EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60139535 0001

Report No.: 26300232 017

Manufacturer:
ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products: (see attachments for products and sites included)

Replaces EC Certificate, Registration No.: DD 60117020 0001

Expiry Date: 2024-05-27

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2019-06-09

Date: 2019-05-27

Notified Body

[redacted]

TÜV Rheinland LGA Products GmbH

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer:
ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

- Sterile and non-sterile cutting gauze
- Non-sterile dressing gauze
- Sterile and non-sterile gauze swabs
  (with or without X-ray thread)
- Sterile and non-sterile gauze lap sponges
  (with X-ray thread/ with X-ray chip)
- Sterile and non-sterile gauze balls
  (with or without X-ray thread)
- Sterile and non-sterile gauze rolls
  (with or without X-ray thread)
- Sterile and non-sterile non-woven swabs
  (with or without X-ray thread)
- Sterile paraffin gauze dressings
- Sterile three-way stopcocks
- Sterile transfusion sets for single use
- Sterile infusion sets for single use
- Sterile extension tubes for infusion pump

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

No [redacted]

Date: 2019-05-27

Ra [redacted]

TÜVRheinland®

Doc. 2/6, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer:
ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

- Sterile endotracheal tubes
- Sterile tracheostomy tubes
- Sterile breathing circuits
- Sterile catheter mounts
- Non-sterile anaesthetic masks
- Sterile laryngeal masks
- Sterile oxygen masks
- Sterile Multi-Vent masks
- Sterile non-rebreath masks
- Sterile nebulizer masks
- Sterile nasal oxygen cannulas
- Sterile nebulizer sets
- Sterile oxygen tubing
- Sterile suction catheters
- Sterile abdominal drains
- Sterile feeding tubes
- Sterile stomach and duodenal tubes
- Sterile urology catheters
- Sterile surgical suction sets
- Sterile surgical suction cannulas
- Sterile syringes for single use

TÜV Rheinland LGA Products GmbH TÜVRheinland Zertifizierungsstelle

Notified Body

[redacted]

Rafa

Date: 2019-05-27

TÜVRheinland®

Doc. 3/6, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer:
ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

- Sterile insulin syringes
- Sterile tuberculin syringes
- Sterile hypodermic needles
- Sterile insulin pen needles
- Sterile blood lancets
- Sterile IV cannulas
- Sterile needle free valves
- Sterile surgical gloves

TÜV Rheinland LGA Products GmbH TÜVRheinland® Zertifizierungsstelle

Notif[redacted]

Date: 2019-05-27

Rafa[redacted]

TÜVRheinland®

Doc. 4/6, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer: ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

For the following medical devices the scope covers only the aspects of manufacture concerned with securing and maintaining sterile conditions:

- Adhesive cannulla fixation dressings
- Adhesive wound dressings
- Eye pads
- Incise films
- Transparent film dressings
- Foam dressings
- Absorbent wound dressings
- Surgical gowns
- Surgical drapes
- Sets of surgical drapes
- Fluid collection pouches
- Nelaton catheters

TÜV Rheinland LGA Products GmbH TÜVRheinland Zertifizierungsstelle

Noti[redacted]

Date: 2019-05-27

Raf[redacted]

TÜVRheinland®

Doc. 5/6, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer: ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

For the following medical devices the scope covers only the aspects of manufacture concerned with securing and maintaining sterile conditions:

- Vaginal speculums
- Cervical brushes
- Urine bags
- Tongue depressors
- Guedel airways
- Intubation stylets
- Endotracheal tube holders
- Suction tubes
- Withdrawal cannulas
- Alginate dressings
- Cannula stoppers
- Umbilical cord clamps

TÜV Rheinland LGA Products GmbH TÜVRheinland Zertifizierungsstelle

Notifie[redacted]

Date: 2019-05-27

Rafal [redacted]

TÜVRheinland®

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60139535 0001
Report No.: 26300232 017

Manufacturer:
ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Polska

Products included:

ZARYS International Group
Spolka z ograniczona odpowiedzialnoscia,
spolka komandytowa
ul. Gustawa Eiffel'a 15
44-109 Gliwice
Poland

Activity: Production

TÜV Rheinland LGA Products GmbH TÜVRheinland Zertifizierungsstelle

Notifie[redacted]

Date: 2019-05-27

Rafal Byczkowski

CE sertifikatas
93/42/EEC Direktyva V Priedas
Gamybos kokybės užtikrinimas
Medicinos prietaisai

Registracijos Nr.: DD 60139535 0001

Ataskaitos Nr.: 26300232 017

Gamintojas: Zarys International Group
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Produktai: (Žr. Priedus dėl produktų ir įtrauktų vietų)
Pakeičia CE sertifikatą Nr. DD 60100191 0001

Galioja iki: 2024-05-27

Notifikuota įstaiga patvirtina minėtos įmonės įdiegtą ir taikomą kokybės valdymo sistemą. Direktyvos 93/42/EEB V Priedo reikalavimai buvo įvykdyti. Minėtas gamintojas yra įdiegės ir taiko kokybės užtikrinimo sistemą, kuri yra periodiškai tikrinama, pagal minėtos direktyvos V Priedo 4 skyrių. IIb ir III klasės prietaisų, minimų šiame sertifikate, perkėlimui į rinką būtinas EB tipo patikros sertifikatas pagal III Priedą.

