

DEKRA Certification GmbH – Handwerkstraße 15 – D-70565 Stuttgart

Joline GmbH & Co. KG  
Mr. Dr. Marian Wenzel  
Neue Rottenburger Str. 50  
72379 Hechingen  
Germany

**DEKRA Certification GmbH**

Handwerkstraße 15  
D-70565 Stuttgart

Contact Stephanie Donner  
Phone -  
Fax +49.711.7861-2615  
Email stephanie.donner@dekra.com

Headquarters  
Phone +49.711.7861-2566  
Fax +49.711.7861-2615

Date 2023-11-29

**Subject: Notified Body Confirmation Letter**

**Our reference: 50565-CoL-01, Rev.0**

**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 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices**

Dear Dr. Wenzel,

This letter confirms that, DEKRA Certification GmbH, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 0124 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:

Joline GmbH & Co. KG  
Neue Rottenburger Str. 50  
72379 Hechingen  
Deutschland

SRN Number: DE-MF-000005494

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables 1 and 2 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 MDD. 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 MDD.

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issued under Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the

DEKRA Certification GmbH  
Handwerkstraße 15  
D-70565 Stuttgart  
www.dekra-certification.de/  
medizinprodukte

Registered at the local court of Stuttgart  
under HRB Nr. 17662  
Bank: Commerzbank AG  
IBAN: DE76 6008 0000 0901 4949 00  
BIC: DRES DE FF 600  
Ust.-ID-Nr. DE 811 976 119

Managing director:  
Dr. Rolf Krökel

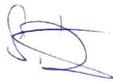
date of MDD certificate expiry; or provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively, by the 20 Mar 2023 for the relevant 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
- 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,

Stephanie Donner  
2023-11-29



Digitally signed by Stephanie  
Donner  
Date: 2023-11-29 15:41:00+01:00

Enclosures:

Confirmation Letter Annex

**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>Product or product group identification acc. to MDD - certificate</b>	<b>MDD Device classification</b>	<b>MDD Certificate and Certificate Annex No. with revision</b>	<b>MDR Application ID</b>
<b>MD 0106</b> <b>Kyphoplasty Systems ALLEVO</b> <ul style="list-style-type: none"> <li>- Kits</li> <li>- Individual Instruments</li> </ul>	Class IIa	Certificate 50565-16-06 Annex Rev. 2	A22051293
<b>MD 0102</b> <b>Dialysis Catheter ST</b> <ul style="list-style-type: none"> <li>- Kits</li> <li>- Catheter</li> </ul>	Class IIa	Certificate 50565-16-06 Annex Rev. 2	A22071097
<b>MD 0203</b> <b>Dialysis Catheter PU-LT</b> <ul style="list-style-type: none"> <li>- Kits</li> <li>- Catheter</li> </ul>	Class III	Certificate 50565-16-06 Annex Rev. 2	A22071097
<b>MD 0203</b> <b>Dialysis Catheter Silicone LT</b> <ul style="list-style-type: none"> <li>- Kits</li> <li>- Catheter</li> </ul>	Class III	Certificate 50565-16-06 Annex Rev. 2	A22071097
<b>MD0101</b> <b>Miniclamp</b>	Class Is	Certificate 50565-17-05 Annex Rev. 0	A22051293
<b>MD 0106</b> <b>Mixer</b>	Class Is	Certificate 50565-17-05 Annex Rev. 0	A21091017