

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

Registration No.: HD 60017553 0001

Report No.: 02421383 002

Manufacturer: Mediaid Inc.,
17517 Fabrica Way, Suite H
Cerritos, CA 90703
USA

Scope: Design, Manufacturing and Sales of Pulse Oximeters and
Pulse Oximeter Sensors

Production facilities: see attachment

Replaces Approval, Registration No.: HD 60006298 0001

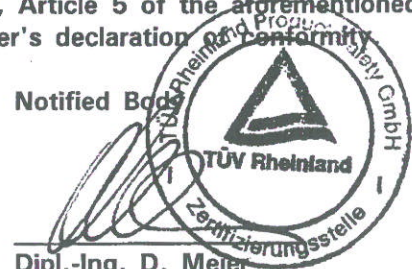
Date of Expiry: 03.03.2012

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

Dipl.-Ing. D. Meier

Cologne, 24.04.2007



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.