

**Kliinilised termomeetrid. Osa 2:
Faasimuundurtüüpi (punktmaatriks)
termomeetrid**

Clinical thermometers - Part 2: Phase change type
(dot matrix) thermometers

EESTI STANDARDI EESSÕNA

NATIONAL FOREWORD

<p>Käesolev Eesti standard EVS-EN 12470-2:2001 sisaldab Euroopa standardi EN 12470-2:2000 ingliskeelset teksti.</p> <p>Käesolev dokument on jõustatud 09.03.2001 ja selle kohta on avaldatud teade Eesti standardiorganisatsiooni ametlikus väljaandes.</p> <p>Standard on kättesaadav Eesti standardiorganisatsioonist.</p>	<p>This Estonian standard EVS-EN 12470-2:2001 consists of the English text of the European standard EN 12470-2:2000.</p> <p>This document is endorsed on 09.03.2001 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:</p> <p>This part of the standard specifies performance requirements and test methods for phase change-type (dot matrix) thermometers for measuring temperature in body cavities.</p> <p>NOTE: A body cavity can be the mouth, rectum or armpit.</p> <p>The standard does not apply to clinical thermometers designed for special applications (e.g. thermometers for hypothermia) which owing to their measurement range, scale interval or maximum permissible error do not meet the requirements specified in this standard.</p>	<p>Scope:</p> <p>This part of the standard specifies performance requirements and test methods for phase change-type (dot matrix) thermometers for measuring temperature in body cavities.</p> <p>NOTE: A body cavity can be the mouth, rectum or armpit.</p> <p>The standard does not apply to clinical thermometers designed for special applications (e.g. thermometers for hypothermia) which owing to their measurement range, scale interval or maximum permissible error do not meet the requirements specified in this standard.</p>
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ICS 17.200.20

Võtmesõnad: clinical thermometers, definition, definitions, fever thermometers, fitness for purpose, measurement duration, medical equipment, operating requirements, specification (approval), specifications, testing, thermometers

ICS 17.200.20

English version

Clinical thermometers

Part 2: Phase change-type (dot matrix) thermometers

Thermomètres médicaux – Partie 2:
Thermomètres à changement de
phase (matrice à points)

Medizinische Thermometer – Teil 2:
Phasenumschlagthermometer
(Punktmatrix)

This European Standard was approved by CEN on 2000-09-16.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration.

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.

The European Standards exist 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, the Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, the Netherlands, Norway, Portugal, Spain, Sweden, Switzerland, and the United Kingdom.

CEN

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

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Foreword

This European Standard has been prepared by Technical Committee CEN/TC 205 "Non-active medical devices", the secretariat of which is held by BSI.

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 April 2001, and conflicting national standards shall be withdrawn at the latest by April 2001.

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

For relationship with EU Directive(s), see informative Annex ZA, which is an integral part of this standard.

This European Standard applies to clinical thermometers which are used for measuring the body temperature of humans.

EN 12470 consists of the following Parts under the general title "Clinical thermometers":

- Part 1: Metallic liquid-in-glass thermometers with maximum device
- Part 2: Phase change type (dot matrix) thermometers
- Part 3: Performance of compact electrical thermometers (non-predictive and predictive) with maximum device
- Part 4: Performance of electrical thermometers for continuous measurement
- Part 5¹ Performance of infra-red ear thermometers (with maximum device)

Annexes A and ZA are informative.

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

¹ In preparation

1 Scope

This Part of EN 12470 specifies performance requirements and test methods for phase change-type (dot matrix) thermometers for measuring temperature in body cavities.

NOTE: A body cavity can be the mouth, rectum or armpit.

This European Standard does not apply to clinical thermometers designed for special applications (e.g. thermometers for hypothermia) which owing to their measurement range, scale interval or maximum permissible error do not meet the requirements specified in this Standard.

2 Normative references

This European Standard incorporates by dated or undated reference, provisions from other publication. These normative references are cited at the appropriate places in the text and the publications are listed hereafter. For dated references, subsequent amendments to or revisions of any of these publications apply to this European Standard only when incorporated in it by amendment or revision. For undated references the latest edition of the publication referred to applies (including amendments).

EN 980	<i>Graphical symbols for use in the labelling of medical devices</i>
EN 1041	<i>Information supplied by the manufacturer with medical devices</i>
EN 556+A1	<i>Sterilization of medical devices - Requirements for terminally-sterilized medical devices to be labelled "Sterile"</i>
ISO 2859-2:1985	<i>Sampling procedures for inspection by attributes - Part 2: Sampling plans indexed by limiting quality (LQ) for isolated lot inspection</i>

3 Terms and definitions

For the purposes of this Part of EN 12470, the following terms and definitions apply:

3.1 measurement time

length of time required to measure body temperature.

3.2 phase change (dot matrix) thermometer

device utilising a change in state of chemical components designed to measure and indicate human body temperature.

3.3 retention time

duration of time for which the optimal signal for reading persists.

3.4 sensor matrix

temperature measuring area consisting of temperature dots.

NOTE: The dots contain different chemical mixtures, which change their state at specific temperatures. This change is accompanied by a change in appearance, e. g. change of colour. When in contact with the temperature site being measured, the change of state takes place in the sequence of dots up to and including the dot corresponding to the temperature of the site. This dot indicates the site temperature.

3.5 temperature offset

designed difference between preadjusted thermometer reading and water bath temperature after reaching thermal equilibrium.