

## Declaration of Conformity

The manufacturer:

St. Jude Medical (Quality System certificate I1 14 05 14607 184)  
Cardiac Rhythm Management Division  
15900 Valley View Court  
Sylmar  
California 91342  
U.S.A.

Fulfills the requirements of Annex 2 of the European Union's Active Implantable Medical Devices Directive, AIMDD, 90/385/EEC / corresponding national legislation.

The Authorized EC Representative is:

St. Jude Medical  
Coordination Center BVBA  
The Corporate Village  
Da Vincilaan 11 Box F1  
1935 Zaventem  
BELGIUM

We, representing the manufacturer, hereby certify that the product(s) listed in the **Attachment** are covered by EC-certificate number **I7 14 05 14607 174** and fulfill(s) applicable requirements including CE-marking and the Essential Requirements of AIMDD 90/385/EEC / corresponding national legislation. The above certificates have been issued by the Notified Body 0123, TÜV SÜD Product Service, Ridlerstr. 65, 80339 Munich, Germany.

We also certify that applicable product(s) listed in the **Attachment** fulfill(s) applicable requirements include CE-marking and other Requirements of the Radio and Tele Terminal Equipment Directive, R&TTED, 1999/5/EC / corresponding national legislation.

Place: Sylmar, USA

Date: 19-Feb-2015



Elisabeth Neely  
Vice President, Global Regulatory Affairs  
St. Jude Medical CRMD

## Attachment to Declaration of Conformity

The following product(s) is/are approved under EC-Certificate number **I7 14 05 14607 174**.

<b>Device Name</b>	<b>Model No. or Version No.</b>	<b>GMDN Code</b>
Accent MRI	PM1224	47267
	PM2224	47265
	PM1124	47267
	PM2124	47265
Accent ST MRI	PM1226	47267
	PM2226	47265
	PM1126	47267
	PM2126	47265
Tendril MRI	LPA1200M	35223
SJM MRI Activator Handheld Device	EX4000	47805