

**Chemical disinfectants  
and antiseptics —  
Quantitative carrier  
test for the evaluation  
of mycobactericidal  
or tuberculocidal  
activity of chemical  
disinfectants used  
for instruments in  
the medical area —  
Test method and  
requirements (phase 2,  
step 2)**

ICS 11.080.20

# National foreword

This British Standard is the UK implementation of EN 14563:2008.

The UK participation in its preparation was entrusted to Technical Committee CH/216, Chemical disinfectants and antiseptics.

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

**Chemical disinfectants and antiseptics - Quantitative carrier test  
for the evaluation of mycobactericidal or tuberculocidal activity of  
chemical disinfectants used for instruments in the medical area -  
Test method and requirements (phase 2, step 2)**

Désinfectants et antiseptiques chimiques - Essai quantitatif  
de porte-germe pour l'évaluation de l'activité  
mycobactéricide ou tuberculocide des désinfectants  
chimiques utilisés pour instruments en médecine humaine -  
Méthode d'essai et prescriptions (phase 2, étape 2)

Chemische Desinfektionsmittel und Antiseptika -  
Quantitativer Keimträgerversuch zur Prüfung der  
mykobakteriziden oder tuberkuloziden Wirkung chemischer  
Desinfektionsmittel für Instrumente im  
humanmedizinischen Bereich - Prüfverfahren und  
Anforderungen (Phase 2, Stufe 2)

This European Standard was approved by CEN on 18 October 2008.

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## Foreword

This document (EN 14563:2008) has been prepared by Technical Committee CEN/TC 216 “Chemical disinfectants and antiseptics”, the secretariat of which is held by AFNOR.

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of an identical text or by endorsement, at the latest by May 2009, and conflicting national standards shall be withdrawn at the latest by May 2009.

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.

This document has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association, and supports essential requirements of EC Directive(s).

For relationship with EU Directive 93/42/EEC, see informative annex ZA, which is an integral part of this document.

Other methods to evaluate the efficacy of chemical disinfectants and antiseptics for different applications in the medical field are in preparation.

A collaborative trial will be undertaken to provide a precision annex to this standard.

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