



APPROVAL

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

Registration No.: DD 60016119 0001

Report No.: 16007614 001

Manufacturer: Foshan Roson
Medical Instrument Co., Ltd.
No. 2, 12th Road, West Section
Lianhe Avenue, Luocun

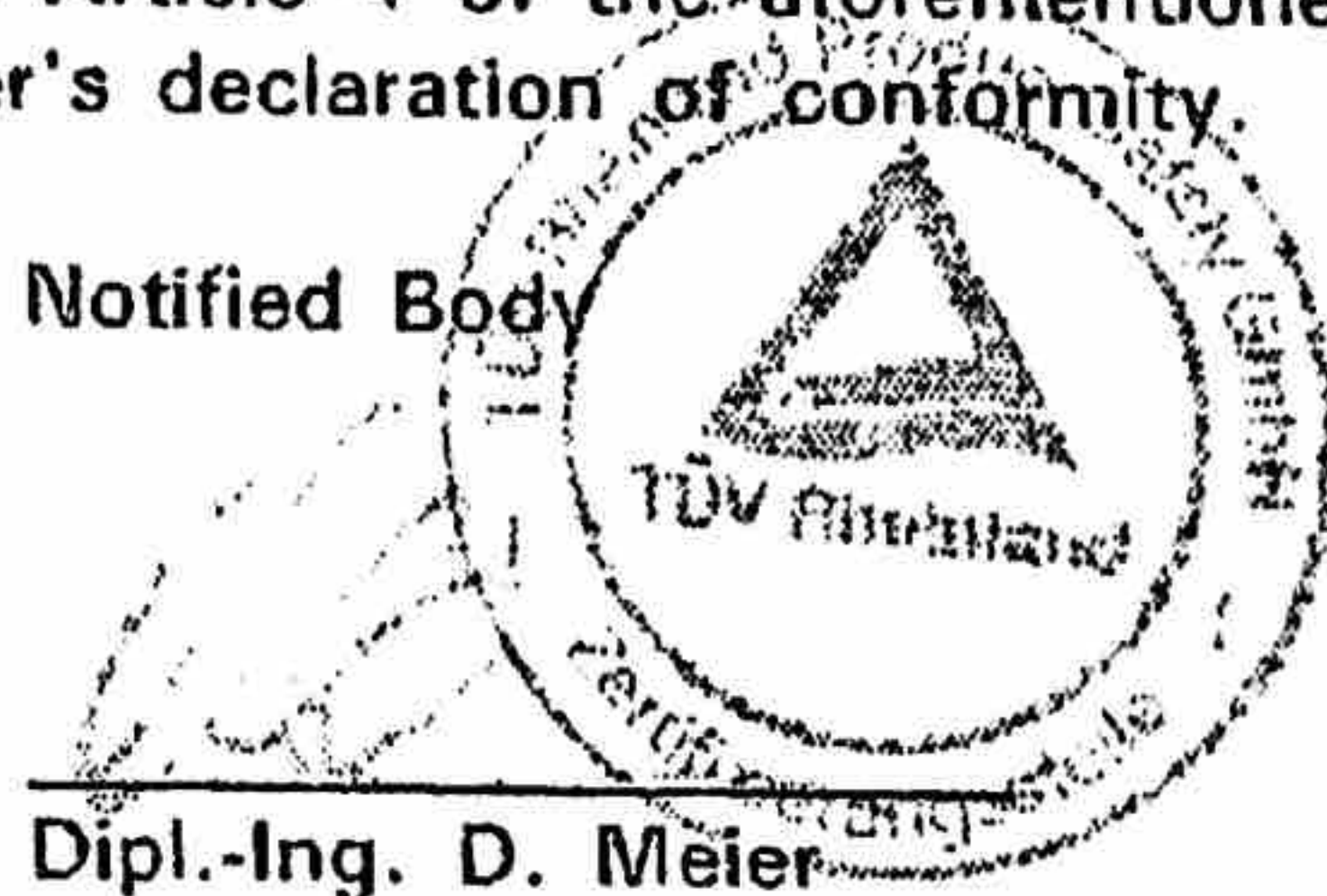
Foshan, Guangdong
China

Scope: Manufacturing and Sales of Dental Units

Date of Expiry: 29.01.2020

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



Cologne, 30.01.2007

TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln
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.

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

