



CERTIFICADO CE
de acuerdo con los Anexos VII punto 5 y V punto 3 de la Directiva 93/42/CEE
EC CERTIFICATE
in accordance with Annexes VII section 5 and V section 3 of Directive 93/42/EEC

Certificado nº/Certificate no	Fecha de validez/Date of validity	ON nº/NB no
2000 06 0272 CP	Desde/From 05-03-2020 Hasta/To 26-05-2024	0318

A favor de/In favour of:

Fabricante/Manufacturer:
Nombre/Name: BECTON DICKINSON S.A.
Dirección/Address: C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España
Representante autorizado ante la UE/ Authorized EU representative: Idem

Para el producto estéril con función de medición / For the sterile product with a measuring function

Categoría/Category: Productos de un solo uso/ Single use products
Grupo genérico/ Generic group: Productos para inyección y/o aspiración de fluidos/ Instruments for injection and/or aspiration of fluids
Tipo/Type: Especificado en Anexos de este Certificado/ Specified in Annexes to this Certificate

Elaborado en/In the facilities:

C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España
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Fecha inicial/ Initial date: 19-06-1995

Fecha de prórroga anterior/ Previous extension date: 05-05-2015

Este certificado se emite en base a la auditoría contenida en el expediente nº 95 05 0005, y garantiza que los aspectos de fabricación relacionados con la obtención y el mantenimiento de las condiciones de esterilidad y con la conformidad de los requisitos metroológicos han sido evaluados de acuerdo con el punto 3 del Anexo V de la Directiva.

This certificate is issued on the basis of the audit contained in dossier no 95 05 0005, and guarantees that the aspects of manufacture concerned with securing and maintaining sterile conditions and with the conformity with the metrological requirements have been assessed in accordance with Annex V section 3 of Directive.

Madrid, 05 de marzo de 2020

DIRECTORA DE LA AGENCIA ESPAÑOLA DE MEDICAMENTOS Y PRODUCTOS SANITARIOS

 **agencia española de medicamentos y productos sanitarios**

Fdo. Mª Jesús Lamas Díaz

Agencia Española de Medicamentos y Productos Sanitarios	Localizador: B 4 G Z K A J 6 B A
Fecha de la firma: 05/03/2020	
<i>Puede comprobar la autenticidad del documento en la sede de la AEMPS: https://sede.aemps.gob.es</i>	

CORREO ELECTRÓNICO
on0318@aemps.es

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C/ CAMPEZO, 1 - EDIFICIO 8
28022 MADRID
Tel.: (+34) 902.101.322 / (+34) 91.822.59.97
Fax: (+34) 91.822.52.89

ORGANISMO NOTIFICADO 0318



ANEXO N°/ANNEX NO: I
CERTIFICADO CE
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Certificado n°/Certificate no 2000 06 0272 CP	Fecha de validez/Date of validity Desde/From 05-03-2020 Hasta/To 26-05-2024	ON n°/NB no 0318
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A favor de/In favor of:

Fabricante/Manufacturer: Nombre/Name: BECTON DICKINSON S.A. Dirección/Address: C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España Representante autorizado ante la UE/Authorized EU representative: Idem
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Tipo de producto / Devices type: Jeringas sin aguja y jeringa con aguja de punta roma/ Syringes without needle and syringes with blunt fill needle.

Clasificación/Classification: I estéril con función de medición/ I sterile with a measuring function.

1. Jeringas estériles sin aguja / Sterile syringes without needle [MD 0104] [MDS 7006]

1.1. Jeringas de dos piezas BD Discardit™ II / Two-piece syringes BD Discardit™ II

- 1.1.a).** Jeringas de:/ Syringes of: 2 ml
- 1.1.b).** Jeringas de:/ Syringes of: 5 ml
- 1.1.c).** Jeringas de:/ Syringes of: 10 ml
- 1.1.d).** Jeringas de:/ Syringes of: 20 ml

1.2. Jeringas de dos piezas con pistón rompible BD SoloMed™/ Two-piece syringes with breakable plunger BD SoloMed™

- 1.2.a).** Jeringas de:/ Syringes of: 2 ml
- 1.2.b).** Jeringas de:/ Syringes of: 5 ml
- 1.2.c).** Jeringas de:/ Syringes of: 10 ml

1.3. Jeringas de tres piezas BD Emerald™ / Sterile three-piece syringes BD Emerald™

- 1.3.a).** Jeringas de:/ Syringes of: 2 ml
- 1.3.b).** Jeringas de:/ Syringes of: 5 ml
- 1.3.c).** Jeringas de:/ Syringes of: 10 ml

Agencia Española de Medicamentos y Productos Sanitarios

Fecha de la firma: 05/03/2020

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1.4. Jeringas de tres piezas BD Emerald™ Pro/ Three-piece syringes BD Emerald™ Pro

1.4.a). Jeringas de:/ Syringes of: 2 ml

1.4.b). Jeringas de:/ Syringes of: 5 ml

1.4.c). Jeringas de:/ Syringes of: 10 ml

1.5. Jeringas de tres piezas estériles BD Emerald™ con cono excéntrico/ Eccentric tip sterile three-piece syringe BD Emerald™

1.5.a). Jeringas de:/ Syringe of: 10ml

2. Jeringas estériles de tres piezas con aguja de punta roma para aspiración de medicamentos desde viales o ampollas/ Sterile three-piece syringes with blunt fill needle to aspirate pharmaceutical fluids from vials or ampoules

2.1. Jeringas de tres piezas estériles con aguja punta roma/ Syringe with blunt fill needle

2.1.a). Jeringa de 2 ml con aguja de punta roma 18G × 1 ½" (1.2 × 40 mm)/ Syringe of 2 ml with 18G × 1 ½" (1.2 × 40 mm) blunt fill needle

2.1.b). Jeringa de 5 ml con aguja de punta roma 18G × 1 ½" (1.2 × 40 mm)/ Syringe of 5 ml with 18G × 1 ½" (1.2 × 40 mm) blunt fill needle

2.1.c). Jeringa de 10 ml con aguja de punta roma 18G × 1 ½" (1.2 × 40 mm)/ Syringe of 10 ml with 18G × 1 ½" (1.2 × 40 mm) blunt fill needle

Este certificado ampara todas las marcas de estos productos incluidas por el fabricante en su declaración de conformidad. This certificate covers all trademarks of these products included by the manufacturer in his declaration of conformity.

Madrid, 05 de marzo de 2020

DIRECTORA DE LA AGENCIA ESPAÑOLA DE MEDICAMENTOS Y PRODUCTOS SANITARIOS

 **agencia española de
medicamentos y
productos sanitarios**

Fdo. M^a Jesús Lamas Díaz

Agencia Española de Medicamentos y Productos Sanitarios

Fecha de la firma: 05/03/2020

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ORGANISMO NOTIFICADO 0318

Becton Dickinson S.A.
Camino de Valdeoliva, s/n
San Agustin del Guadalix
Madrid
28750
Spain

12 March 2024

Notified Body Confirmation Letter
Reference: EU2023-607/ 764888

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, **BSI Group The Netherlands B.V.**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **2797** 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:

Becton Dickinson S.A.
Camino de Valdeoliva, s/n
San Agustin del Guadalix
Madrid
28750
Spain
SRN Number: ES-MF-000016479

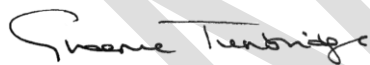
The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below. Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which the NB is also responsible for appropriate surveillance 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 BSI Group The Netherlands B.V.,



Graeme Tunbridge
Senior Vice President, Medical Devices

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
N/A	N/A	N/A	N/A

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
BD Plastipak™ Luer-Lok™ Syringe (without needle) Basic UDI-DI: 038290SYLPEAGDM5 Listed on MDD certificate as: <i>Three-piece syringes without needle;</i> <i>Sterile BD Plastipak™ Luer-Lok™ syringes;</i> <i>Amber color syringes of: 50 ml</i>	Class IIa	N/A	Certificate reference: 95 06 0005 CP; expiry date 2024-05-24, Number: 0318
BD Plastipak™ Luer-Lok™ Convenience Pack (without needle) Basic UDI-DI: 038290UOKRIXNENG Listed on MDD certificate as: <i>Three-piece syringes without needle;</i> <i>Sterile BD Plastipak™ Luer-Lok™ syringes;</i>	Class IIa	'N/A'	Certificate reference: 95 06 0005 CP; expiry date 2024-05-24, NB number 0318

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
<p><i>Sizes:</i></p> <p><i>Convenience Pack of syringes of: 20 ml,</i></p> <p><i>Convenience Pack of syringes of: 50 ml</i></p>			
<p>BD Plastipak™ Catheter Tip Syringe</p> <p>Basic UDI-DI: 038290TWOWNMYHW3</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle, Insulin syringes; Three-piece sterile BD Plastipak™ syringes without needle; Syringes with Catheter Tip connection</i></p> <p><i>Sizes:</i></p> <p><i>Syringes of: 50 ml</i></p> <p><i>Syringes of: 100 ml</i></p>	<p>Class I device placed on the market in sterile condition</p> <p>Class I device with a measuring function</p>	<p>'N/A'</p>	<p>Certificate reference: 2000 06 0273 CP; expiry date 2024-05-24, NB number 0318</p>
<p>BD Plastipak™ Catheter Tip Syringe (Bulk Non Sterile)</p> <p>Basic UDI-DI: 038290IHOCSHLU8L</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle; Non-sterile three-piece syringes without needle BD Plastipak™; Syringes with Catheter Tip connection</i></p> <p><i>Size:</i></p> <p><i>Syringes of: 50 ml</i></p>	<p>Class I device with a measuring function</p>	<p>'N/A'</p>	<p>Certificate reference: MDD/AIMDD Certificate #2019 09 0898 CP; expiry date 2024-05-24, NB# 0318</p>
<p>BD Plastipak™ Luer-Slip Syringe (without needle)</p>	<p>Class I device placed on the</p>	<p>'N/A'</p>	<p>Certificate reference:</p>

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
<p>Basic UDI-DI: 038290SRGVPAAEKC</p> <p>Listed on the MDD certificate as: <i>Syringes without needle, Insulin syringes; Three-piece sterile BD Plastipak™ syringes without needle; Luer-Slip syringes</i></p> <p><i>Sizes:</i> <i>Syringes of: 1 ml</i> <i>Syringes of: 2 ml</i> <i>Syringes of: 5 ml</i> <i>Syringes of: 10 ml</i> <i>Syringes of: 20 ml</i> <i>Syringes of: 30 ml</i> <i>Syringes of: 50 ml</i></p>	<p>market in sterile condition</p> <p>Class I device with a measuring function</p>		<p>#2000 06 0273 CP; expiry date 2024-05-24, NB number 0318</p>
<p>BD Plastipak™ Luer-Slip Syringe (without needle Bulk Non Sterile)</p> <p>Basic UDI-DI: 038290XOKHQIDL</p> <p>Listed on the MDD certificate as: <i>Syringes without needle; Non-sterile three-piece syringes without needle BD Plastipak™; Luer-Slip syringes</i></p> <p><i>Sizes:</i> <i>Syringes of: 1 ml</i> <i>Syringes of: 20 ml</i> <i>Syringes of: 50 ml</i></p>	<p>Class I device with a measuring function</p>	<p>'N/A'</p>	<p>Certificate reference: #2019 09 0898 CP; expiry date 2024-05-24, NB# 0318</p>
<p>BD Whitacre Spinal Needle</p>	<p>Class III</p>	<p>'N/A'</p>	<p>Certificate reference:</p>

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
<p>Basic UDI-DI: 038290JPCVIFOCBR</p> <p>Listed on the MDD certificate as:</p> <p><i>Spinal needles; Whitacre Pencil Point Spinal Needles; Sterile Whitacre Pencil Point Spinal Needle: BD Whitacre Needle</i></p> <p><i>Sizes:</i></p> <p><i>22G × 3.50" (0.7 × 90 mm)</i></p> <p><i>24G × 3.50" (0.55 × 90 mm)</i></p> <p><i>25G × 3.50" (0.50 × 90 mm)</i></p> <p><i>27G × 3.50" (0.40 × 90 mm)</i></p>			<p>#95 06 0005 CP and 2010 02 0700 ET; expiry date 2024-05-24, NB# 0318</p>
<p>BD Whitacre Spinal Needle Set (Spinal needle + introducer)</p> <p>Basic UDI-DI: 038290KITVJVYYLK</p> <p>Listed on the MDD certificates as:</p> <p><i>Sterile Whitacre Pencil Point Spinal Needle with introducer set, BD Whitacre Needle</i></p> <p><i>Sizes:</i></p> <p><i>25G × 3.50" (0.50 × 90 mm) with introducer 20G × 1.25" (0.9 × 32 mm)</i></p> <p><i>25G × 4.06" (0.50 × 103 mm) with introducer 20G × 1.25" (0.9 × 32 mm)</i></p> <p><i>27G × 3.50" (0.40 × 90 mm) with introducer 22G × 1.25" (0.7 × 32 mm)</i></p> <p><i>27G × 4.06"(0.40 × 103 mm) with introducer 22G × 1.25" (0.7 × 32 mm)</i></p>	Class III	'N/A'	<p>Certificate reference:</p> <p>#95 06 0005 CP and 2010 02 0700 ET; expiry date 2024-05-24, NB# 0318, to be read in conjunction with amending AEMPS letter dated 16 November 2021, ON0318/GH/JC/PS/95 04 0005</p>
<p>BD Quincke Spinal Needle</p>	Class III	'N/A'	<p>Certificate reference:</p>

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
<p>Basic UDI-DI: 038290HFPYXWDAGC</p> <p>Listed on the MDD certificate as:</p> <p><i>Spinal needles; Spinal needle Quincke type point; Sterile spinal needle Quincke type point: BD Spinal Needle</i></p> <p><i>Sizes:</i></p> <p><i>18G × 3.00" (1.2 × 75 mm)</i> <i>18G × 3.50" (1.2 × 90 mm)</i> <i>19G × 3.00" (1.1 × 75 mm)</i> <i>19G × 3.50" (1.1 × 90 mm)</i> <i>20G × 1.50" (0.90 × 38 mm)</i> <i>20G × 3.00" (0.90 × 75 mm)</i> <i>20G × 3.50" (0.90 × 90 mm)</i> <i>22G × 1.50" (0.7 × 38 mm)</i> <i>22G × 2.50" (0.7 × 63 mm)</i> <i>22G × 3.00" (0.7 × 75 mm)</i> <i>22G × 3.50" (0.7 × 90 mm)</i> <i>23G × 3.50" (0.64 × 90 mm)</i> <i>25G × 3.00" (0.50 × 75 mm)</i> <i>25G × 3.50" (0.50 × 90 mm)</i> <i>26G × 3.50" (0.45 × 90 mm)</i> <i>27G × 3.50" (0.40 × 90 mm)</i></p>			<p>#95 06 0005 CP and 2010 02 0701 ET; expiry date 2024-05-24, NB# 0318</p>
<p>BD Quincke Spinal Needle Set (Spinal needle + introducer)</p> <p>Basic UDI-DI: 038290JBDWOAQF5Y</p> <p>Listed on the MDD certificate as:</p>	Class III	'N/A'	<p>Certificate reference:</p> <p>MDD/AIMDD Certificate #95 06 0005 CP and 2010 02 0701 ET; expiry date 2024-05-24, NB# 0318, to be read in conjunction with amending AEMPS letter dated 15</p>

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
<p><i>Spinal needles; Spinal needle Quincke type point; Sterile spinal needle Quincke type point with introducer set; BD Spinal Needle</i></p> <p><i>Sizes:</i></p> <p><i>25G × 3.50" (0.50 × 90 mm) con introducer / with introducer 20G × 1.25" (0.9 × 32 mm)</i></p> <p><i>26G × 3.50" (0.45 × 90 mm) con introducer / with introducer 20G × 1.25" (0.9 × 32 mm)</i></p> <p><i>27G × 3.50" (0.40 × 90 mm) con introducer / with introducer 22G × 1.25" (0.7 × 32 mm)</i></p>			November 2021 S/REF CNCps ID: 90811
<p>BD Quincke Spinal NRFit™ Needle</p> <p>Basic UDI-DI: 038290WAHENDUTA4</p> <p>Listed on the MDD certificate as:</p> <p><i>Spinal needles; BD Quincke Spinal NRFit™ Needles; Sterile BD Quincke Spinal NRFit™ Needles BD Quincke Spinal NRFit™ Needle</i></p> <p><i>Sizes:</i></p> <p><i>25G × 3.00" (0.5 × 76.2 mm)</i></p> <p><i>18G × 3.00" (1.2 × 76.2 mm)</i></p> <p><i>18G × 3.50" (1.2 × 88.9 mm)</i></p> <p><i>22G × 1.50" (0.7 × 38.1 mm)</i></p> <p><i>22G × 2.50" (0.7 × 63.5 mm)</i></p> <p><i>22G × 3.00" (0.7 × 76.2 mm)</i></p> <p><i>22G × 3.50" (0.7 × 88.9 mm)</i></p> <p><i>25G × 3.50" (0.5 × 88.9 mm)</i></p> <p><i>25G × 2.00" (0.5 × 50.8 mm)</i></p> <p><i>19G × 3.00" (1.1 × 76.2 mm)</i></p>	Class III	N/A	Certificate reference: #95 06 0005 CP and 2020 04 0913 ET; expiry date 2024-05-24, NB# 0318

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
<p>19G × 3.50" (1.1 × 88.9 mm) 20G × 1.50" (0.9 × 38.1 mm) 20G × 3.00" (0.9 × 76.2 mm) 20G × 3.50" (0.9 × 88.9 mm) 23G × 3.50" (0.6 × 88.9 mm) 26G × 3.50" (0.45 × 88.9 mm) 27G × 3.50" (0.4 × 88.9 mm)</p>			
<p>BD Quincke Spinal NRFit™ Needle Set (NRFit Spinal Needle + introducer) Basic UDI-DI: 038290MIVCNRJEZ</p> <p>Listed on the MDD certificate as: <i>Spinal needles; BD Quincke Spinal NRFit™ Needles; Sterile BD Quincke spinal NRFit™ Needle with introducer set. BD Quincke Spinal NRFit™ Needle Set</i></p> <p>Sizes: <i>25G × 3.50" (0.50 × 88.9 mm) con introducer / with introducer 20 G × 1.25" (0.90 × 31.8 mm)</i> <i>27G × 3.50" (0.40 × 88.9 mm) con introducer / with introducer 22 G × 1.25" (0.70 × 31.8 mm)</i> <i>26G × 3.50" (0.45 × 88.9 mm) with introducer / with introducer 20G x 1.25" (0.90 x 31.8 mm)</i></p>	Class III	N/A	<p>Certificate reference: #95 06 0005 CP and 2020 04 0913 ET; expiry date 2024-05-24, NB# 0318, to be read in conjunction with amending AEMPS letter dated 15 november 2021, S/REF CNCps ID: 92262</p>
<p>BD Whitacre Spinal NRFit™ Needle Basic UDI-DI: 038290RFMMCXCCX</p>	Class III	N/A	<p>Certificate reference: #95 06 005 CP and 2020 04 0912 ET; expiry date 2024-05-24, NB# 0318</p>

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
<p>Listed on the MDD certificate as:</p> <p><i>Spinal needles; BD Whitacre Spinal NRFit™ Needles; Sterile BD Whitacre Spinal NRFit™ Needle BD Whitacre Spinal NRFit™ Needle</i></p> <p><i>Sizes:</i></p> <p><i>25G × 4.06" (0.50 × 103.2 mm)</i></p> <p><i>25G × 4.70" (0.50 × 119.1 mm)</i></p> <p><i>27G × 4.06" (0.40 × 103.2 mm)</i></p> <p><i>27G × 4.70" (0.40 × 119.1 mm)</i></p> <p><i>22G × 3.50" (0.70 × 88.9 mm)</i></p> <p><i>24G × 3.50" (0.55 × 88.9 mm)</i></p> <p><i>25G × 3.50" (0.50 × 88.9 mm)</i></p> <p><i>27G × 3.50" (0.40 × 88.9 mm)</i></p>			
<p>BD Whitacre Spinal NRFit™ Needle Set (NRFit Spinal Needle + introducer)</p> <p>Basic UDI-DI: 038290EDURMMPSBE</p> <p>Listed on the MDD certificate as:</p> <p><i>Spinal needles; BD Whitacre Spinal NRFit™ Needles; Sterile BD Whitacre Spinal NRFit™ Needle with introducer set. BD Whitacre Spinal NRFit™ Needle Set.</i></p> <p><i>Sizes:</i></p> <p><i>27G × 3.50" (0.40 × 88.9 mm) with introducer 22G × 1.25" (0.70 × 31.8 mm)</i></p> <p><i>25G × 3.50" (0.50 × 88.9 mm) with introducer 20G × 1.25" (0.90 × 31.8 mm)</i></p>	Class III	N/A	<p>Certificate reference:</p> <p>#95 06 005 CP and 2020 04 0912 ET; expiry date 2024-05-24, NB# 0318</p>

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
<p><i>25G × 4.06" (0.50 × 103.2 mm) with introducer 20G × 1.25" (0.90 × 31.8 mm)</i></p> <p><i>27G × 4.06" (0.40 × 103.2 mm) with introducer 22G × 1.25" (0.70 × 31.8 mm)</i></p>			
<p>BD Spinal Introducer NRFit Needle</p> <p>Basic UDI-DI (Sterile): 038290XGKVVVIXRA</p> <p>Listed on the MDD certificate as:</p> <p><i>Needle introducer; Sterile BD Spinal needle introducer NRFit™ BD Spinal needle introducer NRFit™</i></p> <p>Sizes:</p> <p><i>20G × 1.25" (0.90 × 31.8 mm)</i></p> <p><i>22G × 1.25" (0.70 × 31.8 mm)</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>#95 06 0005 CP; expiry date 2024-05-24, NB# 0318</p>
<p>BD Discardit II Syringe (with Needle)</p> <p>Basic UDI-DI (Microlance needle): 038290HLQCKJFL8D</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes with needle; Sterile two-piece syringes with needle BD Discardit™ II</i></p> <p>Sizes:</p> <p>Syringes of: 2 ml:</p> <p><i>With needle: 21G × 5/8" (0.8 × 16 mm)</i></p> <p><i>With needle: 22G × 1 1/2" (0.7 × 40 mm)</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p><i>With needle: 24G × 1" (0.55 × 25 mm)</i></p> <p><i>Syringes of: 5 ml</i></p> <p><i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>Syringes of: 10 ml</i></p> <p><i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>With needle: 21G × 1" (0.8 × 25 mm)</i></p> <p><i>Syringes of: 20 ml</i></p> <p><i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i></p>			
<p>BD Discardit II Non Sterile Bulk Syringe (without needle)</p> <p>Basic UDI-DI: 038290IWCCNEZKAT</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle; Non-sterile syringes without needle; Two-piece syringes BD Discardit™ II</i></p> <p><i>sizes:</i></p> <p><i>Syringes of: 2 ml</i></p> <p><i>Syringes of: 5 ml</i></p> <p><i>Syringes of: 10 ml</i></p> <p><i>Syringes of: 20 ml</i></p>	Class I device with a measuring function	N/A	<p>Certificate reference:</p> <p>MDD/AIMDD Certificate #99 03 0213 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>
<p>BD Discardit II Syringe (without needle)</p> <p>Basic UDI-DI: 038290DDEODFOLXL</p>	Class I device placed on the market in sterile condition	N/A	<p>Certificate reference:</p> <p>MDD/AIMDD Certificate #2000 06 0272 CP; expiry date 2024-05-26, NB# 0318</p>

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
<p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle and syringes with blunt fill needle; Sterile syringes without needle; Two-piece syringes BD Discardit™ II</i></p> <p><i>Syringes of: 2 ml</i></p> <p><i>Syringes of: 5 ml</i></p> <p><i>Syringes of: 10 ml</i></p> <p><i>Syringes of: 20 ml</i></p>	Class I device with a measuring function		<p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>
<p>BD Microlance™ 3 Needle</p> <p>Basic UDI-DI: 038290MLFUVFGJET</p> <p><i>Listed on the MDD certificate as:</i></p> <p><i>Needles; Sterile hypodermic needles BD Microlance™ 3</i></p> <p><i>Sizes:</i></p> <p><i>Needle 20G × 1 ½" (0.9 × 40 mm)</i></p> <p><i>Needle 20G × 1" (0.9 × 25 mm)</i></p> <p><i>Needle 21G × 1" (0.8 × 25 mm)</i></p> <p><i>Needle 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>Needle 21G × ⅝" (0.8 × 16 mm)</i></p> <p><i>Needle 22G × 1 ¼" (0.7 × 30 mm)</i></p> <p><i>Needle 22G × 1 ½" (0.7 × 40 mm)</i></p> <p><i>Needle 22G × 1" (0.7 × 25 mm)</i></p> <p><i>Needle 23G × 1 ¼" (0.6 × 30 mm)</i></p> <p><i>Needle 23G × 1" (0.6 × 25 mm)</i></p> <p><i>Needle 22G × 2" (0.7 × 50 mm)</i></p> <p><i>Needle 26G × ⅜" (0.45 × 10 mm)</i></p> <p><i>Needle 25G × 1" (0.5 × 25 mm)</i></p> <p><i>Needle 25G × ⅝" (0.5 × 16 mm)</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>MDD/AIMDD Certificate #95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p><i>Needle 27G × ½" (0.4 × 13 mm)</i></p> <p><i>Needle 16G × 1 ½" (1.6 × 40 mm)</i></p> <p><i>Needle 21G × 2" (0.8 × 50 mm)</i></p> <p><i>Needle 19G × 1 ½" (1.1 × 40 mm)</i></p> <p><i>Needle 19G × 1" (1.1 × 25 mm)</i></p> <p><i>Needle 19G × 2" (1.1 × 50 mm)</i></p> <p><i>Needle 18G × 2" (1.2 × 50 mm)</i></p> <p><i>Needle 27G × ¾" (0.4 × 19 mm)</i></p> <p><i>Needle 26G × ½" (0.45 × 13 mm)</i></p> <p><i>Needle 30G × ½" (0.3 × 13 mm)</i></p> <p><i>Needle 24G × 1" (0.55 × 25 mm)</i></p> <p><i>Needle 26G × ⅝" (0.45 × 16 mm)</i></p> <p><i>Needle 18G × 1 ½" (1.2 × 40 mm)</i></p> <p><i>Needle 27G × ⅜" (0.4 × 10 mm)</i></p> <p><i>Needle 23G × 1 ½" (0.6 × 40 mm)</i></p>			
<p>BD Microlance TM 3 Non Sterile Bulk Needle</p> <p>Basic UDI-DI: 038290MVVIHDPPLC</p> <p>Listed on the MDD certificate as:</p> <p><i>Needles; Non-sterile hypodermic needles BD Microlance™ 3</i></p> <p><i>Needle 25G × ⅝" (0.5 × 16 mm)</i></p> <p><i>Needle 21G × ⅝" (0.8 × 16 mm)</i></p> <p><i>Needle 25G × 1" (0.5 × 25 mm)</i></p> <p><i>Needle 21G × 1" (0.8 × 25 mm)</i></p> <p><i>Needle 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>Needle 20G × 1 ½" (0.9 × 40 mm)</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318, to be read in conjunction with amending AEMPS letter dated 10 March 2022 regarding and change submitted 22 December 2021, S/REF 94331</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p><i>Needle 22G x 1 ¼" (0.7 x 30 mm)</i></p> <p><i>Needle 30G x ½" (0.3 x 13 mm)</i></p> <p><i>Needle 27G x ½" (0.4 x 13 mm)</i></p> <p><i>Needle 19G x 1 ½" (1.1 x 40 mm)</i></p> <p><i>Needle 14G x 1 ¼" (2.1 x 30 mm)</i></p> <p><i>Needle 26G x ½" (0.45 x 13 mm)</i></p> <p><i>Needle 26G x ⅜" (0.45 x 10 mm)</i></p> <p><i>Needle 23G x 1" (0.6 x 25 mm)</i></p> <p><i>Needle 21G x 2" (0.8 x 50 mm)</i></p> <p><i>Needle 18G x 2" (1.2 x 50 mm)</i></p> <p><i>Needle 27G x ¾" (0.4 x 19 mm)</i></p> <p><i>Needle 24G x 1" (0.55 x 25 mm)</i></p> <p><i>Needle 18G x 1 ½" (1.2 x 40 mm)</i></p> <p><i>Needle 22G x 1" (0.7 x 25 mm)</i></p> <p><i>Needle 22G x 1 ½" (0.7 x 40 mm)</i></p> <p><i>Needle 23G x 1 ¼" (0.6 x 30 mm)</i></p> <p><i>New non-sterile variant of the Microlance™ 3 27G x ⅜" (0.4 x 10 mm) needle</i></p>			
<p>BD SoloShot™ Mini Syringe</p> <p>Basic UDI-DI: 038290CDXRROBXD6</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes with needle; Sterile auto-disable two-piece syringes with integrated cannula BD SoloShot™ Mini</i></p> <p><i>Sizes:</i></p> <p><i>2.6.a Syringes of: 0.5 ml</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p> <p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p>2.6.a.1 With needle: 23G × 1" (0.6 × 25 mm)</p> <p>2.6.a.2 With needle: 24G × ¾" (0.55 × 19 mm)</p> <p>2.6.a.3 With needle: 25G × ½" (0.5 × 16 mm)</p> <p>2.6.a.4 With needle: 25G × 1" (0.5 × 25 mm)</p> <p>2.6.b Syringes of: 0.1 ml</p> <p>2.6.b.1 With needle: 27G × ¾" (0.4 × 10 mm)</p> <p>2.6.c Syringes of: 0.05 ml</p> <p>2.6.c.1 With needle: 27G × ¾" (0.4 × 10 mm)</p> <p>2.5.d. Syringes of 0.2 ml</p> <p>2.5.d.1. With needle: 23G × 1" (0.6 × 25 mm)</p> <p>2.5.e. Syringes of 0.3 ml</p> <p>2.5.e.1. With needle: 23G × 1" (0.6 × 25 mm)</p> <p>2.5.f. Syringes of 0.4 ml</p> <p>2.5.f.1. With needle: 23G × 1" (0.6 × 25 mm)</p>			
<p>BD SoloShot™ IX Syringe</p> <p>Basic UDI-DI: 038290MJTMIQJVL</p>	Class IIa	N/A	<p>Certificate reference: #95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A.</p>

