TÜVRheinland®

APPROVAL

EC Directive 93/42/EEC Annex V, Article 3
Quality Assurance System Production

Registration No.: DD 60037333 0001

Report No.: 17018508 001

Manufacturer: Foshan Roson Medical
Instrument Co., Ltd.
No.9 Henggui ZhongRoad
Lianhe Avenue, Luocun
Nanhai District, Foshan City

528226 Guangdong Province
China

Scope: Manufacture of Dental Units

Replaces Approval, Registration No.: DD 60016119 0001

Date of Expiry: 11.02.2016

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex V, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex V, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Date 21.03.2011

Dipl.-Ing. D. Meier

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. 0197 to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. CE