical laboratories - Reduction of error through risk
management and continual improvement (ISO/TS 22367:2008,
including Cor 1:2009)**

Laboratoires médicaux - Réduction d'erreurs par gestion du
risque et amélioration continue (ISO/TS 22367:2008, Cor
1:2009 inclus)

Medizinische Laboratorien - Fehlerrückführung durch
Risikomanagement und ständige Verbesserung (ISO/TS
22367:2008, einschließlich Cor 1:2009)

This Technical Specification (CEN/TS) was approved by CEN on 4 January 2010 for provisional application.

The period of validity of this CEN/TS is limited initially to three years. After two years the members of CEN will be requested to submit their comments, particularly on the question whether the CEN/TS can be converted into a European Standard.

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Foreword

The text of ISO 22367:2008, including Cor 1:2009 has been prepared by Technical Committee ISO/TC 212 "Clinical laboratory testing and in vitro diagnostic test systems" of the International Organization for Standardization (ISO) and has been taken over as CEN ISO/TS 22367:2010 by Technical Committee CEN/TC 140 "In vitro diagnostic medical devices" the secretariat of which is held by DIN.

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

The text of ISO 22367:2008, including Cor 1:2009 has been approved by CEN as a CEN ISO/TS 22367:2010 without any modification.

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TECHNICAL SPECIFICATION

ISO/TS 22367

First edition
2008-05-01

Medical laboratories — Reduction of error through risk management and continual improvement

*Laboratoires médicaux — Réduction d'erreurs par gestion du risque et
amélioration continue*

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ISO/TS 22367:2008(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.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

In other circumstances, particularly when there is an urgent market requirement for such documents, a technical committee may decide to publish other types of document:

- an ISO Publicly Available Specification (ISO/PAS) represents an agreement between technical experts in an ISO working group and is accepted for publication if it is approved by more than 50 % of the members of the parent committee casting a vote;
- an ISO Technical Specification (ISO/TS) represents an agreement between the members of a technical committee and is accepted for publication if it is approved by 2/3 of the members of the committee casting a vote.

An ISO/PAS or ISO/TS is reviewed after three years in order to decide whether it will be confirmed for a further three years, revised to become an International Standard, or withdrawn. If the ISO/PAS or ISO/TS is confirmed, it is reviewed again after a further three years, at which time it must either be transformed into an International Standard or be withdrawn.

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.

ISO/TS 22367 was prepared by Technical Committee ISO/TC 212, *Clinical laboratory testing and in vitro diagnostic test systems*.

Introduction

It is a requirement of ISO 15189 that laboratories have an investigative process to identify aspects that do not conform with their own procedures or with predetermined requirements in the quality management system. ISO 15189 specifies that this be linked both to corrective actions and to preventive actions. In addition, it specifies that management review the suitability and effectiveness of the system and its activities in support of patient care, and that they introduce necessary changes. This can best be done by considering potential risks introduced at each step of each process.

Preventive actions are planned and appropriate anticipatory processes, based upon verifiable information, are undertaken to prevent a potential action from occurring. Corrective actions are similarly planned together with appropriate reactive processes; however, these are undertaken to amend identified problems and to avoid their recurrence. Risk management is a planned process that is part of preventive actions and corrective actions.

Preventive actions and corrective actions can be more effectively directed when they are based upon information that is well-organized; classification systems and risk management analysis are two processes that provide well-organized information.

In the context of organizational management, risk has been described as a multidimensional concern about stability and predictability of outcome. Organizational risk involves components that affect the operational, technical, liability and business aspects of the laboratory. In the context of continual improvement, the risk elements of potential for loss are considered with higher priority than the elements of gain. Consideration of risk necessarily includes the linked but different elements of likelihood of occurrence and severity of impact. Factors that impact upon risk can act either directly or indirectly.

The framework of risk management can be described as consisting of the following steps:

- a) planning for risk,
- b) identifying risk and its impacts,
- c) developing risk-handling strategies, and
- d) monitoring for risk control.

These steps are consistent with the management requirements described in ISO 15189, including:

- identifying and controlling non-conformities,
- establishing preventive actions and corrective actions,
- carrying out internal audits and management reviews, and
- implementing continual improvement.

This Technical Specification is intended to provide the first steps to introduce risk management into the structure, organization, operation and quality management system of the medical laboratory.

Classification of laboratory non-conformities, errors and incidents is useful for monitoring purposes and allows the laboratory to determine their criticality, to set priorities in addressing them and to identify underlying causative factors that contribute to errors.

Considerations contained within local, regional and national regulations normally apply.