EC Certificate

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

TÜVRheinland®

Registration No.: HD 60135854 0001

Report No.: 17037792 010

Manufacturer: Taishan Anson
Electrical Appliances Co., Ltd.
Dragon Mountain
Industrial Estate, Duanfen
529245 Taishan, Guangdong province
China

Products: Phototherapy Units

Replaces Approval, Registration No.: HD 60092937 0001

Expiry Date: 2024-04-24

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: 2019-04-25

Date: 2019-03-15

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle
Notified Body
TÜVRheinland®

S. Liu

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