

DECLARATION OF CONFORMITY

Manufacturer: Abbott Vascular

Address: 3200 Lakeside Drive
 Santa Clara, California 95054, USA

Manufacturing Sites: 26531 Ynez Road
 Temecula, California 92591, USA

30590 Cochise Circle
 Murrieta, California 92563, USA

Device Name: HI-TORQUE® EXTRA S'PORT™ 0.014'' Guide Wire

Device Classification: Class III

GMDN Code: 35094 – Cardiac catheter guidewire, single-use

Classification Rationale: The following Annex IX definition(s) apply to the HI-TORQUE® EXTRA S'PORT™ 0.014'' Guide Wire for purposes of classifications: Per Rule 6, Annex IX, all surgically invasive devices intended for transient use are in Class IIa unless they are: Intended specifically to control, diagnose, monitor, or correct a defect of the heart or of the central circulatory system through direct contact with these parts of the body, in which case they are in Class III.

Authorized European Representative: Abbott Vascular International BVBA
 Park Lane
 Culliganlaan 2B
 1831 Diegem, Belgium

Model Numbers: HI-TORQUE® EXTRA S'PORT™ 0.014'' Guide Wire

FG#	Tip Shape
2222XM	Straight Tip Shape
2222XMJ	'J' Tip Shape
2223XM	Straight Tip Shape
2223XMJ	'J' Tip Shape
22225M-DCA	Straight Tip Shape
22235M-DCA	Straight Tip Shape
11125M-DCA	Straight Tip Shape
11135M-DCA	Straight Tip Shape

I, the undersigned, hereby declare that the medical devices specified above conform with the applicable *Essential Requirements* listed in Annex I and Annex II Part 4 of EC Council Directive 93/42/EEC.

Directive 2006/42/EC on Machinery and directive 1989/686/EC on Personal Protective Equipment do not apply.

This declaration is supported by an EC quality system and design examination approval listed below.

Quality Management System - ISO 13485:2003, Certificate: FM72377

EC Design Examination Certificate: CE 01497

Full Quality Assurance Certificate: CE 510108

Notified Body: British Standards Institution, a notified body authorized by the United Kingdom Competent Authority, Notified Body Identification Number 0086.

British Standards Institution
Kitemark Court
Davy Avenue
Knowlhill
Milton Keynes
MK5 8PP
United Kingdom

This Declaration of Conformity expires upon revision or with the revision or expiration of any of the supporting certificates listed above.

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

By: Susan Slane Date: 8/15/14

Susan Slane
Divisional VP, Global Quality and Compliance
Abbott Vascular

Place of issue: Temecula Date of issue: 8/15/14

Effective Date: 8/15/14