

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

**Registration No.:** HD 60115150 0001

**Report No.:** 21256460 001

**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
- X-ray application devices (without radiation components)

(see attachment for products included)

Replaces Certificate, Registration No.: HD 60103173 0001

**Expiry Date:**

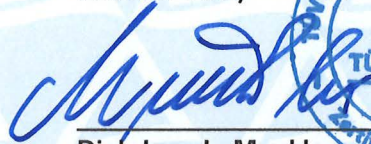

2021-11-18

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:** 2017-08-02

**Date:** 2017-08-02

Notified Body

  
Dipl.-Ing. I. Munkler  


**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**

Doc. 1/3, Rev. 2

**Attachment to  
Certificate**

**Registration No.:** HD 60115150 0001  
**Report No.:** 21256460 011

**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
  - StörM-Touch
  - MONITORARM

**Date:** 2019-06-03

Notified Body



**Dipl.-Ing. I. Munkler**



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

Doc. 2/3, Rev. 2

**Attachment to  
Certificate**

**Registration No.:** HD 60115150 0001  
**Report No.:** 21256460 011

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

X-ray application devices (without radiation components):

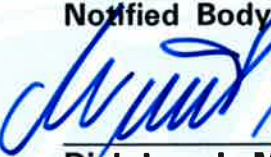
- C-MX

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: 2019-06-03**

**Notified Body**



**Dipl.-Ing. I. Munkler**

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

Doc. 3/3, Rev. 2

**Attachment to  
Certificate**

**Registration No.:** HD 60115150 0001  
**Report No.:** 21256460 011

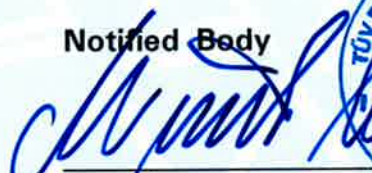
**Manufacturer:** **STORZ MEDICAL AG**  
**Lohstampfstr. 8**  
**8274 Tägerwilen**  
**Schweiz**

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

**Date:** 2019-06-03

**Notified Body**

  
**Dipl.-Ing. I. Munkler**

