

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

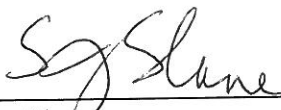
Model Numbers: Absolute Pro Peripheral Self-Expanding Stent System

Description (stent diameter, system length)	Part Number	Suffix-XXX (stent length, mm)						
		-020	-030	-040	-060	-080	-100	
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
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Davy Avenue
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Milton Keynes
MK5 8PP
United Kingdom

By:  Date: 4/18/12
Susan Slane
Divisional VP, Global Quality and Compliance
Abbott Vascular