Galioja nuo: 2019-06-09

Data: 2019-05-27

Notifikuota įstaiga

/parašas/ /antspaudas/
Rafal Byczkowski

TÜV Rheinland LGA Products GmbH – Tillystrasse 2 – 90431 Nürnberg

TÜV Rheinland LGA Products GmbH yra Notifikuota įstaiga pagal Direktyvą 93/42/EEB dėl medicinos prietaisų, kurios identifikavimo numeris: 0197.

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60117020 0001**

Ataskaitos Nr.: **26300232 005**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Įtraukti Produktai:

- Sterili ir nesterili kerpama marlė
- Nesterilūs marliniai tvarsčiai
- Sterilus/nesterilus marlinis tvarstis

(su/be rentgenokontrastine linija)

- Sterilūs ir nesterilūs marlės rutuliai

(su/be rentgenokontrastine linija)

- Sterilūs ir nesterilūs marlės ritiniai

(su/be rentgenokontrastine linija)

- Sterilūs ir nesterilūs neaustinės medžiagos tvarsčiai

(su/be rentgenokontrastine linija)

- Sterilūs parafininiai tvarsčiai
- Sterilūs trijų krypčių kraneliai
- Sterilios vienkartinės transfuzinės sistemos
- Sterilios vienkartinės infuzinės sistemos
- Sterilios prailginimo linijos infuzinėms pompoms

Data: 2019-05-27

Notifikuota įstaiga

/parašas/ /antspaudas/
Rafal Byczkowski

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60117020 0001**

Ataskaitos Nr.: **26300232 005**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Įtraukti Produktai:

- Sterilūs endotrachėjiniai vamzdeliai
- Sterilūs tracheostominiai vamzdeliai
- Sterilio kvėpavimo grandinės
- Sterilūs prailgintojai prie intub. Vamzdelio
- Sterilios laringinės kaukės
- Sterilios deguonies kaukės
- Sterili kaukė su rezervuaru
- sterili aerozolinė kaukė
- Sterlios nosies deguonies kaniulės
- Sterilūs aerozoliniai rinkiniai
- Sterilūs deguonies vamzdeliai
- Sterilūs atsiurbimo kateteriai
- Sterilūs chirurginiai atsiurbimo rinkiniai
- Sterilios chirurginės atsiurbimo kaniulės
- Sterilūs maitinimo zondai
- Sterilūs skrandžio ir duodenaliniai zondai
- Sterilūs urologiniai kateteriai
- Sterilūs vienkartiniai švirkštai

Data: 2019-05-27 **Notifikuota įstaiga**

/parašas/ /antspaudas/
Rafal Byczkowski

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60117020 0001**

Ataskaitos Nr.: **26300232 005**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Įtraukti Produktai:

- Sterilūs insulininiai švirkštai
- Sterilūs tuberkuliniai švirkštai
- Sterilios hipoderm· nės adatos
- Sterilios insulininių penų adatos
- Sterilūs kraujo lancetai
- Sterilios intraveninės kaniulės
- Sterilūs beadatiniai vožtuvai
- Sterilios chirurginės pirštinės

Data: 2019-05-27

Notifikuota įstaiga

/parašas/ /antspaudas/
Rafal Byczkowski

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60117020 0001**

Ataskaitos Nr.: **26300232 005**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Žemiau nurodytiems medicinos prietaisams reikalavimai taikomi tik gamybos aspektams, susijusiems su sterilių sąlygų kūrimų bei išlaikymu:

- Lipnus kaniulių fiksatoriai
- Lipnūs žaizdų tvarsčiai
- Akių tamponėliai
- Pjūvio juosta
- Permatomi juostiniai tvarsčiai
- Putų tvarsčiai
- Absorbuojantys žaizdų tvarsčiai
- Chirurginiai chalatai
- Chirurginiai dangalai
- Skysčių rinkimo talpos
- Nelaton kateteriai

Data: 2019-05-27

Notifikuota įstaiga

/parašas/ /antspaudas/
Rafal Byczkowski

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60117020 0001**

Ataskaitos Nr.: **26300232 005**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Įtraukti Produktai:
Žemiau nurodytiems medicinos prietaisams reikalavimai taikomi tik gamybos aspektams, susijusiems su sterilių sąlygų kūrimų bei išlaikymu:

- Vaginalinės spekulės
- Gimdos kaklelio šepetėliai
- Šlapimo maišeliai
- Liežuvio prispaudėjai
- Orofaringiniai vamzdeliai
- Intubaciniai stiletai
- Tracheostominio vamzdelio laikikliai
- Atsiurbimo vamzdeliai
- Ištraukimo kaniulės
- Alignato tvarsčiai
- Kaniulių kamštukai
- Umbilikaliniai spaustukai

Data: 2019-05-27 **Notifikuota įstaiga**

/parašas/ /antspaudas/

Rafal Byczkowski

TÜV Rheinland
LGA Products GmbH
Tillystrasse 2, 90431 Nürnberg

Priedas prie sertifikato
Registracijos Nr.: **DD60139535 0001**

Ataskaitos Nr.: **26300232 017**

Gamintojas: **Zarys International Group**
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Pod Borem 18
41-808 Zabrze
Lenkija

Įtrauktos vietos:

Zarys International Group
Spolka z organiczona
odpowiedzialnoscia
spolka komandytowa
ul. Gustawa Eiffel'a 15
44-109 Gliwice
Lenkija

Data: 2019-05-27

Notifikuota įstaiga

/parašas/ /antspaudas/
Rafal Byczkowski

Business Stream Products
Certification Department

TÜVRheinland®
LGA
Precisely Right.

