

Clinical laboratory testing and in vitro diagnostic test systems — Susceptibility testing of infectious agents and evaluation of performance of antimicrobial susceptibility test devices —

Part 2: Evaluation of performance of antimicrobial susceptibility test devices

The European Standard EN ISO 20776-2:2007 has the status of a British Standard

ICS 11.100.20

National foreword

This British Standard is the UK implementation of EN ISO 20776-2:2007.
The UK participation in its preparation was entrusted to Technical Committee CH/212, IVDs.
A list of organizations represented on this committee can be obtained on request to its secretary.
This publication does not purport to include all the necessary provisions of a contract. Users are responsible for its correct application.
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English Version

Clinical laboratory testing and in vitro diagnostic test systems -
Susceptibility testing of infectious agents and evaluation of
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Evaluation of performance of antimicrobial susceptibility test
devices (ISO 20776-2:2007)

Systèmes d'essais en laboratoire et de diagnostic in vitro -
Sensibilité in vitro des agents infectieux et évaluation des
performances des dispositifs pour antibiogrammes - Partie
2: Évaluation des performances des dispositifs pour
antibiogrammes (ISO 20776-2:2007)

Labormedizinische Untersuchungen und In-vitro-
Diagnostika-Systeme - Empfindlichkeitsprüfung von
Infektionserregern und Evaluation von Geräten zur
antimikrobiellen Empfindlichkeitsprüfung - Teil 2: Evaluation
der Leistung einer Vorrichtung zur antimikrobiellen
Empfindlichkeitsprüfung (ISO 20776-2:2007)

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Management Centre: rue de Stassart, 36 B-1050 Brussels

Foreword

This document (EN ISO 20776-2:2007) has been prepared by Technical Committee CEN/TC 140 "In vitro diagnostic medical devices", the secretariat of which is held by DIN, in collaboration with Technical Committee ISO/TC 212 "Clinical laboratory testing and in vitro diagnostic test systems".

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of an identical text or by endorsement, at the latest by January 2008, and conflicting national standards shall be withdrawn at the latest by January 2008.

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INTERNATIONAL STANDARD

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*Systèmes d'essais en laboratoire et de diagnostic in vitro — Sensibilité
in vitro des agents infectieux et évaluation des performances des
dispositifs pour antibiogrammes —*

*Partie 2: Évaluation des performances des dispositifs pour
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