

**In vitro diagnostikasüsteemid.
Mikrobioloogilisel eesmärgil
laboraatorsetes tingimustes
kasvatatavad mikroobid. Terminid ja
määratlused**

In vitro diagnostic systems - Culture media for
microbiology - Terms and definitions

EESTI STANDARDI EESSÕNA

NATIONAL FOREWORD

<p>Käesolev Eesti standard EVS-EN 1659:1999 sisaldab Euroopa standardi EN 1659:1996 ingliskeelset teksti.</p> <p>Käesolev dokument on jõustatud 12.12.1999 ja selle kohta on avaldatud teade Eesti standardiorganisatsiooni ametlikus väljaandes.</p> <p>Standard on kättesaadav Eesti standardiorganisatsioonist.</p>	<p>This Estonian standard EVS-EN 1659:1999 consists of the English text of the European standard EN 1659:1996.</p> <p>This document is endorsed on 12.12.1999 with the notification being published in the official publication of the Estonian national standardisation organisation.</p> <p>The standard is available from Estonian standardisation organisation.</p>
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<p>Käsitlusala: See Euroopa standard esitab eri klassifikatsioonide terminid laboratoorsetes tingimustes kasvatatavate mikroobide kohta, mida kasutatakse mikrobioloogias (bakterioloogias ja mükoloogias).</p>	<p>Scope:</p>
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ICS 01.040.07, 07.100.10

Võtmesõnad: bioproov, meditsiin, mikrobioloogiline analüüs, sõnastik, teatud tingimustes kasvatatud mikroobid

ICS 01.040.07; 07.100.10

Descriptors: In vitro diagnostic systems, culture media, microbiology, terminology.

English version

In vitro diagnostic systems
Culture media for microbiology
Terms and definitions

Systèmes de diagnostic in vitro – Milieux
de culture de microbiologie – Termes et
définitions

In-vitro-Diagnostik/Diagnostika –
Kulturmedien für die Mikrobiologie:
Begriffe

This European Standard was approved by CEN on 1996-10-19.

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Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the Central Secretariat or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the Central Secretariat has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Netherlands, Norway, Portugal, Spain, Sweden, Switzerland, and the United Kingdom.

CEN

European Committee for Standardization
Comité Européen de Normalisation
Europäisches Komitee für Normung

Central Secretariat: rue de Stassart 36, B-1050 Brussels

Foreword

This European Standard has been prepared by Technical Committee CEN/TC 140 "In vitro diagnostic systems" the secretariat of which is held by DIN.

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 1997, and conflicting national standards shall be withdrawn at the latest by May 1997.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Netherlands, Norway, Portugal, Spain, Sweden, Switzerland and the United Kingdom.

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