



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

Registration No.: HD 60007286 0001

Report No.: 02271485 002

Manufacturer: Defibtech, LLC  
753 Boston Post Road, Suite 102  
Guilford, CT 06437  
USA

Scope: Design, Production, Sales and Services for  
Semi Automatic External Defibrillator

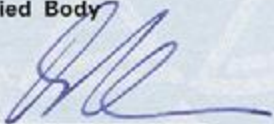
Date of Expiry: 10.02.2009

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, 11.02.2004



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