STN

Neinvazívne sfygmomanometre Časť 2: Klinické vyšetrenie typu prerušovaného automatizovaného merania (ISO 81060-2: 2018/Amd 2: 2024) Zmena A2

STN EN ISO 81060-2/A2

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Non-invasive sphygmomanometers - Part 2: Clinical investigation of intermittent automated measurement type (ISO 81060-2:2018)

Táto norma obsahuje anglickú verziu európskej normy. This standard includes the English version of the European Standard.

Táto norma bola oznámená vo Vestníku ÚNMS SR č. 06/24

Obsahuje: EN ISO 81060-2:2019/A2:2024, ISO 81060-2:2018/Amd 2:2024

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English Version

Non-invasive sphygmomanometers - Part 2: Clinical investigation of intermittent automated measurement type - Amendment 2 (ISO 81060-2:2018/Amd 2:2024)

Sphygmomanomètres non invasifs - Partie 2: Investigation clinique pour type ponctuel à mesurage automatique - Amendment 2 (ISO 81060-2:2018/Amd 2:2024) Nichtinvasive Blutdruckmessgeräte - Teil 2: Klinische Prüfung der intermittierenden automatisierten Bauart - Änderung 2 (ISO 81060-2:2018/Amd 2:2024)

This amendment A2 modifies the European Standard EN ISO 81060-2:2019; it was approved by CEN on 26 February 2024.

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European foreword

This document (EN ISO 81060-2:2019/A2:2024) has been prepared by Technical Committee ISO/TC 121 "Anaesthetic and respiratory equipment" in collaboration with Technical Committee CEN/TC 205 "Non-active medical devices" the secretariat of which is held by DIN.

This Amendment to the European Standard EN ISO 81060-2:2019 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by October 2024, and conflicting national standards shall be withdrawn at the latest by October 2024.

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Endorsement notice

The text of ISO $81060-2:2018/Amd\ 2:2024$ has been approved by CEN as EN ISO 81060-2:2019/A2:2024 without any modification.



International Standard

ISO 81060-2

Non-invasive sphygmomanometers —

Part 2:

Clinical investigation of intermittent automated measurement type

AMENDMENT 2

Sphygmomanomètres non invasifs —

Partie 2: Investigation clinique pour type ponctuel à mesurage automatique

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This document was prepared jointly by Technical Committee ISO/TC 121, Anaesthetic and respiratory equipment, Subcommittee SC 3, Respiratory devices and related equipment used for patient care, and Technical Committee IEC/TC 62, Medical equipment, software, and systems, Subcommittee SC 62D, Particular medical equipment, software, and systems, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 205, Non-active medical devices, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

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