

Bastos Viegas S.A. Avenida da Fábrica, 298, 4560-164 Guilhufe-Penafiel. Portugal

01st September 2023

Confirmation Letter Reference: CLNB1639 - ES/MAD/300002696

To whom it may concern,

Confirmation of receipt of a formal application and conclusion of written agreement 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, SGS Belgium NV, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 1639 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:

Bastos Viegas S.A.
Avenida da Fábrica, 298,
4560-164
Guilhufe-Penafiel.
Portugal
SRN: PT-MF-000002795 + PT-PT-000002808 + PT-IM-000002801

The devices covered by the formal application and the written agreement mentioned above are shown at the end of this letter.

In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired prior to the publication of the Regulation EU 2023/607 on 15th March 2023. this letter also confirms that:

- The manufacturer submitted the MDR application and signed the written agreement by the date of MDD/AIMDD certificate expiry;
- The certificates expired after 26th May 2021 by course of time and were valid at the date of their expiry neither having been suspended nor withdrawn.

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 120.3 of MDR (as amended by EU 2023/607), are shown below:

• 26th May 2026 for Class III custom-made implantable devices

SGS Belgium NV

Certification and Business Enhancement Registered Office: Noorderlaan 87 BE-2030 Antwerpen t +32 (0)3 545 48 48 f +32 (0)3 545 48 49 Boulevard International/Internationalelaan 55D BE-1070 Brussels t+32 (0)2 556 00 40 f +32 (0)3 545 48 49 www.be.sgs.com



- 31st December 2027 for Class III devices and Class IIb implantable devices excluding Wellestablished technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31st December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- 31st 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 SGS Belgium NV 1639,

pp [Jérôme JADOT]

Virginie SILORET

Global Medical Device Certification Manager

ser letter Regulation Confirmation letter Regulation Email: Virginie.siloret@sgs.com Phone: +41 22 739 98 58



Devices covered by this letter:

Device name	MDR Device classification & name	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
Sterile and non-sterile, single use non-woven dressings, with or without X-Ray thread.	Class IIa: Sterile and non-sterile, single use, invasive, nonwoven dressings, with or without X-Ray thread. Class IIa: Sterile and non-sterile, single use, invasive, texart dressings.	N/A	Certificate ES19/86750; NB1639
Sterile and non-sterile, single use gauze dressings, with or without X-Ray thread.	Class IIa: Sterile and non-sterile, single use, invasive gauze dressings, with or without X-Ray thread. Class IIa: Sterile and non-sterile, single use, ribbon gauze dressings. Class IIa: Sterile and non-sterile, single use, vaginal tampons.	N/A	Certificate ES19/86750; NB1639
Sterile and non-sterile single use plastic instruments: stylet intended to monitor and explore wounds; curette intended for scraping, debriding and/or cleaning of	Class IIa: Sterile and non-sterile, single use, stylet. Class IIa: Sterile and non-sterile, single use, curettes. Class IIa:	N/A	Certificate ES19/86750; NB1639

SGS Belgium NV

Certification and Business Enhancement Registered Office: Noorderlaan 87 BE-2030 Antwerpen t+32 (0)3 545 48 48 f+32 (0)3 545 48 49 Boulevard International/Internationalelaan 55D BE-1070 Brussels t+32 (0)2 556 00 40 f+32 (0)3 545 48 49 www.be.sqs.com



Device name	MDR Device classification & name	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
biological tissues and wounds; needle-holders intended to hold and guide suture needles securely for suturing; roux retractor intended to hold back tissues and organs, in order to expose the surgical field; invasive forceps for surgical procedures.	Sterile and non-sterile, single use, needle holders. Class IIa: Sterile and non-sterile, singleuse, retractors. Class IIa: Sterile and non-sterile single use haemostatic forceps. Class IIa: Sterile and non-sterile, singleuse, auxiliary forceps. Class IIa: Sterile and non-sterile, singleuse, standard forceps. Class IIa: Sterile and non-sterile, singleuse, standard forceps. Class IIa: Sterile and non-sterile, singleuse, sponge forceps. Class IIa: Sterile and non-sterile, singleuse, sponge forceps.	Regulation	
Sterile and non-sterile, single use surgical bowls, trays, basin, basin liner, pitcher, lids and bowl sets for temporary storage and transport of organs and tissues during surgical procedures.	Class IIa: Sterile and non-sterile, single use, surgical bowls, trays, basins, basin liners, pitchers and lids.	N/A	Certificate ES19/86750; NB1639
	Class III: Sterile, single use, procedure packs.	N/A	Certificate ES19/86750; NB1639
Sterile, single use surgical and procedure sets and packs.	Class III: Sterile, single use, procedure packs.	N/A	Certificate ES19/86750; NB1639
Sterile, single use procedure sets		N/A	Certificate ES19/86752; NB1639

SGS Belgium NV

Certification and Business Enhancement Registered Office: Noorderlaan 87 BE-2030 Antwerpen t+32 (0)3 545 48 48 f+32 (0)3 545 48 49 Boulevard International/Internationalelaan 55D BE-1070 Brussels t+32 (0)2 556 00 40 f+32 (0)3 545 48 49 www.be.sqs.com



Device name	MDR Device classification & name	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
Sterile, single use non-	Class Is:	N/A	Certificate
invasive non-woven	Sterile, single use, non-	N/A	ES19/86752;
dressings.	invasive, nonwoven dressings.		NB1639
Sterile, single use orthopedic padding, elastic and tubular bandages.	Class Is: Sterile, single use, orthopaedic paddings. Class Is: Sterile, single use, conforming bandages. Class Is: Sterile, single use, support bandages. Class Is: Sterile, single use, crepe bandages. Class Is: Sterile, single use, short stretch bandages. Class Is: Sterile, single use, short stretch bandages. Class Is: Sterile, single use, stockinettes.	N/A SUITATION	Certificate ES19/86752; NB1639
Sterile, single use non-	Class Is:		Certificate
adherent wound	Sterile, single use, non-	N/A	ES19/86752;
dressings.	adherent dressings.		NB1639
Sterile, single use absorbent pads, maternity pads and first aid dressings.	Class Is: Sterile, single use, absorbent pads. Class Is: Sterile, single use, maternity pads. Class Is: Sterile, single use, first-aid dressings.	N/A	Certificate ES19/86752; NB1639
Sterile, single use non- invasive, gauze dressings, eye pads.	Class Is: Sterile, single use, eye pads. Class Is:	N/A	Certificate ES19/86752; NB1639



Device name	MDR Device classification & name	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
	Sterile, single use, gauze bandages.		23/0
Sterile, single use, non- invasive forceps and umbilical cord clamps.	Class Is: Sterile, single-use, non-invasive forceps, towel clamps and tube clamps.	N/A	Certificate ES19/86752; NB1639
	Class Is: Sterile, single-use, umbilical cord clamp.	N/A	Certificate ES19/86752; NB1639
Sterile, single use, tongue depressor.	Class Is: Sterile, single use, tongue depressors.	N/A	Certificate ES19/86752; NB1639
Sterile, single use eye shield.	Class Is: Sterile, single use, eye sh <mark>ield</mark> s.	N/A	Certificate ES19/86752; NB1639
Sterile, single use surgical drapes and draping sets.	Class Is: Sterile, single use, draping sets. Class Is: Sterile, single use, surgical drapes. Class Is: Sterile, single use, medical equipment covers.	N/A	Certificate ES19/86752; NB1639
Sterile, single use operating room towels, towel clamps, incise drapes, instruments pouches, fluid pouches, adhesive tape for operations and surgical absorbent pads.	Class Is: Sterile, single use, non- invasive, nonwoven and Texart OR towels and absorbent pads. Class Is: Sterile, single-use, non-invasive forceps, towel clamp and tube clamp. Class Is: Sterile, single use, incise drapes. Class Is:	N/A	Certificate ES19/86752; NB1639



Device name	MDR Device classification & name	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
	Sterile, single use, instrument and fluid pouches. Class Is: Sterile, single use, operation tape.		11/2033/0
. • .	Class Is: Sterile, single use, protection sheets. Class Is: Sterile, single use, baby blankets.	N/A	Certificate ES19/86752; NB1639
Sterile, single use disinfectant applicators.	Class Is: Sterile, single use, foam applicators.	N/A	Certificate ES19/86752; NB1639
Sterile, single use plastic skin staple remover.	Class Is: Sterile, single use, skin staple remover.	N/A	Certificate ES19/86752; NB1639
Sterile, single use guidewire bowls for preparation and temporary storage of guidewire while keeping it in place and hydrated.	Class Is: Sterile, single use, guidewire bowls.	N/A	Certificate ES19/86752; NB1639
Sterile, single use, measuring medicine cups for medicine administration.	Class Is/m & class Im: Sterile and non-sterile, single use, medicine cups.	N/A	Certificate ES19/86753; NB1639
Non sterile, single use, measuring medicine cups for medicine administration.		N/A	Certificate ES19/86751; NB1639

Confirmation Letter Revision History



Date	NB internal reference traceable to each version of the letter	Action
01/09/2023	Version 1	Initial issue

GGS HB 1639. Confirmation Letter Regulation HU 2023/601

SGS Belgium NV

Certification and Business Enhancement Registered Office: Noorderlaan 87 BE-2030 Antwerpen t +32 (0)3 545 48 48 f +32 (0)3 545 48 49 Boulevard International/Internationalelaan 55D BE-1070 Brussels t+32 (0)2 556 00 40 f +32 (0)3 545 48 49 www.be.sgs.com

Member of the SGS Group