EU Certificate

Technical Documentation Assessment REGULATION (EU) 2017/745 on Medical Devices, Annex IX Chapter II, Section 4



Registration No.:

IZ 2320977-1

Manufacturer:

SP Medical A/S

Møllevej 1 4653 Karise Denmark

EUDAMED Single Registration No.:

DK-MF-000019782

General product group

name:

Products of class III:

C040101 - CORONARY ARTERY GUIDEWIRES

CORONARY ARTERY DIAGNOSTIC GUIDEWIRES

C040102 - CORONARY ARTERY GUIDEWIRES

CORONARY ARTERY THERAPEUTIC GUIDEWIRES

C040201 - PERIPHERAL VASCULAR GUIDEWIRES

PERIPHERAL VASCULAR DIAGNOSTIC GUIDEWIRES

C040202 - PERIPHERAL VASCULAR GUIDEWIRES

PERIPHERAL VASCULAR THERAPEUTIC GUIDEWIRES

Models and types:

- Seldinger - Straight

- Seldinger - J-tip

- Seldinger - Double ended - Seldinger Movable - Straight

- Seldinger Movable - J-tip

Basic UDI-DI:

5710086-GWAC

The Notified Body hereby declares that the requirements of Annex IX, Chapter II, Section 4 of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and maintains a technical documentation defined by Annex IX, Chapter II, Section 4 of the aforementioned regulation. In addition to this certificate an EU quality management system certificate is required before placing the listed products on the market.

Report No.:

84952932-10

Effective date:

2022-03-29

Expiry date:

2027-03-28

Issue date:

2022-03-29





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TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

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Registration No.:

IZ 2320977-1

Manufacturer:

SP Medical A/S

Møllevej 1 4653 Karise Denmark

Intended use:

General intravascular and coronary arterial use to aid in the selective placement

of interventional devices during diagnostic and/or therapeutic procedures.

Product name:

Accoat Guide Wires

Authorised

N/A

representative(s):

Certificate history		
Revision:	Description:	Issue date:
1	Initial revision.	2022-03-29

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