Abbott Vascular

3200 Lakeside Drive
Santa Clara, CA 95054

Tel 408-845-3000
Fax 408-845-3333

DECLARATION OF CONFORMITY

Manufacturer:	Abbott Vascular
Address:	3200 Lakeside Drive Santa Clara, California 95054, USA
Additional Manufacturing Sites:	26531 Ynez Road Temecula, California 92591, USA
Device Name:	**Absolute Pro Peripheral Self-Expanding Stent System**
Device Classification:	Class IIb
GMDN Code:	47932 - Multiple peripheral artery stent
Classification Rationale:	The following Annex IX definition(s) apply to the Absolute Pro Peripheral Stent System for purposes of classification: Per Rule 8, Annex IX, all implantable devices and long-term surgically invasive devices are in Class IIb unless they are intended: to be used in direct contact with the heart, the central circulatory system, or the central nervous system, in which case they are in Class III.
Authorized European Representative:	Abbott Vascular International BVBA Park Lane Culliganlaan 2B 1831 Diegem, Belgium

Abbott Vascular

3200 Lakeside Drive
Santa Clara, CA 95054

Tel 408-845-3000
Fax 408-845-3333

Model Numbers: **Absolute Pro Peripheral Self-Expanding Stent System**

Description (stent diameter, system length)	**Part Number**	**Suffix-XXX** (stent length, mm)					
ABSOLUTE PRO 5mm, 80cm	1011914	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 6mm, 80cm	1011915	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 7mm, 80cm	1011916	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 8mm, 80cm	1011917	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 9mm, 80cm	1011918	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 10mm, 80cm	1011919	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 5mm, 135cm	1011920	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 6mm, 135cm	1011921	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 7mm, 135cm	1011922	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 8mm, 135cm	1011923	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 9mm, 135cm	1011924	-020	-030	-040	-060	-080	-100
ABSOLUTE PRO 10mm, 135cm	1011925	-020	-030	-040	-060	-080	-100

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 3 of EC Council Directive 93/42/EEC.

This declaration is supported by an EC quality system (Annex II) under the supervision of the British Standards Institution, a notified body authorized by the United Kingdom Competent Authority (EC Quality System Certificate Number: CE 510108).

British Standards Institution – Notified Body Identification Number 0086
Kitemark Court
Davy Avenue
Knowlhill
Milton Keynes
MK5 8PP
United Kingdom

By: Sg Slane Date: 4/18/12

Susan Slane
Divisional VP, Global Quality and Compliance
Abbott Vascular