

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

**Registration No.:** HD 60146581 0001

**Report No.:** 26300490 002

**Manufacturer:** Astar Spółka z Ograniczoną  
Odpowiedzialnością  
ul. Świt 33  
43-382 Bielsko-Biała  
Polska

**Products:** see attachment for products and site included

**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:** 2020-02-12

**Date:** 2020-02-12

**Notified Body**

*Maciej Sciera*

**Maciej Sciera**



**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. 1/1, Rev. 0

**Attachment to  
Certificate**

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ul. Świt 33  
43-382 Bielsko-Biała  
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**Products included:**

- Electrotherapy devices
- Laser therapy devices
- IR therapy devices
- Magnetic field therapy devices
- Ultrasound therapy devices
- Ultrasound therapy combined with electrotherapy devices
- Vacuum therapy devices
- Shock wave physical therapy devices

**Additional site included:**

Astar Sp. z o.o.  
ul. Pod Mlynska Kepa 748  
43-384 Jaworze, Poland

**Activity:** Manufacture

**Date:** 2020-02-12

**Notified Body**

*Sciera Maciej*  
**Maciej Sciera**

