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Declaration of Conformity

Certificate No:1768

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 Intermittent electronic patient thermometer
GMDN code:	45617 (UMDNS 16-157), 14035 (UMDNS 14-032)
Product type:	Infrared patient thermometer, skin Infrared patient thermometer, ear
GMDN code:	17888 (UMDNS 14-036), 17887 (UMDNS 14-036)
Product class:	Class IIa
Brand name:	MICROLIFE

Blood Pressure Monitor GMDN 45617	
Customers type no.:	**Manufacturers type no.:**
BP A2 Basic	BP A2 Basic (ERP BP3GQ1-3P)
BP A2 Classic	BP A2 Classic (ERP BP3UG1-2E)
BP A2 Easy	BP A2 Easy (ERP BP3GQ1-1P)
BP A3 Easy	BP A3 Easy (ERP BP3GX1-2N)
BP A3 Plus	BP A3 Plus (ERP BP3GX1-5N)
BP A6 Advanced Easy	BP A6 Advanced Easy (ERP BP3GU1-3E)
BP A6 BT	BP A6 BT (ERP BP3GU1-7B)
BP A6 PC	BP A6 PC (ERP BP3GU1-8Y)
BP A7 Touch	BP A7 Touch (ERP BP3GT1-6Y)
BP B1 Classic	BP B1 Classic (ERP BP3KE1-3E)
BP B3 AFIB	BP B3 AFIB (ERP BP3KT1-3N)
BP B6 Connect BT	BP B6 Connect (ERP BP3KV1-5X)
BP W100	BP W100 (ERP BP3MK1)
BP A150 AFIB	BP A150 AFIB (ERP BP3MS1-2D)

Digital Thermometer GMDN 14035	
Customers type no.:	**Manufacturers type no.:**
MT 16C2	MT 16C2
MT 16F1	MT 16F1
MT 1931	MT 1931
MT 400	MT 400 (ERP MT1P11)
MT 700	MT 700 (ERP MT17K1)
MT 850	MT 850 (ERP MT1PG1)

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Infrared Thermometer GMDN 17888, 17887	
Customers type no.:	**Manufacturers type no.:**
NC 150	NC 150 (ERP FR1MF1)
NC 150 BT	NC 150 BT (ERP FR1MF1-B)
NC 200	NC 200 (ERP FR1DG1)
IR 150	IR 150 (ERP IR1DF1-1)
IFR 100	IFR 100 (ERP IFR1DU1)

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:
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, 20th January 2022

for the Manufacturer: **for the Importer:**

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Regulatory Manager