

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

**Registration No.:** HD 60149031 0001

**Report No.:** 21273496 008

**Manufacturer:** VacuTec Meßtechnik GmbH  
Dornblüthstr. 14 a  
01277 Dresden  
Deutschland

**Products:** Detectors and systems for the detection and measurement of ionizing radiation  
  
(see attachment for products included)

Replaces Certificate, Registration No.: HD 60127716 0001

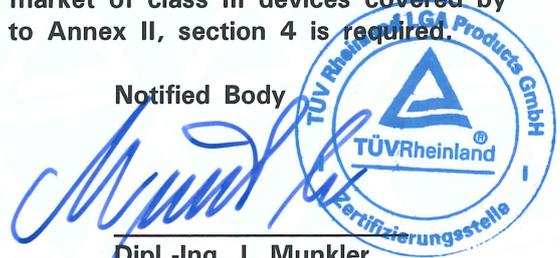
**Expiry Date:** 2024-05-26

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:** 2020-04-28

**Date:** 2020-04-28

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**

**Attachment to  
Certificate**

**Registration No.:** HD 60149031 0001  
**Report No.:** 21273496 008

**Manufacturer:** VacuTec Meßtechnik GmbH  
Dornblüthstr. 14 a  
01277 Dresden  
Deutschland

**Products included:**

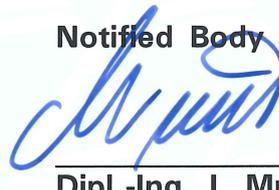
- Ionization chambers for automatic X-ray exposure control

For the following devices the scope covers only the aspects of manufacture concerned with conformity of the products with the metrological requirements:

- Dose area product meters

**Date:** 2020-04-28

**Notified Body**



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