SLOVENSKI STANDARD
SIST EN 1060-1:2000/A1:2002

01-november-2002

Neinvazivni sfigmomanometri - 1. del: Splošne zahteve

Non-invasive sphygmomanometers - Part 1: General requirements

Nichtinvasive Blutdruckmessgeräte - Teil 1: Allgemeine Anforderungen

Tensiometres non invasifs - Partie 1: Exigences générales

Ta slovenski standard je istoveten z: EN 1060-1:1995/A1:2002

ICS:

11.040.55 Diagnostiļ na oprema Diagnostic equipment

SIST EN 1060-1:2000/A1:2002 **en**

EUROPEAN STANDARD

NORME EUROPÉENNE

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English version

Non-invasive sphygmomanometers - Part 1: General requirements

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Nichtinvasive Blutdruckmessgeräte - Teil 1: Allgemeine Anforderungen

This amendment A1 modifies the European Standard EN 1060-1:1995; it was approved by CEN on 6 April 2002.

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

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Contents

Foreword

This document (EN 1060-1:1995/A1:2002) 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of an identical text or by endorsement, at the latest by November 2002, and conflicting national standards shall be withdrawn at the latest by November 2002.

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 EU Directive(s).

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, Malta, Netherlands, Norway, Portugal, Spain, Sweden, Switzerland and the United Kingdom.

Amended text

9.3 Marking of the device

Amend item a) to read 'indication of the correct positioning for the cuff over the artery;'.