Lloyd's Register
LRQA

# EC CERTIFICATE – PRODUCTION QUALITY ASSURANCE

## In accordance with the requirements of the Medical Devices Directive 93/42/EEC and the Medical Devices Regulations 2002, UK Statutory Instrument 2002 No. 618

This is to certify that the Quality Management System of:

**MEDlight GmbH**
**Werrestrasse 94,**
**D-32049 Herford,**
**Germany**

has been assessed against the requirements of Annex V of the Medical Devices Directive 93/42/EEC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown on the attached schedule.

Approval is subject to the maintenance of the quality system in accordance with the requirements of the above Directive and Regulations.

Authorisation is hereby given to use the LRQA Notified Body Registration Number in accordance with the requirements of the specified Directives/Regulations in relation to the products as identified above.

| | |
|---|---|
| Certificate No: | LRQ 4001336/A |
| Original Approval: | 7 January 2004 |
| Current Certificate: | 1 April 2016 |
| Certificate Expiry: | 31 March 2019 |

LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited

1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom

Lloyd's Register Group Limited, its affiliates and subsidiaries, including Lloyd's Register Quality Assurance Limited (LRQA), and their respective officers, employees or agents are, individually and collectively, referred to in this clause as 'Lloyd's Register'. Lloyd's Register assumes no responsibility and shall not be liable to any person for any loss, damage or expense caused by reliance on the information or advice in this document or howsoever provided, unless that person has signed a contract with the relevant Lloyd's Register entity for the provision of this information or advice and in that case any responsibility or liability is exclusively on the terms and conditions set out in that contract.

Lloyd's Register
LRQA

# EC CERTIFICATE – PRODUCTION QUALITY ASSURANCE CERTIFICATE LRQ 4001336/A SCHEDULE

## In accordance with the requirements of the Medical Devices Directive 93/42/EEC and the Medical Devices Regulations 2002, UK Statutory Instrument 2002 No. 618

**MEDlight GmbH**
**Werrestrasse 94,**
**D-32049 Herford,**
**Germany**

**Class IIa Products**
Phototherapy Devices
Photodiagnostic Devices
BEAMpro Series

Schedule Issue: 01

Date of Schedule Issue: 1 April 2016

LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited



1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom

Lloyd's Register Group Limited, its affiliates and subsidiaries, including Lloyd's Register Quality Assurance Limited (LRQA), and their respective officers, employees or agents are, individually and collectively, referred to in this clause as 'Lloyd's Register'. Lloyd's Register assumes no responsibility and shall not be liable to any person for any loss, damage or expense caused by reliance on the information or advice in this document or howsoever provided, unless that person has signed a contract with the relevant Lloyd's Register entity for the provision of this information or advice and in that case any responsibility or liability is exclusively on the terms and conditions set out in that contract.