OffiSoins

CONTROLE

Automatische polsbloeddrukmeter

METING VAN DE DIASTOLISCHE EN SYSTOLISCHE BLOEDDRUK EN HARTSLAG TER HOOGTE VAN DE POLS

+ 2 jaar garantie
+ Eenvoudige meting met één knop
+ Meting volgens de oscillometrische methode
+ Detectie van hartritmestoornissen
+ Klinisch gevalideerd
+ Groot display
+ Grote manchet (13,5 tot 21,5 cm)
+ 2 gebruikers x 199 geheugenplaatsen per gebruiker
+ Datum- en tijdsfunctie inbegrepen bij meetgegevens
+ Gemiddelde berekening van 3 metingen
+ Automatische uitschakeling en energiebesparing
+ Meetbereiken:

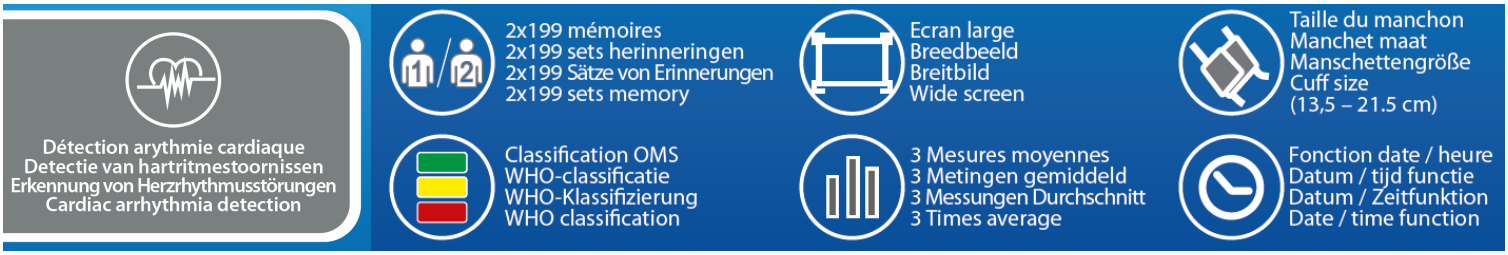
  Dia. / Sys. : 0 - 299 mmHg
  Puls. : 40 - 199 puls./min

+ Meetnauwkeurigheid:

  Dia. / Sys. : ± 3 mmHg
  Puls. : ± 5%

+ Wordt geleverd met 2 AAA-batterijen
+ CE 0123

4613519



Omschrijving	EAN13	CNK	Verpakt per
OFFISOINS Automatische polsbloeddrukmeter	7863124802331	4613519	1

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 078672 0014 Rev. 01

Manufacturer:

Shenzhen Urion Technology Co., Ltd.
Floor 4-6th of Building D
Jiale Science & Technology Industrial Zone
No.3, ChuangWei Road
Heshuikou Community, MaTian Street
GuangMing New District
518106 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Digital Blood Pressure Monitors, Infrared Thermometer, Compressor Nebulizers**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G1 078672 0014 Rev. 01

Report No.: GZ2117902

Valid from: 2021-04-12
Valid until: 2024-05-26

Date, 2021-04-12

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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