



Reg. Numero / <i>Reg. Number</i>	MED 23002	Revisione / <i>Revision</i>	10
Primo rilascio / <i>First issue date</i>	2003-09-17	Valido da / <i>Valid from</i>	2018-09-14
Scadenza / <i>Valid until</i>	2023-09-16	Ultima modifica / <i>Last change date</i>	2019-03-04

Pagina / Page 1 di / of 5

## Certificato CE del Sistema di Garanzia della Qualità *EC Quality Assurance System Certificate*

Si certifica che, sulla base dei risultati degli audit effettuati, il Sistema completo di garanzia di Qualità dell'Organizzazione/ *We certify that, on the basis of the audits carried out, the full Quality Assurance System of the Organization:*

### **ADRIA S.r.l. con Unico Socio**

**Sede Operativa / Operational headquarter:**

Via San Francesco d'Assisi, 15  
46020 Pegognaga, MN - Italia

**Sede Legale / Registered Headquarter**

Via Modena, 46  
40017 San Giovanni in Persiceto, BO - Italia

è conforme ai requisiti applicabili della Direttiva 93/42/CEE e successive modifiche ed integrazioni, Allegato II escluso il pto 4, attuata in Italia con Dlgs. 46 del 1997/02/24 e successive modifiche ed integrazioni per le seguenti tipologie di Dispositivi Medici/ *Is in compliance with the applicable requirements of 93/42/EEC Directive as amended, Annex II without point 4, transposed in Italy by Dlgs. 46 of 1997/02/24 as amended for the following Medical Devices:*

Accessori per l'infusione di cementi / *Accessories for infusion of cements*

Aghi e set per biopsia / *Biopsy needles and sets*

Cateteri da drenaggio, perfusione, dilatazione / *Drainage, perfusion, dilatation catheters*

Cateteri idrofilici / *Hydrophilic catheters*

Dispositivo per discectomia discale percutanea e accessori / *Percutaneous discectomy device*

Set da drenaggio / *Drainage set*

Set da infusione per radiologia / *Products for radiology*

Kiwa Cermet Italia S.p.A.  
Società con socio unico, soggetta  
all'attività di direzione e coordinamento  
di Kiwa Italia Holding S.r.l.  
Via Cadriano, 23  
40057 Granarolo dell'Emilia (BO)  
Tel +39.051.459.3.111  
Fax +39.051.763.382  
E-mail: info@kiwacermet.it  
www.kiwacermet.it

Rif. rapporto di audit/ *Ref. audit report:* del/dated 16-20/01/2018

**Chief Operating Officer**

*Giampiero Belcredi*



Organismo Notificato n. 0476  
*Notified Body nr. 0476*



Reg. Numero /  
Reg. Number MED 23002

Primo rilascio /  
First issue date 2003-09-17

Scadenza /  
Valid until 2023-09-16

Revisione /  
Revision 10

Valido da /  
Valid from 2018-09-14

Ultima modifica /  
Last change date 2019-03-04

# CERTIFICATE

## Allegato tecnico al Certificato/ Technical sheet enclosed to the Certificate

### Identificazione dei Dispositivi Medici/ Identification of Medical Devices:

#### Tipologia / Medical Devices:

Accessori per l'infusione di cementi / Accessories for infusion of cements

#### Classe di rischio / Risk class:

II a

#### Codice NANDO / NANDO codes:

MD 0102, MDS 7006 Ethylene oxide gas sterilization (EOG)

#### Marca / Brandname:

ADRIA

#### Modello / Model:

Aghi e set da infusione per trattamenti ossei / Needles for infusion and infusion set for osseous treatments

#### Codici / Codes:

49xxx; 490xxxxx

#### Marca / Brandname:

TEKNIMED

#### Modello / Model:

Aghi e set da infusione per trattamenti ossei / Needles for infusion and infusion set for osseous treatments

