

EC Certificate Directive 93/42/EEC Annex V

Production Quality Assurance
Medical Devices

Registration No.: DD 6

DD 60141729 0001

Report No.:

21246563 023

Manufacturer:

MED TRUST Handelsges.m.b.H.

Gewerbepark 10

7221 Marz Österreich

Products:

(see attachment for products included)

Expiry Date:

2024-05-26

The Notified Body hereby declares that the requirements of Annex V 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date:

2019-09-24

Date:

2019-09-24

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.

Notified Body

Dipl.-Ing. I. Munkler invierung

ÜVRheinland



Doc. 1/1, Rev. 0

TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Attachment to Certificate

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Manufacturer:

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Österreich

Products included:

- Disposable Syringes
- Blood Lancets
- Hypodermic Needles
- Electronic Blood Pressure Monitors

Date: 2019-09-24

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