Hammarplast Medical AB
Kartåsgatan 8, SE-531 40 Lidköping
Sweden

28 February 2024

Notified Body Confirmation Letter
Reference: *MDD Cert No. 41319092-05 - CN00413-01*

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, Intertek Medical Notified Body AB, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 2862 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:

Hammarplast Medical AB
Kartåsgatan 8, SE-531 40 Lidköping
Sweden

SRN Number (if available): SE-MF-000009420

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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that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,

Brian Mather
Certification Manager
Intertek Medical Notified Body AB

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
Medicine measure	I(m)	Listed as 01250-01254	41319092-05; NB0413
Medicine measure	I(m)	Listed as 01250-01254	41319092-05; NB0413
Medicine measure	I(m)	Listed as 01250-01254	41319092-05; NB0413
Medicine measure	I(m)	Listed as 01250-01254	41319092-05; NB0413
Medicine measure	I(m)	Listed as 01250-01254	41319092-05; NB0413
Medicine measure ECO	I(m)	Listed as 01350-01354	41319092-05; NB0413
Medicine measure ECO	I(m)	Listed as 01350-01354	41319092-05; NB0413
Medicine measure ECO	I(m)	Listed as 01350-01354	41319092-05; NB0413
Medicine measure ECO	I(m)	Listed as 01350-01354	41319092-05; NB0413
Medicine measure ECO	I(m)	Listed as 01350-01354	41319092-05; NB0413
Medicine measure ECO+	I(m)	Listed as 02250-02254	41319092-05; NB0413
Medicine measure ECO+	I(m)	Listed as 02250-02254	41319092-05; NB0413
Medicine measure ECO+	I(m)	Listed as 02250-02254	41319092-05; NB0413
Medicine measure ECO+	I(m)	Listed as 02250-02254	41319092-05; NB0413
Medicine measure ECO+	I(m)	Listed as 02250-02254	41319092-05; NB0413
Medicine measure	I(m)	Listed as 10300-10304	41319092-05; NB0413
Medicine measure	I(m)	Listed as 10300-10304	41319092-05; NB0413
Medicine measure	I(m)	Listed as 10300-10304	41319092-05; NB0413
Medicine measure	I(m)	Listed as 10300-10304	41319092-05; NB0413
Medicine measure	I(m)	Listed as 10300-10304	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413

Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Disposable tourniquet cuffs	I(s)	N/A	41319092-05; NB0413
Medicine measure JONI	I(m)	Listed as J10823-J10833	41319092-05; NB0413
Medicine measure JONI	I(m)	Listed as J10823-J10833	41319092-05; NB0413
Medicine measure JONI	I(m)	Listed as J10823-J10833	41319092-05; NB0413
Medicine measure JONI	I(m)	Listed as J10823-J10833	41319092-05; NB0413

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

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action