EC Certificate Full Quality Assurance System: Certificate CH19/0996

The management system of

Harald Nordin SA

Zone Industrielle La Foge,
CH - 1816 Chailly-sur-Montreux, Switzerland

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

• Glass Fiber Post for endodontic application
• Fiber Carbon Post for endodontic application
• Peeso & Gates, endo reamers for endodontic applications
• Cement for restorative dentistry (Dropsin)
• Metallic screw post: Gold, Titanium, Stein still and Ancorex for endodontic applications
• Titanium dentine retention pins
(Devices intended to support the dental restoration)

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 26 June 2020 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 4 July 2008
and first certified by SGS Belgium NV since 16 January 2020

Certification is based on reports numbered CH/GE 3302011

Authorised by

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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