#### Codici / Codes:

T060xxx

#### Tipologia / Medical Devices:

Aghi e set per biopsia / Biopsy needles and sets

#### Classe di rischio / Risk class:

II a

#### Codice NANDO / NANDO codes:

MD 0106, MDS 7006 Ethylene oxide gas sterilization (EOG)

#### Modello / Model:

Aghi da biopsia ossea / Bone marrow biopsy needles

#### Codici / Codes:

aaa(a) xxx(x) ; xx(x)(x)yyy(y)

Kiwa Cermet Italia S.p.A.  
Società con socio unico, soggetta  
all'attività di direzione e coordinamento  
di Kiwa Italia Holding S.r.l.

Via Cadriano, 23  
40057 Granarolo dell'Emilia (BO)  
Tel +39.051.459.3.111  
Fax +39.051.763.382  
E-mail: info@kiwacermet.it  
www.kiwacermet.it

Chief Operating Officer

Giampiero Belcredi





Reg. Numero / <i>Reg. Number</i>	MED 23002	Revisione / <i>Revision</i>	10
Primo rilascio / <i>First issue date</i>	2003-09-17	Valido da / <i>Valid from</i>	2018-09-14
Scadenza / <i>Valid until</i>	2023-09-16	Ultima modifica / <i>Last change date</i>	2019-03-04

**Allegato tecnico al Certificato/  
*Technical sheet enclosed to the Certificate***

**Identificazione dei Dispositivi Medici/ *Identification of Medical Devices:***

**Tipologia / *Medical Devices:***

Aghi e set per biopsia / *Biopsy needles and sets*

**Modello / Model:**

Aghi per biopsia tessuti molli e accessori e introduttori / *Soft tissues needles and accessories and introductors*

**Codici / Codes:**

aaa(a) xxx(x) ; xx(x)(x)yyy(y)

**Modello / Model:**

Kit da cifoplastica e accessori / *Kiphoplastic kit and accessories*

**Codici / Codes:**

495xxyzz

**Modello / Model:**

Reperi mammari / *Breast localization*

**Codici / Codes:**

aaa(a) xxx(x) ; xx(x)(x)yyy(y)

**Tipologia / *Medical Devices:***

Cateteri da drenaggio, perfusione, dilatazione / *Drainage, perfusion, dilatation catheters*

**Classe di rischio / *Risk class:***

II a

**Codice NANDO / *NANDO codes:***

MD 0101, MDS 7006 Ethylene oxide gas sterilization (EOG)

**Modello / Model:**

Cateteri da dilatazione ed accessori per il gonfiaggio / *Dilatation catheters and high pressure inflation devices*

**Codici / Codes:**

xx(x)/yyy(y) ; xxxxyyyyyy(y)

**Classe di rischio / *Risk class:***

II b

**Codice NANDO / *NANDO codes:***

MD 0101, MDS 7006 Ethylene oxide gas sterilization (EOG)

**Chief Operating Officer**

*Giampiero Belcredi*





Reg. Numero / <i>Reg. Number</i>	MED 23002	Revisione / <i>Revision</i>	10
Primo rilascio / <i>First issue date</i>	2003-09-17	Valido da / <i>Valid from</i>	2018-09-14
Scadenza / <i>Valid until</i>	2023-09-16	Ultima modifica / <i>Last change date</i>	2019-03-04

Pagina / Page 4 di / of 5

## Allegato tecnico al Certificato/ *Technical sheet enclosed to the Certificate*

### Identificazione dei Dispositivi Medici/ *Identification of Medical Devices:*

#### Tipologia / *Medical Devices:*

Cateteri da drenaggio, perfusione, dilatazione / *Drainage, perfusion, dilatation catheters*

#### Modello / *Model:*

Cateteri da infusione e drenaggio e accessori / *Catheters for infusion and drainage and accessories*

#### Codici / *Codes:*

xx(x)/yyy(y) ; xxxxxxxxxx(y)

#### Tipologia / *Medical Devices:*

Cateteri idrofilici / *Hydrophilic catheters*

#### Classe di rischio / *Risk class:*

II a

#### Codice NANDO / *NANDO codes:*

MD 0101, MDS 7006 Ethylene oxide gas sterilization (EOG)

#### Modello / *Model:*

Cateteri idrofilici intermittenti / *Hydrophilic intermittent catheters*

#### Codici / *Codes:*

xxx(x) xxx(x)

#### Tipologia / *Medical Devices:*

Dispositivo per discectomia discale percutanea e accessori / *Percutaneous discectomy device*

#### Classe di rischio / *Risk class:*

II b

#### Codice NANDO / *NANDO codes:*

MD 1104, MDS 7006 Ethylene oxide gas sterilization (EOG)

#### Modello / *Model:*

Dispositivo per discectomia discale percutanea e accessori / *Percutaneous discectomy device*

#### Codici / *Codes:*

509xxyy

Kiwa Cermet Italia S.p.A.  
Società con socio unico, soggetta  
all'attività di direzione e coordinamento  
di Kiwa Italia Holding S.r.l.  
Via Cadriano, 23  
40057 Granarolo dell'Emilia (BO)  
Tel +39.051.459.3.111  
Fax +39.051.763.382  
E-mail: info@kiwacermet.it  
www.kiwacermet.it

Chief Operating Officer  
*Giampiero Belcredi*



Organismo Notificato n. 0476  
*Notified Body nr. 0476*



Reg. Numero /  
*Reg. Number* MED 23002

Primo rilascio /  
*First issue date* 2003-09-17

Scadenza /  
*Valid until* 2023-09-16

Revisione /  
*Revision* 10

Valido da /  
*Valid from* 2018-09-14

Ultima modifica /  
*Last change date* 2019-03-04

Pagina / Page 5 di / of 5

CERTIFICATE

**Allegato tecnico al Certificato/  
*Technical sheet enclosed to the Certificate***

**Identificazione dei Dispositivi Medici/ *Identification of Medical Devices:***

**Tipologia / *Medical Devices:***  
Set da drenaggio / *Drainage set*

**Classe di rischio / *Risk class:***  
II a

**Codice NANDO / *NANDO codes:***  
MD 0101, MDS 7006 Ethylene oxide gas sterilization (EOG)

**Modello / *Model:***  
Paracentesi / *paracensis*

**Codici / *Codes:***  
xx(x)yyy(y)

**Tipologia / *Medical Devices:***  
Set da infusione per radiologia / *Products for radiology*

**Classe di rischio / *Risk class:***  
II a

**Codice NANDO / *NANDO codes:***  
MD 0102, MDS 7005

**Modello / *Model:***  
Set da infusione per radiologia / *Products for radiology*

**Codici / *Codes:***  
xx(x)yyy(y)

La lista completa dei codici, relativi ai modelli certificati, è disponibile presso Kiwa Cermet Italia./ *The complete list of the codes related to the certificated models is available at Kiwa Cermet Italia.* Il presente Certificato è soggetto al rispetto dei requisiti contrattuali di Kiwa Cermet Italia ed è valido solo per le tipologie di dispositivi sopra identificate soggette a sorveglianza/ *This Certificate is subject to Kiwa Cermet Italia regulations and it is valid only for the above mentioned Medical Devices that are subject to survey.* L'allegato tecnico è parte integrante del presente Certificato./ *The technical sheet is an integrating part of this Certificate.*

Kiwa Cermet Italia S.p.A.  
Società con socio unico, soggetta  
all'attività di direzione e coordinamento  
di Kiwa Italia Holding S.r.l.  
Via Cadriano, 23  
40057 Granarolo dell'Emilia (BO)  
Tel +39.051.459.3.111  
Fax +39.051.763.382  
E-mail: info@kiwacermet.it  
www.kiwacermet.it

**Chief Operating Officer**  
*Giampiero Belcredi*



Organismo Notificato n. 0476  
*Notified Body nr. 0476*



To

**ADRIA S.r.l.**

**VIA MODENA, 46 40017 SAN GIOVANNI  
IN PERSICETO BO IT – Italia**

Sesto San Giovanni, 11.07.2023

**Notified Body Confirmation Letter**

**Reference: 000119756**

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, ICIM SPA, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 0425 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:

**ADRIA S.r.l.**

**VIA MODENA, 46 40017 SAN GIOVANNI IN PERSICETO BO IT – Italia**

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)

A On behalf of the Notified Body,  
 ICIM SPA  
 Piazza don Enrico Mapelli, 75  
 20099 Sesto San Giovanni Milano  
 Nando Code: 0425

**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
Biopsy needles and sets	Class IIa	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Accessories for infusion of cements	Class IIa	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Drainage, perfusion, dilatation, catheters	Class IIa	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Hydrophilic catheters	Class IIa	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Percutaneous discectomy device	Class IIb	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Drainage set	Class Is	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476
Product for radiology	Class IIa	N.A.	Certificate n. MED 23002 Valid until 16.09.2023 Notified Body n. 0476

**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
N.A.	N.A.	N.A.	N.A.
N.A.	N.A.	N.A.	N.A.
N.A.	N.A.	N.A.	N.A.

### Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
11.07.2023	000119756	Initial issue

While remaining at your disposal for any clarification on the content of this offer, we take this opportunity to extend our best regards.

Dott. Edoardo Dossena  
 Product Sales Manager Certificazione  
 Prodotto, Ispezioni e Direttive  
  
 ICIM S.p.A.

Dott. Dario Bruno  
 Direttore Commerciale  
  
 ICIM S.p.A.



## Manufacturer's Declaration

in relation to 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, in particular with respect to

- the validity of certificates issued under Council Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD) or Council Directive 93/42/EEC on Medical Devices (MDD) (Directive Certificates) *and/or*<sup>1</sup>
- the compliance of the devices and us as their manufacturer with the conditions for the continued placing on the market and putting into service

Manufacturer name	ADRIA SRL
Manufacturer address and contact details	Registered Headquarter: via Modena 46, 40017 San Giovanni in Persiceto (BO) - Italy Operational Headquarter: via San Francesco d' Assisi 15, 46020 Pegognaga (MN) – Italy Ph. +39 0376 559772 Email:customer-service@adriamedical.com
Single Registration Number (SRN) (if available)	IT-MF-000027852

Authorised Representative name (if applicable)	N.A.
Authorised Representative address and contact details	N.A.
Single Registration Number (SRN) (if available)	N.A.

Notified body name (if applicable)	KIWA CERMET ITALIA
------------------------------------	--------------------

<sup>1</sup> The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body.



		X See attached schedule
Notified body number (if applicable)	0476	X See attached schedule
Directive Certificate number(s) to which this confirmation is made (if applicable)	MED 23002	X See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	2023-09-16	X See attached schedule
End date of extended validity/transition period	2028-12-31	X See attached schedule

We, as the manufacturer declare under our sole responsibility:

- for the above listed **Directive Certificate** (or see attached schedule, if multiple certificates) the conditions for the legal extension of validity as required in Article 120.2 of the MDR are met *and/or*<sup>2</sup>
- the listed **device(s)** in the attached schedule and we as their manufacturer are in compliance with the conditions listed in Article 120.3c of the MDR for continued placing on the market and putting into service,

namely by fulfilling the following conditions:

➤ **Directive Certificate(s)** as listed above or in the attached schedule

- Directive Certificate(s) covering the listed device(s) was/were issued after 25 May 2017, was/were valid on 26 May 2021 and have not been withdrawn afterwards.

