EC Certificate Full Quality Assurance System: Certificate CH19/0963

The management system of

Intensiv SA

Via al Molino 107
CH-6926 Montagnola

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

Rotating diamond coated instruments for conservative and restorative dentistry
Oscillating diamond coated instruments for conservative, restorative and orthodontic dentistry

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 09 February 2021 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 3. Certified since 11 March 1998

Certification is based on reports numbered CH/GE 3205732

Authorised by

Global Medical Devices Certification Manager

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

Intensiv SA
Via al Molino 107,
6926 Montagnola
Switzerland

August 20th, 2023

Confirmation Letter Reference: CLNB1639 – CH/GE 3205732

To whom it may concern,

Confirmation of receipt of a formal application and conclusion of written agreement 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

Intensiv SA
Via al Molino 107,
6926 Montagnola
Switzerland
SRN Number (if available): CH-MF-000036274

Intensiv Trade GmbH
Hoheneichstrasse 12
D-75210 Keltern
Germany
The SRN Number: DE-AR-000013123

The devices covered by the formal application and the written agreement mentioned above are shown at the end of this letter.

In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired prior to the publication of the Regulation EU 2023/607 on 15th March 2023, this letter also confirms that:

- The manufacturer submitted the MDR application and signed the written agreement 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 1639,

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

Devices covered by this letter:

Device name / Basic UDI-DI	**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
Rotating diamond coated instruments for conservative and restorative dentistry/++D959DIAROTINSRASF	Class IIa	N/A	Certificate CH19/0963 / NB1639

Device name / Basic UDI-DI	**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
++D959DIAROTINSFGRP			
Oscillating diamond coated instruments for conservative, restorative and orthodontic dentistry/++D959DIAOSCINSOSLQ ++D959DIAOSCINSFILCA	Class IIa	N/A	Certificate CH19/0963 / NB1639

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2023/08/20	Version 1	Initial issue
2024/01/15	Version 2	Adding SRN Number CH-MF-000036274 Adding references ++D959DIAROTINSFGRP Adding references ++D959DIAOSCINSFILCA
2024/01/24	Version 3	Correction typo in the address “Hoheneichstrasse” instead of “Hocheneichstrasse”