

Annex to the EC Certificate No. 50736-16-07

Revision status: 1

Valid from 2021-01-25 to 2023-03-26

Devices/device categories included in the certificate:

Class II a:

- E-wire Guide Wire
- E-expand Stent Graft Balloon Catheter

Class II b:

- FlowLine Bipore ePTFE Vascular graft
- E-liac Stent Graft System
- E-ventus BX Peripheral Stent Graft System

Class III:

- Textile vascular grafts: FlowWeave, FlowNit, FlowWeave Bioseal, FlowNit Bioseal
- FlowLine Bipore Heparin ePTFE Vascular graft
- E-vita thoracic 3G Stent Graft System
- E-vita open plus Stent Graft System
- E-tegra Stent Graft System
- E-nside TAAA Multibranch Stent Graft System
- E-nya Thoracic Stentgraft System
- E-vita OPEN NEO

For the placing on the market of class III devices covered by this certificate an EC design-examination certificate according to directive 93/42/EEC annex II (4) is required.



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