EC Certificate

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

TÜVRheinland®

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

TÜV Rheinland LGA Products GmbH
TÜV Rheinland®
III III
Zertifizierungsstelle

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.