

Number: 3903009TD09

EU Technical Documentation Assessment Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter II and III

Manufacturer:

Biosense Webster, Inc

31 Technology Drive, Suite 200

Irvine, CA 92618

USA

SRN ID.: US-MF-000014219

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

0344

Supplement to certificate: 3826381CN

**Authorized Representative: Biosense Webster, A Division of Johnson & Johnson Medical NV/SA,
Leonardo da Vincilaan 15, 1831 Diegem, Belgium**

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer/ authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.M. McKenzie
Principal Certification Manager

First Issued: **23-11-2023**

Date: **23-11-2023**

Expiry date: **01-11-2028**

© Integral publication of this certificate and adjoining reports is allowed

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra-product-safety.com Company registration 09085396

Number: 3903009TD09

EU Technical Documentation Assessment Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter II and III

This certificate covers the following device(s) / groups of device(s):

Class III	
<p>Basic UDI-DI: 08468350a0022EX</p> <p>Device Name: THERMOCOOL™ Catheters Type: C020301 - Cardiac Tissue Ablation Electrocatheters, Radiofrequency Model: CELSIUS™ THERMOCOOL™ Catheter D-1189-XX-S (XX= 01-04) NAVISTAR™ THERMOCOOL™ Catheter D-1197-XX-S (XX= 14-18) EZ STEER™ THERMOCOOL™ Bi-Directional Catheter D- 1294-XX-S (XX= 01-05) EZ STEER™ THERMOCOOL™ NAV Bi-Directional Catheter D-1292-XX-S (XX=-01-05)</p>	<p><i>Intended Purpose: The THERMOCOOL™ Catheters are intended for use in cardiac electrophysiological mapping (stimulation and recording) and, when used in conjunction with an RF generator, for cardiac ablation.</i></p>
<p>Basic UDI : 08468350a0021EV</p> <p>Device Name: THERMOCOOL™ RMT Catheters Type: C020301 - Cardiac Tissue Ablation Electrocatheters, Radiofrequency Model: NAVISTAR™ RMT THERMOCOOL™ Catheter D-1266- 01-S CELSIUS™ RMT THERMOCOOL™ Catheter D-1302- 01-S</p>	<p><i>Intended Purpose: The THERMOCOOL™ RMT Catheters are intended for use in cardiac electrophysiological mapping (stimulation and recording) and, when used in conjunction with an RF generator, for cardiac ablation.</i></p>

First Issued: 23-11-2023

Date: 23-11-2023

Expiry date: 01-11-2028

© Integral publication of this certificate and adjoining reports is allowed

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra-product-safety.com Company registration 09085396

Number: 3903009TD09

EU Technical Documentation Assessment Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter II and III

Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Revision	Date of Issue certificate	Certification Notice Reference	Action
0	23-11-2023	3826381CN35	First issue

First Issued: 23-11-2023

Date: 23-11-2023

Expiry date: 01-11-2028

© Integral publication of this certificate and adjoining reports is allowed

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra-product-safety.com Company registration 09085396