

**EC Certificate**  
**Directive 93/42/EEC Annex V**  
**Production Quality Assurance**  
**Medical Devices**

**Registration No.:** DD 60145315 0001

**Report No.:** 21190672 020

**Manufacturer:** Argen Dental GmbH  
Fritz-Vomfelde-Str. 12  
40547 Düsseldorf  
Deutschland

**Products:** Dental materials:  
- Acrylic teeth  
- Dental PMMA-discs for CAD/CAM processing

Replaces Certificate, Registration No.: DD 60124682 0001



**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:** 2019-12-13

**Date:** 2019-12-13

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.