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
<p>Listed on the MDD certificate as:</p> <p><i>Syringes with needle; Sterile auto-disable two piece syringes with integrated cannula BD SoloShot TM IX</i></p> <p>Sizes:</p> <p><i>Syringes of: 0.5 ml</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>With needle: 24G × ¾" (0.55 × 19 mm)</i></p> <p><i>With needle: 24G × 1" (0.55 × 25 mm)</i></p> <p><i>With needle: 25G × ⅝" (0.5 × 16 mm)</i></p> <p><i>With needle: 25G × 1" (0.5 × 25 mm)</i></p> <p><i>Syringes of: 1 ml</i></p> <p><i>With needles: 22G × 1" (0.7 × 25 mm)</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p>			<p>C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>
<p>BD Flu+ Syringe</p> <p>Basic UDI-DI: 038290VFCLOUDWGE</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes with needle; Sterile two-piece syringes with integrated cannula BD Flu+</i></p> <p>Sizes:</p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>With needle: 25G × ⅝" (0.5 × 16 mm)</i></p> <p><i>With needle: 25G × 1" (0.5 × 25 mm)</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>0.1 – 1 ml</i></p>	<p>Class IIa</p>	<p>N/A</p>	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318, to be read in conjunction with amending AEMPS letter dated 20 October 2021 regarding and change submitted 20 August 2021, S / REF: 90802</p> <p>MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p><i>With needle: 25G x 1" (0.5 x 25 mm) 0.1 – 1 ml</i></p>			
<p>BD Emerald TM Syringe (without needle)</p> <p>Basic UDI-DI: 038290PDVJVUZUKV</p> <p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle and syringes with blunt fill needle; Sterile syringes without needle; Sterile three-piece syringes BD Emerald™</i></p> <p><i>Sizes:</i></p> <p><i>Syringes of: 2 ml</i></p> <p><i>Syringes of: 5 ml</i></p>	<p>Class I device placed on the market in sterile condition</p> <p>Class I device with a measuring function</p>	<p>N/A</p>	<p>Certificate reference:</p> <p>#2000 06 0272 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>
<p>BD Emerald™ Syringe (with needle)</p> <p>Basic UDI-DI (Microlance needle): 038290JGXPPFFCE5</p> <p>Listed on the MDD certificate as:</p> <p><i>Sterile three-piece syringes with needle BD Emerald™;</i></p> <p><i>Syringes of 2 ml and needle side by side:</i></p> <p><i>With needle: 21G x 1 ½" (0.8 x 40 mm)</i></p> <p><i>With needle: 23G x 1" (0.6 x 25 mm)</i></p> <p><i>Syringes of 2 ml and pre-attached needle</i></p> <p><i>With needle: 21G x 1 ½" (0.8 x 40 mm)</i></p>	<p>Class IIa</p>	<p>N/A</p>	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<p><i>With needle: 23G × 1" (0.6 × 25 mm)</i> <i>With needle: 25G × 5/8" (0.5 × 16 mm)</i></p> <p><i>Syringes of 5 ml and needle side by side</i> <i>With needle: 21G × 1 1/2" (0.8 × 40 mm)</i> <i>With needle: 23G × 1" (0.6 × 25 mm)</i> <i>With needle: 24G × 1" (0.55 × 25 mm)</i></p> <p><i>Syringes of 5 ml and pre-attached needle</i> <i>With needle: 21G × 1 1/2" (0.8 × 40 mm)</i> <i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>Syringes of 10 ml and needle side by side</i> <i>With needle: 21G × 1 1/2" (0.8 × 40 mm)</i> <i>With needle: 21G × 1" (0.8 × 25 mm)</i></p> <p><i>Syringes of 10 ml and pre-attached needle</i> <i>With needle: 21G × 1 1/2" (0.8 × 40 mm)</i></p>			
<p>BD Emerald™ Non Sterile Bulk Syringe (without needle)</p> <p>Basic UDI-DI: 038290ZDNJQVHDJV</p>	<p>Class I device with a measuring function</p>	<p>N/A</p>	<p>Certificate reference: #99 03 0213 CP; expiry date 2024-05-26, NB# 0318</p>

