DECLARATION OF CONFORMITY

Legal Manufacturer: Stryker Neurovascular
47900 Bayside Parkway
Fremont, CA 94538
USA

Manufacturing Site(s): Stryker Neurovascular
Business and Technology Park
Model Farm Road
Cork, Ireland

European Representative: Stryker European Operations B.V.
Herikerbergweg 110
Amsterdam
1101 CM
Netherlands

Product: Excelsior® 1018® Reinforced Microcatheter
Design Dossier NV00017604

Product Category: Microcatheters

Classification: Class III, Rule 7 according to Annex IX of the MDD

We declare that the products identified above are in conformity with all relevant provisions of the Council Directive 93/42/EEC of 14 June 1993 concerning Medical Devices. This declaration is made under Annex I of this directive. All supporting information is retained under the control of the Legal Manufacturer.
Notified Body:

BSI
Kitemark Court, Davy Avenue,
Knowlhill, Milton Keynes, MK5 8PP,
United Kingdom

Identification number 0086

Quality Certificate: BSI EN ISO 13485 Certificate # MD 638383
First issued 23 November 2015

EC Certificate: BSI Certificate # CE 635352
First issued 01 June 2015

DE Certificate: BSI Certificate # CE 635354
First issued 01 June 2015

This Declaration of Conformity is issued under the sole responsibility of Stryker Neurovascular.

Signature, Date of Issue:

Signature: Katherine Mack  Date: Feb 27, 2017
Name: Katherine Mack
Senior Director of Quality
Stryker Neurovascular, Fremont, CA

Signature: Jennifer Mateus  Date: Feb 20, 2017
Name: Jennifer Mateus
Senior Director of Regulatory Affairs
Stryker Neurovascular, Fremont, CA
Attachment to the EC Declaration of Conformity for Excelsior® 1018® Reinforced Microcatheters

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<th>Description</th>
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<tr>
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<td>Excelsior® 1018® Reinforced Microcatheter, 150 cm/ 6 cm, 1-tip</td>
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<td>M0031441890</td>
<td>Excelsior® 1018® Reinforced Microcatheter, 150 cm/ 6 cm, 2-tip</td>
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