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Medcare Saglik Urunleri San. ve Tic. A.S.  
Fatih Mah. Camlik Cad. No:54 35414 Gaziemir/İzmir TÜRKİYE

**Notified Body Confirmation Letter Reference: CERBO0291222**

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, Kiwa Cermet Italia, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 0476 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:

Medcare Saglik Urunleri San. ve Tic. A.S.  
Fatih Mah. Camlik Cad. No:54 35414 Gaziemir/İzmir TÜRKİYE  
**SRN Number (if available): TR-MF-000023643**

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issued under Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD certificate expiry; or provided evidence that a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



**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	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
<b>Sterile Disposable Gowns -Standard Surgical Gown</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input type="checkbox"/> Same <input checked="" type="checkbox"/> Substitute  MDD Product Name: Sterile Disposable Gowns	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Gowns -Reinforced Surgical Gown</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input type="checkbox"/> Same <input checked="" type="checkbox"/> Substitute  MDD Product Name: Sterile Disposable Gowns	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Drapes (e.g. Laparotomy Drape)</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input checked="" type="checkbox"/> Same <input type="checkbox"/> Substitute	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Drape Packs (e.g. General Surgery Drape Pack)</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input checked="" type="checkbox"/> Same <input type="checkbox"/> Substitute	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Equipment Covers - Microscope cover</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input type="checkbox"/> Same <input checked="" type="checkbox"/> Substitute  MDD Product Name: Sterile Disposable Equipment Covers	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Equipment Covers -Cardboard camera cover</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD	Certificate 1984-MDD-20-649;



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	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
		<input type="checkbox"/> Same <input checked="" type="checkbox"/> Substitute  MDD Product Name: Sterile Disposable Equipment Covers	NB:Kiwa Belgelendirme Hizmetleri-1984
<b>Sterile Disposable Equipment Covers -Fluoroscopy cover</b>	Class I devices placed on the market in sterile condition	Identification of the corresponding device under MDD  <input type="checkbox"/> Same <input checked="" type="checkbox"/> Substitute  MDD Product Name: Sterile Disposable Equipment Covers	Certificate 1984-MDD-20-649; NB:Kiwa Belgelendirme Hizmetleri-1984

**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	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification

#### Confirmation Letter Revision History

Rev. Rev.	Date Date	Action Azione
00	2023/12/20	Initial issue: Sterile Disposable Gowns -Standard Surgical Gown, Sterile Disposable Gowns -Reinforced Surgical Gown, Sterile Disposable Drapes (e.g. Laparotomy Drape), Sterile Disposable Drape Packs (e.g. General Surgery Drape Pack), Sterile Disposable Equipment Covers - Microscope cover, Sterile Disposable Equipment Covers -Cardboard camera cover,
01	2024/09/26	<b>The MDD appropriate surveillance offer signed.</b>

For further information on the content of the letter or verification of the validity of the letter please contact [medical@kiwa.com](mailto:medical@kiwa.com) or phone at +39.051.4593.111

