

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

**Registration No.:** HD 60128931 0001

**Report No.:** 16802202 007

**Manufacturer:** Liaoning Upcera Co.,Ltd.  
No.122 Xianghuai Road  
Economic Development Zone  
Benxi  
117004 Liaoning  
China

**Products:** Dental Zirconia Ceramics, Dental Lithium Disilicate Glass Ceramics, Color for the staining of Zirconia Ceramics, Dental Filling/Restorative Polymer Based Block, Glaze Paste

Replaces Approval, Registration No.: DD 60111717 0001

**Expiry Date:** 2023-06-12

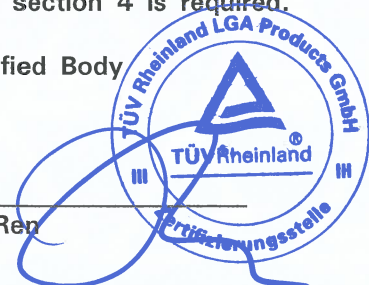
The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, 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:** 2018-06-13

**Date:** 2018-06-06

Notified Body

X. Ren



**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.