Certificate

TÜVRheinland®

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

Terumo Medical Corporation
950 Elkton Boulevard
Elkton MD 21921
USA

has established and maintained a

Quality Management System
according to
ISO 13485:2003

Audit Report No.: 31193020 001
Certificate Registration No.: 74 500 4146
Expiry Date: February 16, 2015

For the Design and Development and Manufacturing of

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

(see attachment for additional products under registration)

SCC Accredited CB-MS™

Certification Officer: B. Ludovico

TUV Rheinland of North America, Inc.
Newtown, Connecticut

Effective Date: February 27, 2012

TÜVRheinland®

Doc. 1/1, Rev. 0

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
- Injection Plugs
- Guiding Sheaths
- Support Catheters
- Micro Blood Collection Devices
- Lancets

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Effective Date: February 27, 2012

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