

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

**Registration No.:** DD 60148985 0001

**Report No.:** 17062485 009

**Manufacturer:** Biocare Enterprise Limited  
Flat 1A, 9/F, Brill Plaza, No.84  
Tokwawan Road  
Kowloon  
Hong Kong

**Products:** Low-intensity Laser Devices  
Replaces Approval, Registration No.: DD 60138420 0001

**Expiry Date:** 2024-05-26

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:** 2020-09-17

**Date:** 2020-09-17

Notified Body



Dipl.-Ing. W. Hsu

**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.