



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 17 08 99651 002

Manufacturer:**TriMedika Ltd**

Unit 3 E3 Building, Springvale Campus
398 Springfield Road
Co Antrim
Belfast, Northern Ireland
BT12 7DU
UNITED KINGDOM

**Facility(ies):**

TriMedika Ltd
Unit 3 E3 Building, Springvale Campus, 398 Springfield Road, Co
Antrim, Belfast, Northern Ireland, BT12 7DU, UNITED KINGDOM

**Product
Category(ies):****Infra Red Thermometers
for Clinical use**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 75938830

Valid from: 2017-10-02

Valid until: 2022-10-01

Date, 2017-10-02

Stefan Preiß



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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