



**Notified Body Confirmation Letter Reference: C682954**

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, DNV Product Assurance AS, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number NB 2460 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:

MEDICLUS CO., LTD.

No. 1210, 134, Gongdan-ro, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do, Korea

SRN Number (if available): KR-MF-000022584

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issued under Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD certificate expiry; or provided evidence that a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

Place and date:  
Høvik, 2024/09/24



For the Issuing office:  
DNV Product Assurance AS – Notified Body 2460  
Veritasveien 1, 1363 Høvik, Norway

**Rajesh Kumar Chellappan**  
Management Representative

Lack of fulfilment of conditions as set out in the Certification Agreement may render this letter invalid.

NOTIFIED BODY 2460: DNV Product Assurance AS, Veritasveien 1, 1363 Høvik, Norway, Tel +47 67 57 88 00, www.dnv.com



- 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)

**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:**

<b>Device name and Basic UDI-DI (under MDR application)</b>	<b>MDR Device classification (as proposed by the manufacturer and verified at the quotation request review stage)</b>	<b>If the MDR device is a substitute device, identification of the corresponding MDD device</b>	<b>MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification</b>
Bioceramic Root Canal Sealer (One-Fil)  / Basic UDI DI: 880951834EN03JV	Ila	Dental root canal sealer (One-Fil) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)
Dental Temporary Filling Material (Once-Fil Flow)  / Basic UDI DI: 880951834TP02NE	Ila	N/A	0068/QCO-DM-312-2021, MTIC (CE 0068)
Any-Com Flow Dental Restoration Procedure Pack (Any-Com Flow)  / Basic UDI DI: 880951834RS03NH	Ila	Dental Composite Resin (Any-Com Flow) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)
Light Cured Temporary Crown and Bridge Resin (Ezi-Crown)  / Basic UDI DI: 880951834TP05NL	Ila	Dental temporary filling material (Ezi-Crown) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)
Dental Temporary Cement (Any-Temp NE)  / Basic UDI DI: 880951834TP03NG	Ila	Dental Temporary Cement (Any-Temp) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)
Dual Cured Dental Composite Resin (Any-Core)  / Basic UDI DI: 880951834RS02NF	Ila	Dental composite resin (Any-Core) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)
Resin-based Root Canal Sealer (Any-Seal RC)  / Basic UDI DI: 880951834EN05JZ	Ila	Dental root canal sealer (Any-Seal RC) (Name change only)	0068/QCO-DM-312-2021, MTIC (CE 0068)



**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:**

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

**Confirmation Letter Revision History**

<b>Date</b>	<b>NB internal reference traceable to each version of the letter</b>	<b>Action</b>
2024/09/24	C682954	Initial issue

**Lack of fulfilment of conditions**

The following may render this letter of confirmation invalid:

- Lack of compliance to the requirements of Regulation (EU) 2023/607.
- Significant changes to design or intended purpose of the devices.
- Changes in the quality system affecting production.
- Periodical audits not held within the timeframe.