

# Certificate

The Certification Body of  
**TÜV Rheinland Product Safety GmbH**

hereby certifies that the organization  
**TechDevice Corporation**  
**650 Pleasant St.**  
**Watertown, MA 02472**  
**USA**

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

**Design and Development and Manufacture of Guidewires,  
Balloon Dilatation Catheters and Latex Balloon Catheters  
Replaces Certificate, Registration No.: SX 60013912 0001**

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 60019520 0001

An audit was performed. Report No.: 30792400 001

This Certificate is valid until: 19.12.2012

Certification Body



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

Cologne, 01.04.2008



*B. Ludovico*  
B. Ludovico

**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>