

ITENA CLINICAL
Central Parc, Batiment B 97 Allée De La Louve
93420 Villepinte Fr - France
09/07/2024

Notified Body Confirmation Letter
Reference: 145501

To whom it may concern,

Confirmation 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, ICIM SPA, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 0425 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:

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The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below. Table 1 identifies the devices for 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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively, by the 20 Mar 2023 for the relevant devices.

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 120.3c of MDR (as amended by (EU) 2023/607), are shown below:

- 26 May 2026 for Class III custom-made implantable devices
- 31 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)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- 31 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,
ICIM SPA
Piazza Don Enrico Mapelli, 75
2099 Sesto San Giovanni MI

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:

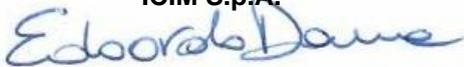
Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Dental glass fiber post for endodontic restoration - DENTOCLIC - Glass fiber posts	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Dental titanium posts for endodontic restoration - DENTOCLIC - Titanium and stainless steel posts	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Dental drills for endodontic restoration - DENTOCLIC - Drills and Reamers	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Polymer-based filling restorative materials - REFLECTYS and REFLECTYS FLOW	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Glass fiber ceramic disk and blocs - NUMERYS GF	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Hybrid ceramic disk and blocs - NUMERYS HC	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Bioceramic reparative cement - MTA BIOREP	Class IIb excluding Class IIb implantable	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Bioceramic root canal filler - MTA BIOSEAL	Class IIb excluding Class IIb implantable	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Temporary crowns and bridges - DENTOCROWN HD	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Temporary cement - PROVITEMP and DENTOTEMP	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
Dentocore - Restorative Cement	Class IIa	N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425

Device name or Basic UDI-DI (under MDR application)	MDR classification proposed by the manufacturer and verified at the pre-application stage)	Device (as proposed by the manufacturer and verified at the pre-application stage)	If the 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
Dentocore Body - Restorative Cement	Class IIa		N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
PREVENT SEAL Self-etching light cured pit & fissure sealant	Class IIa		N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
DENTAL COMPOSITE RESIN	Class IIa		N/A	Certificate N. MED-004026-00; Letter N. BO/2022/003657/LG-md NB ICIM SPA CE 0425
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Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
08/05/2023	REV.00	Initial issue
19/03/2024	REV.01	Addition of device: PREVENT SEAL Self-etching light cured pit & fissure sealant
04/06/2024	REV.02	Addition of device: DENTAL COMPOSITE RESIN
09/07/2024	REV.03	Added Clearer specification for the device DENTAL COMPOSITE RESIN

Mr. Edoardo Dossena
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Miss. Flavia Lepore
Sales Director
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