

	
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
<p>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.</p>	
Cologne, <u>11.02.2004</u>	 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	