

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

Registration No.: HD 60146857 0001

Report No.: 21256460 016

Manufacturer: STORZ MEDICAL AG
Lohstampfestr. 8
8274 Tägerwilen
Schweiz

Products:

- Equipment for the extracorporeal induced shock wave and pressure pulse therapy for stationary and mobile use
- Equipment for the extracorporeal magnetotransduction therapy
- X-ray application devices (without radiation components) (see attachment for products included)

Replaces Certificate, Registration No.: HD 60140661 0001

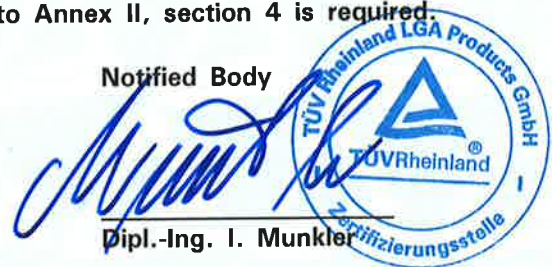
Expiry Date: 2024-05-26

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: 2020-02-18

Date: 2020-02-18

Notified Body



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.

**TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg**

**Attachment to
Certificate**

Registration No.: HD 60146857 0001
Report No.: 21256460 018

Manufacturer: **STORZ MEDICAL AG**
Lohstampfestr. 8
8274 Tägerwilen
Schweiz

Equipment for the extracorporeal induced shock wave therapy
for stationary and mobile use:

- MODULITH SLK
with options
 - LITHOTRACK
 - US-SET

- MODULITH SLX-F2
with options
 - C-ARM C-MX
 - US-SET
 - StorM-Touch
 - MONITORARM

Equipment for the extracorporeal induced cardiac shock wave
therapy for treatment of ischemic heart disease:

- MODULITH SLC

Date: 2020-05-06

Notified Body

Roland Gruber



TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: HD 60146857 0001
Report No.: 21256460 018

Manufacturer: **STORZ MEDICAL AG**
Lohstampfestr. 8
8274 Tägerwilen
Schweiz

X-ray application devices (without radiation components):
- C-MX

Equipment for the extracorporeal pneumatically operated
ballistic pressure wave therapy:

- MASTERPULS MP100
- MASTERPULS MP200
- MASTERPULS MP50
- MASTERPULS ONE
- D-ACTOR 100
- D-ACTOR 200
- D-ACTOR 50
- D-ACTOR ONE
- CHATTANOOGA MOBILE RPW - 2805
- CHATTANOOGA MOBILE 2 RPW - 2905
- CHATTANOOGA INTELECT RPW Lite

Date: 2020-05-06

Notified Body

Roland Gruber



TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: HD 60146857 0001
Report No.: 21256460 018

Manufacturer: **STORZ MEDICAL AG**
Lohstampfestr. 8
8274 Tägerwilen
Schweiz

Equipment for the extracorporeal induced shock and pressure wave therapy for stationary and mobile use:

- DUOLITH SD1 T-Top [001x]
- DUOLITH SD1 T-Top [010x]
- DUOLITH SD1 Tower
- CELLACTOR SC1 T-Top
- CELLACTOR SC1 Tower
- CELLIMPACT
- CHATTANOOGA INTELECT F-SW - 21095
- W-MEDICAL SHOCKWAVE F1
- NEUROLITH
- eSparkPower

Equipment for the extracorporeal magnetotransduction therapy

- MAGNETOLITH
- Gymna Magnacure 300

Date: 2020-05-06

Notified Body

Roland Gruber

