EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60153957 0001

Report No.: 16801929 013

Manufacturer:
Lights Medical Manufacture Co.,Ltd.
No. 19, Quanda Road,
Wuqing Development Area
Tianjin, 301700
P.R. China

Products: Wet Pack Products

Replaces Approval, Registration No.: DD 60143690 0001

Digitally signed by Jighili Tatiana
Date: 2023.01.05 14:45:07 EET
Reason: MoldSign Signature
Location: Moldova

Expiry Date: 2023-04-17

The Notified Body hereby declares that the requirements of Annex V 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2021-04-01

Date: 2021-04-01

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Notified Body

Wenxiang Zhang

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.