

EC Certificate

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

Registration No.:

HD 1499629-1

Manufacturer:

LASOTRONIX Sp. z o.o.

Elektroniczna 2A 05-500 Piaseczno

Poland

Products:

- Medical diode lasers

Replaces EC Certificate number HD 1499629-1 dated 2020-07-16

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.

Report No.:

84953464-20

Effective date:

2021-05-25

Expiry date:

2024-05-26

Issue date:

2021-05-25

Jarosław Pyclik TÜV Rheinland LGA Products GmbH Tillystraße 2 · 90431 Nürnberg · Germany

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

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SMART M laser, models:

- SMART M ST1

- SMART M ST2

- SMART M ST3

- SMART M DR1

- SMART M DR2

- SMART M DR3

- SMART M SG1

- SMART M SG2

- SMART M SG3

- SMART M SG4

- SMART M SG5

- SMART M GN1

- SMART M GN2

- SMART M GN3

- SMART M PH1

- SMART M PH2

- SMART M PH3

- SIVIART IVI FITS

- SMART M PH4

- SMART M LG1

- SMART M LG2

- SMART M NS1

- SMART M VT1

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