

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

Registration No.: HD 60145717 0001

Report No.: 21192387 016

Manufacturer: Schulze & Böhm GmbH
Hermülheimer Str. 10
50321 Brühl
Deutschland

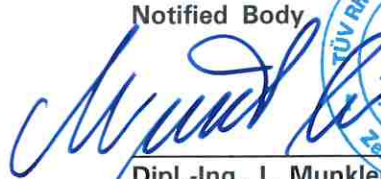

Products: Phototherapy equipment and Balneo-Phototherapy equipment
Replaces Certificate, Registration No.: HD 60128600 0001

Expiry Date: 2024-05-26

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: 2020-01-07

Date: 2020-01-07

Notified Body

Dipl.-Ing. I. Munkler


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.