microlife®

Declaration of Conformity

Certificate No:1767

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 Blood Pressure Long-Term recorder ambulatory recorder

GMDN code: 45617 (UMDNS 16-157), 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 N	WatchBP Home N (ERP BP3MX1-4)
WatchBP Home S	WatchBP Home S (ERP BP3MX1-5)
WatchBP 03	WatchBP 03 (ERP BP3MZ1-1)
WatchBP 03 AFIB	WatchBP 03 AFIB (ERP BP3MZ1-1A)
WatchBP Office	WatchBP Office (ERP TWIN 200)
WatchBP Office AFIB	WatchBP Office AFIB (ERP TWIN 200AFS)
WatchBP Office ABI	WatchBP Office ABI (ERP TWIN 200ABI)
WatchBP Office Central	WatchBP Office Central (ERP Twin 200CBP)

Manufacturing plant: ONBO ELECTRONIC (SHENZHEN) CO. LTD.

Head office: microlife Corporation
9F, 431, GuiGang Road, Nei Hu, Taipei 11492, Taiwan, R.O.C.

Manufacturer: Microlife AG
Espenstrasse 139, 9443 Widnau, Switzerland

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

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

Responsible importer: Colpharma s.r.l.
Via A. M Vicenzi, 19/a
43124 Parma
ITALY

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, 19th January 2022

for the Manufacturer:

Teresa Meile
Regulatory Manager

for the Importer: