# 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. Rep.

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.