



**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

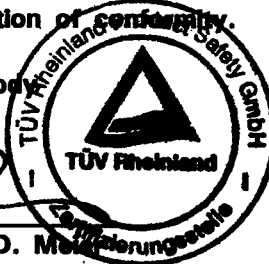
**Date of Expiry:** 15.08.2011

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.

Cologne, 16.08.2006

Notified Body

Dipl.-Ing. D. Meier



**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