

Certificate



TUV Rheinland of North America, Inc., a CMDCAS
recognized registrar, certifies that

Terumo Medical Corporation
950 Elkton Blvd
Elkton, MD 21921
USA

has established and maintained a

Quality Management System
according to
ISO 13485:2003

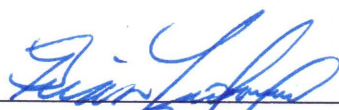
Audit Report No.: 30690450 003
Certificate Registration No.: 74 500 3619
Expiry Date: March 04, 2012

for the Design and Development and Manufacturing of

Disposable Medical Devices, In-Vitro Diagnostic
Devices and Sterile Tubing Welders

(see attachment for additional products covered by this registration)





Certification Officer: Brian Ludovico
TUV Rheinland of North America, Inc.
Newtown, Connecticut
Effective Date: April 07, 2009

Attachment

**Quality Management System
according to ISO 13485:2003**

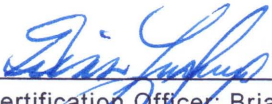
for

**Terumo Medical Corporation
950 Elkton Blvd
Elkton, MD 21921
USA**

The scope of the registration includes the following products:

- Syringes
- Needles
- Intravenous Catheters
- Introducer Kits
- Guiding Sheaths
- Injection Plugs
- Capillary Blood Collection Devices
- Sterile Tubing Welders
- Lancets

Audit Report No.: 30690450 003
Certificate Registration No.: 74 500 3619
Expiry Date: March 04, 2012



Certification Officer: Brian Ludovico
TUV Rheinland of North America, Inc.
Newtown, Connecticut
Effective Date: April 07, 2009