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

SGS Belgium NV, Notified Body 1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 **f** +32 (0)3 545-48-49 www.sgs.com

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

Harald Nordin SA
Zone Industrielle La Foge
Route des Châtaigniers 12
1816 Chailly-sur-Montreux
Switzerland

May 23rd, 2024

Confirmation Letter Reference: CLNB1639 – CH/GE 3205780

To whom it may concern,

Confirmation of receipt of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

This letter confirms that, SGS Belgium NV, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 1639 on NANDO, has received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the following manufacturer

Harald Nordin SA
Zone Industrielle La Foge
Route des Châtaigniers 12
1816 Chailly-sur-Montreux
Switzerland
SRN Number CH-MF-000016069

Authorized representatives
WMDE BV
Bergerweg 18
6085 AT Horn
The Netherlands
NL-AR-000002062

SL MEDREP SAS
22 Avenue de la Gare
25160 Labergement Ste Marie
France
SRN FR-AR-000014715

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below . Table 1 identifies the devices which an MDR application has been received, written agreement concluded and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive. Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but the NB has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that:

- The manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry;
- The certificates expired after 26th May 2021 by course of time and were valid at the date of their expiry neither having been suspended nor withdrawn.

The transition timelines that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120.3 of MDR (as amended by EU 2023/607), are shown below:

- 26th May 2026 for Class III custom-made implantable devices
- 31st December 2027 for Class III devices and Class IIb implantable devices excluding Well-established technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31st December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- 31st December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

On behalf of the Notified Body SGS Belgium NV NB1639,

Virginie SILORET
Global Medical Device Certification Manager
Email: Virginie.siloret@sgs.com
Phone: +41 22 739 98 58

Table 1: Devices covered by this letter and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive**:**

Device name or Basic UDI-DI	**MDR Device classification**	**MDD Device name**	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	**MDD/AIMDD Certificate Reference(s)** of the devices under MDR application, and the NB Identification
Metal Screw Posts 764034297SCREWPOST2U	Class IIa	Metal Screw Posts	N/A	Certificate CH19/0996/NB1639
Dental Reamers 764034297REAMERSQN	Class IIa	Dental Reamers	N/A	Certificate CH19/0996/NB1639
Glass Fiber Posts 764034297GLASSFIBERCP	Class IIa	Glass Fiber Posts	N/A	Certificate CH19/0996/NB1639
Dentine Pins & Drills 764034297DENTINEPINS8R	Class IIa	Dentine Pins & Drills	N/A	Certificate CH19/0996/NB1639

Table 2: Devices covered by this letter and for which the NB is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under MDR application)	**MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)**	**If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device**	**MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification**
N/A	N/A	N/A	N/A

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2024/05/23	Version 1	Initial issue