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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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ZLG-BS-244.10.08



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)

(Devices in Class IIa, IIb or III)

No. G1 058654 0011 Rev. 01

Manufacturer:

SpineVision S.A.S.

Bâtiment E

10 Rue de la Renaissance

92160 Antony

FRANCE

Product Category(ies): Spinal implants and instruments

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G1058654 0011 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:G10586540011Rev.01)

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2024-05-26

Date,

2021-05-12

Christoph Dicks

Head of Certification/Notified Body