microlife®

Declaration of Conformity

Certificate No: 1600

The below listed medical product(s) complies with the

Medical Devices Directive No. 93/42/EEC

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

Annex V

Product type:	Automatic-inflation electronic sphygmomanometer, portable, arm/wrist
GMDN code:	45617 (UMDNS 16-157)
Product class:	Class IIa
Brand name:	MICROLIFE

Customers type no.:	Manufacturers type no.:
BP B3 AFIB	BP B3 AFIB (ERP BP3KT1-3N)

Manufacturing plant ONBO ELECTRONIC (SHENZHEN) CO. LTD.

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

Manufacturer & European Subsidiary:
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.

Responsible importer:
Egli AG Import und Vertrieb
Im Langhang 11
8307 Effretikon
SWITZERLAND

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, 28th July 2020

for the Manufacturer:

microlife
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9443 Widnau / Switzerland
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Teresa Meile
Regulatory Manager

for the Importer: