

Certificate

The Certification Body of
TÜV Rheinland Product Safety GmbH

hereby certifies that the organization

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

has established and applies a quality management system for medical devices
for the following scope:

**Design, Manufacturing and Sales of Patient Monitors,
Pulse Oximeter and Pulse Oximeter Sensors
Production facilities: see attachment**

Proof has been furnished that the requirements specified in

EN ISO 13485:2003

are fulfilled. The quality management system is subject to yearly surveillance.

Certificate Registration No.: SX 60013278 0001

An audit was performed. Report No.: 02421383 002

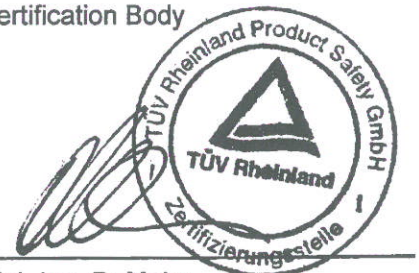
This Certificate is valid until: 03.03.2012



Akkreditiert durch
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln
und Medizinprodukten
ZLG-ZQ-995.00.01-46

Cologne, 24.04.2007

Certification Body



Dipl.-Ing. D. Meier

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

Tel.: (+49/221) 806 - 1371 Fax: (+49/221) 806 - 3935 e-mail: cert-validity@de.tuv.com <http://www.tuv.com/safety>