SAO Y BẢN CHÍNH
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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

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
Qua Thất Chuân

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

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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.

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