



# Ente Certificazione Macchine

Notified Body n. 1282 - Testing Laboratory – Nr. 121697 PJLA

Authorized Training Body n. 6737 - Inspection Body

---



CONFIRMATION LETTER IN THE FRAMEWORK OF  
REGULATION EU 2023/607  
FOR "LEGACY" DEVICES  
ACCORDING TO DIRECTIVE 93/42/EEC

**REFERENCE N° 18/F**

---

ENTE CERTIFICAZIONE MACCHINE SRL

Via Ca' Bella, 243 – Loc. Castello di Serravalle – 40053 Valsamoggia (BO) – Italy

☎ 051.6705141 📠 051.6705156 ✉ info@entecerma.it [www.entecerma.it](http://www.entecerma.it)



ENTE CERTIFICAZIONE MACCHINE

---

---

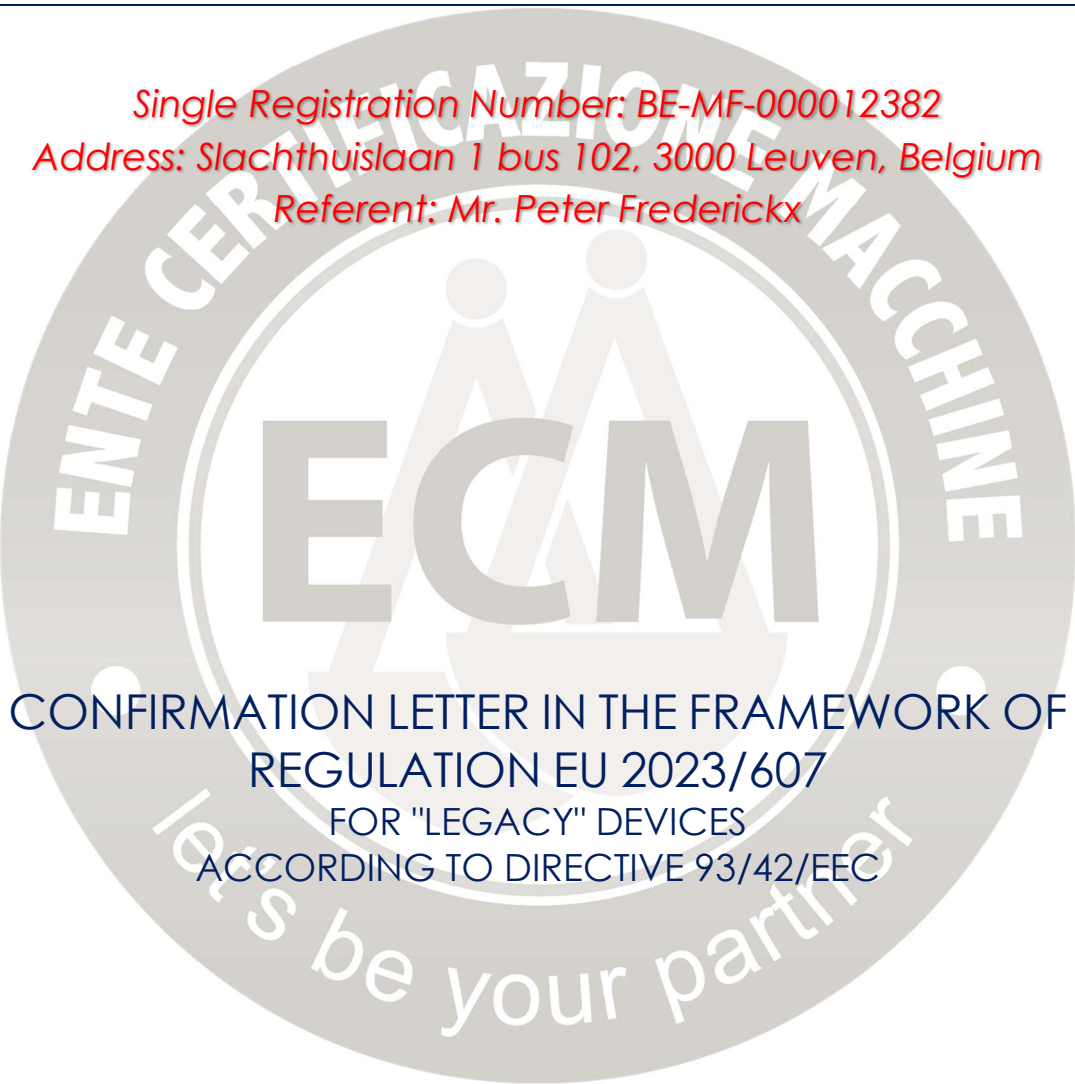
# MedXpress Pro

---

Single Registration Number: BE-MF-000012382

Address: Slachthuislaan 1 bus 102, 3000 Leuven, Belgium

Referent: Mr. Peter Frederickx



CONFIRMATION LETTER IN THE FRAMEWORK OF  
REGULATION EU 2023/607  
FOR "LEGACY" DEVICES  
ACCORDING TO DIRECTIVE 93/42/EEC



## ENTE CERTIFICAZIONE MACCHINE

### Confirmation of the status of a formal application, written agreement, and appropriate surveillance activity 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

**Ente Certificazione Macchine srl (ECM)**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **1282 on NANDO**, confirms to have received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and to have signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with **MedXpress Pro**.

In addition, this letter confirms that **ECM**, where relevant, has signed a written agreement with **MedXpress Pro** governing transfer of the surveillance activity in accordance with Article 120, paragraph 3e of MDR as amended by Regulation (EU) 2023/607.

The devices covered by the formal application and the written agreements mentioned above are identified in the Tables below. **Table 1** identifies devices for which an MDR application has been received, written agreement concluded, and for which **ECM** is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive. **Table 2** identifies devices for which an MDR application has been received and a written agreement concluded, but **ECM** 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 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 120.3 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)

**ENTE CERTIFICAZIONE MACCHINE SRL**

LUCA BEDONNI



## ENTE CERTIFICAZIONE MACCHINE

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

Device name	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application
EZCOVER® probe covers	<input type="checkbox"/> Class IIb device <input type="checkbox"/> Class IIa device <input checked="" type="checkbox"/> Class I devices placed on the market in sterile condition <input type="checkbox"/> Class I devices with a measuring function <input type="checkbox"/> Class I devices that qualify as re-usable surgical instruments	<input type="checkbox"/> MDD/AIMDD device identification: Device name <input checked="" type="checkbox"/> Not applicable	<input checked="" type="checkbox"/> MDD/AIMDD Certificate ECM20MDD010 rev.1 issued by Ente Certificazione Macchine srl, NB number 1282 <input type="checkbox"/> Not applicable

**Table 2 Devices covered by this letter and for which ECM is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive: N.A.**