microlife Declaration of Conformity

Certificate No: 1863

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 16-173)

36888 (UMDNS 12-386)

Product class: Class IIa
Brand name: WATCHBP

Automatic-inflation electronic sphygmomanometer, portable, arm/wrist		
Customers type no.:	Manufacturers type no.:	
WatchBP Office vascular	WatchBP Office vascular (ERP TWIN200VSR)	
WatchBP Office 2G	WatchBP Office 2G (ERP BP3SK1-3B)	
WatchBP Home A	WatchBP Home A (ERP BP3MX1-3)	
WatchBP Home S	WatchBP Home S (ERP BP3MX1-5)	

Blood Pressure Long-Term recorder ambulatory recorder		
Customers type no.:	Manufacturers type no.:	
WatchBP 03 2G	WatchBP 03 2G (ERP BP3SZ1-1)	

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

Head office: microlife Corporation

9F, 431, RuiGuang Road, Taipei 114, 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: Microlife France

118 rue de Rivoli, 75001 Paris, France

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: for the Manufacturer:

Widnau, 20th December 2022

for the Importer:

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Sascha Breuss Regulatory Manager