# EC Certificate

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

**Registration No.:** DD 60138420 0001

**Report No.:** 17062485 004

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

**Products:** Low-intensity Laser Devices

**Expiry Date:** 2022-03-08

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:** 2019-10-24

**Date:** 2019-10-24

Notified Body
S. Liu

**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.