



# 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 101902 0004 Rev. 00**

**SAO Y BẢN CHÍNH**  
Ngày.....tháng.....năm 20....

**Manufacturer:** **BPL MEDICAL TECHNOLOGIES PVT LTD**  
11th km, Bannerghatta Road, Arakere  
Karnataka  
Bangalore 560076  
INDIA

**Product Category(ies):** **Electrocardiograph, Syringe & Volumetric Pump, Non-contact Infrared Thermometer, Finger-tip Pulse Oximeter, Foetal Doppler, Patient Monitor, Ultra sound scanner, Holter Monitoring System.**

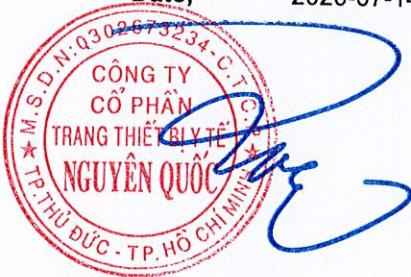
The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms 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.:** IND20190113

**Valid from:** 2020-07-14

**Valid until:** 2024-05-26

**Date,** 2020-07-14



Christoph Dicks  
Head of Certification/Notified Body



**TỔNG GIÁM ĐỐC**  
*Lưu Thất Chuẩn*



# EC Certificate

Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)

**No. G2 101902 0005 Rev. 00**

**Manufacturer:**

**BPL MEDICAL TECHNOLOGIES PVT LTD**

11th km, Bannerghatta Road, Arakere  
Karnataka  
Bangalore 560076  
INDIA

**Product  
Category(ies):**

**Non Invasive Blood Pressure Monitor, Oxygen concentrator,  
Surgical Gloves**

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

**Report No.:** IND20190113

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