*Choose applicable statements:*

- Expired *before* 20 March 2023:
  - Before the original date of expiry as indicated on the Directive Certificate(s), we and the notified body have signed written agreement(s) in accordance with Section 4.3, second subparagraph of Annex VII to this Regulation for the conformity assessment(s) in respect of the device(s) covered by the expired certificate(s) or in respect of a device(s) intended to substitute that/those device(s), or

---

<sup>2</sup> The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body



- A Competent Authority has granted a derogation from the applicable conformity assessment procedure in accordance with Article 59(1) MDR (may be provided upon request), or
- A Competent Authority has required the manufacturer, in accordance with Article 97(1) MDR, to carry out the applicable conformity assessment procedure (may be provided upon request)

*Choose one of the following statements only if a derogation per Article 59(1) or a requirement per Article 97(1) has been granted by a Competent Authority:*

- Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.
- Expired/expires *after* 20 March 2023:

*Choose one applicable statement:*

- Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

#### ➤ **Upclassified devices**

In case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body:

*Choose one applicable statement:*

- Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitutes and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

ADRIA S.r.l.  
Headquarter: via Modena 46, 40017 San Giovanni in Persiceto (BO) - Italy  
Warehouse and logistics: via San Francesco d'Assisi 15, 46020 Pegognaga (MN) - Italy  
VAT Reg. id. 02042571204 - Tel. +39 0376 559772  
www.adriamedical.com - E-mail: customer-service@adriamedical.com



➤ **Quality Management System (QMS)**

*Choose one applicable statement:*

- A QMS in accordance with Article 10(9) MDR will be put in place by no later than 26 May 2024.
- A QMS in accordance with Article 10(9) MDR is in place.
- A notified body has issued the attached certificate for the MDR-compliant QMS.

➤ **Device(s) as listed in the attached schedule**

- The device(s) continue to comply with the AIMDD or MDD.
- There are no significant changes in the design and intended purpose.
- The device(s) do not present an unacceptable risk to health or safety of patients, users or other persons, or to other aspects of the protection of public health.

**Signed for and on behalf of the manufacturer:**

Full Company Name: ADRIA SRL

Location & Date: San Giovanni in Persiceto, 04/08/2023

Signature, Print Name, Title: Maria Vittoria Avaltroni, QA Responsible

Contact Details: [qa@adriamedical.com](mailto:qa@adriamedical.com)

**ADRIA srl**  
Via Modena, 46  
40017 S. GIOVANNI IN PERSICETO (BO) Italy  
Tel. +39 0376559772  
PEC: [adriamedical.pec@legalmail.it](mailto:adriamedical.pec@legalmail.it)  
E-mail: [customer-service@adriamedical.com](mailto:customer-service@adriamedical.com)  
Partita IVA 02042571204 - Codice SDI M5UXCR1

ADRIA S.r.l.  
Headquarter: via Modena 46, 40017 San Giovanni in Persiceto (BO) - Italy  
Warehouse and logistics: via San Francesco d'Assisi 15, 46020 Pegognaga (MN) - Italy  
VAT Reg. id. 02042571204 - Tel. +39 0376 559772  
[www.adriamedical.com](http://www.adriamedical.com) - E-mail: [customer-service@adriamedical.com](mailto:customer-service@adriamedical.com)



## Schedule of Devices

The above Manufacturer's Declaration is valid for the following devices:

Identification of the device(s) <sup>3</sup> (e.g., device name, family/group name, device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made (if applicable)	Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity (if applicable)	Notified Body name and number that issued the Directive Certificate (if applicable)	Notified Body name and number where the MDR application was lodged/contract signed (if applicable)	End date of extended validity / transition period	Substitute Device(s) (if applicable)
<b>Biopsy needles and sets</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Accessories for infusion of cements</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Drainage, perfusion, dilatation, catheters</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Hydrophilic catheters</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Percutaneous discectomy device</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Drainage set</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.
<b>Products for radiology</b>	MED 23002	2023-09-16	KIWA CERMET ITALIA n. 0476	ICIM SPA n. 0425	2028-12-31	N.A.

<sup>3</sup> for devices with AIMDD/MDD certificate(s) the identification should be as in the certificate, and only if the certificate has a generic scope it should be as defined above)