EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60144433 0001

Report No.: 17054024 008

Manufacturer:
Shenzhen Upnmed Equipment
Co., Ltd.
4th Floor, Building #1 East
Huihuang Industrial Area
Xitian Community, Gongming Town
Guangming New District
Shenzhen
518107 Guangdong
China

Products:
Oximeter Probes

Replaces Approval, Registration No.: HD 60126823 0001

Digitally signed by Ermicev Alexandr
Date: 2021.12.20 16:23:38 EET
Reason: MoldSign Signature
Location: Moldova

Expiry Date: 2024-05-27

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2019-12-02

Date: 2019-12-02

Notified Body

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.