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
<p>Listed on the MDD certificate as:</p> <p><i>Syringes without needle; Non-sterile syringes without needle; Three-piece bulk syringes BD Emerald™</i></p> <p>Sizes:</p> <p><i>Syringes of: 2 ml</i></p> <p><i>Syringes of: 5 ml</i></p> <p><i>Syringes of: 10 ml</i></p>			<p>MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>
<p>BD Emerald™ PRO Syringe (with needle)</p> <p>Basic UDI-DI (Microlance needle): 038290GRPEWKKVF5</p> <p>Listed on the MDD certificate as:</p> <p><i>Sterile three-piece syringes with reuse prevention and needle BD Emerald™ PRO;</i></p> <p><i>Syringes of 2 ml and needle side by side</i></p> <p><i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>Syringes of 5 ml and needle side by side</i></p> <p><i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i></p> <p><i>With needle: 23G × 1" (0.6 × 25 mm)</i></p> <p><i>Syringes of 10 ml and needle side by side</i></p>	Class IIa	N/A	<p>Certificate reference:</p> <p>#95 06 0006 CP; expiry date 2024-05-26, NB# 0318</p> <p>MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)</p>

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
<i>With needle: 21G × 1 ½" (0.8 × 40 mm)</i>			
BD Blunt Fill Needle Basic UDI-DI: 38290NGGTJRVACW Listed on the MDD certificate as: <i>Blunt fill needle; Sterile needle to aspirate pharmaceutical fluids from vials or ampoules; Needle BD Blunt Fill Needle</i> <i>Size: 18G × 1 ½" (1.2 × 40 mm)</i>	Class I device placed on the market in sterile condition	N/A	Certificate reference: #2015 03 0838 CP; expiry date 2024-05-26, NB# 0318 MDD Certificate issued to BECTON DICKINSON S.A., C/ Mequinenza, s/n. E-22520 FRAGA (Huesca) España (Part of the same larger organisation)
N/A	N/A	N/A	N/A

Confirmation Letter Revision History

Date	Action
2024/03/12	Initial issue