



Declaration of Conformity

Certificate No:1716

The below listed medical product(s) complies with the

Medical Devices Directive No. 93/42/EEC

Annex V

The product(s) therefore fulfill all essential requirements for the application of the CE-conformity mark.

Product type: Automatic-inflation electronic sphygmomanometer, portable, arm/wrist
GMDN code: 45617 (UMDNS 16-157)
Product type: Blood Pressure Long-Term recorder ambulatory recorder
GMDNS Code: 36888 (UMDNS 12-386)
Product class: Class IIa
Brand name: WATCH BP

Customers type no.:	Manufacturers type no.:
WatchBP Home	WatchBP Home (ERP BP3MX1-1)
WatchBP Home A	WatchBP Home A (ERP BP3MX1-3)
WatchBP Home S	WatchBP Home S (ERP BP3MX1-5)
WatchBP Office AFIB	WatchBP Office AFIB (ERP TWIN200 AFS)
WatchBP Office ABI	WatchBP Office ABI (ERP TWIN200 ABI)
WatchBP Office Target	WatchBP Office Target (ERP BP3MD1-4)

Blood pressure long-term ambulatory recorder GMDNS 36888 (UMDNS 12-386)

Customers type no.:	Manufacturers type no.:
WatchBP 03	WatchBP 03 (ERP BP3MZ1-1)

Manufacturing plant ONBO ELECTRONIC (SHENZHEN) CO. LTD.

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Manufacturer Microlife AG
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Declare under our sole responsibility that the above product fulfills the essential requirements of the Directive 93/42/EEC, including all amendments.

microlife®

EU Representative: Microlife UAB, P. Lukšio g. 32, 08222 Vilnius, Lithuania

Responsible importer: Microlife UAB
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LITHUANIA

Remarks: *Notified Body according to the Medical Devices Directive is TÜV NORD CERT GmbH, Langemarckstrasse 20, 45141 Essen, Germany. Notified Body ID. No. 0044 Certificate No: 04 235 001845, valid until: 20. December 2023*

Place and Date of Issue: Widnau, 5th July 2021
for the Manufacturer:

for the Importer:

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