APPROVAL

EC Directive 93/42/EEC Annex II, Article 3
Full Quality Assurance System
Medical Devices

Registration No.: HD 60015343 0001

Report No.: 17004876 001

Manufacturer:
Unimed Medical Supplies, Inc.
The 5th floor, NO 6 Building
Nanshui Industrial Zone
NO 5 Industrial Road, Shekou

Shenzhen 518067
China

Scope: Design, Manufacturing and Sales of Oximeter Probes

Date of Expiry: 15.08.2011

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex II, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex II, Article 5 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body

TÜV Rheinland Product Safety GmbH
TÜV Rheinland

Cologne, 16.08.2006

Dipl.-Ing. D. Me

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.

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