TÜV Rheinland LGA Products GmbH • 51105 Köln

ZARYS International Group spółka z ograniczoną odpowiedzialnością spółka komandytowa
ul. Pod Borem 18,
41-808 Zabrze,
Poland

Contact

Tel. +49 911 655-5225
Mail: medical-products@de.tuv.com

Date May 15, 2024

Notified Body Confirmation Letter
Reference. : ZARYS_PLA0_HZ_2024-05-10/ 84965323

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 **TÜV Rheinland LGA Products GmbH**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **0197** 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:

ZARYS International Group spółka z ograniczoną odpowiedzialnością spółka komandytowa
ul. Pod Borem 18,
41-808 Zabrze,
Poland
SRN Number: PL-MF-000000410

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 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 May 26, 2021 but before March 20, 2023 without having been withdrawn, this letter also confirms that the manufacturer either 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 March 20, 2023 for the relevant devices.

TÜV Rheinland
LGA Products GmbH

Am Grauen Stein
51105 Köln
Germany

Headquarter

Tillystraße 2
90431 Nuremberg

Phone. +49 911 655 5225
Fax +49 911 655 5226
service@de.tuv.com
www.tuv.com/safety

Board of Management

Dipl.-Ing.
Thomas Weigand, Spokesman

Dipl.-Kfm.
Dr. Jörg Schlösser

Nuremberg HRB 26013
VAT No.: DE 811835490

Chairman of the
Supervisory Board

Dr.-Ing. Michael Fübi

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:

- May 26, 2026 for Class III custom-made implantable devices
- December 31, 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)
- December 31, 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- December 31, 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

[redacted] Malgorzata Blazniak
[redacted] 4.05.15 13:38:01
[redacted] 00'

AUDIT_CERT_REVIEW
Certification body

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
GAZA lux S **Cutting gauze, sterile** **Basic UDI-DI:** **59079968M02010101-ERR**	Class IIa	GAZA lux S Cutting gauze, sterile	DD 1023663-1 NB 0197
GAZA lux S **Cutting gauze, sterile** **Basic UDI-DI:** **59079968M02010101-SSM**	Class IIa	GAZA lux S Cutting gauze, sterile	DD 1023663-1 NB 0197
GAZA lux **Cutting gauze, non-sterile** **Basic UDI-DI:** **59079968M02010101-NSB**	Class IIa	GAZA lux Cutting gauze, non-sterile	DD 1023663-1 NB 0197
GAZA lux **Dressing gauze, non-sterile** **Basic UDI-DI:** **59079968M020107DG**	Class IIa	GAZA lux Dressing gauze, non-sterile	DD 1023663-1 NB 0197
KOMPRI lux S	Class IIa	KOMPRI lux S	DD 1023663-1 NB 0197

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**
Gauze swabs without X-ray thread, sterile **Basic UDI-DI: 59079968M02010201-ES4**		Gauze swabs without X-ray thread, sterile	
KOMPRI lux S **Gauze swabs without X-ray thread, sterile** **Basic UDI-DI: 59079968M02010201-SSY**	Class IIa	KOMPRI lux S Gauze swabs without X-ray thread, sterile	DD 1023663-1 NB 0197
KOMPRI lux S **Gauze swabs with X-ray thread, sterile** **Basic UDI-DI: 59079968M02010202-ES9**	Class IIa	KOMPRI lux S Gauze swabs with X-ray thread, sterile	DD 1023663-1 NB 0197
KOMPRI lux S **Gauze swabs with X-ray thread, sterile** **Basic UDI-DI: 59079968M02010202-ST5**	Class IIa	KOMPRI lux S Gauze swabs with X-ray thread, sterile	DD 1023663-1 NB 0197
KOMPRI lux **Gauze swabs without X-ray thread, non-sterile** **Basic UDI-DI: 59079968M02010201-NSN**	Class IIa	KOMPRI lux Gauze swabs without X-ray thread, non-sterile	DD 1023663-1 NB 0197
KOMPRI lux **Gauze swabs with X-ray thread, non-sterile** **Basic UDI-DI: 59079968M02010202-NST**	Class IIa	KOMPRI lux Gauze swabs with X-ray thread, non-sterile	DD 1023663-1 NB 0197
SERVI lux S **Gauze lap sponges with X-ray thread, sterile** **Basic UDI-DI: 59079968M02010302-ESL**	Class IIa	SERVI lux S Gauze lap sponges with X-ray thread, sterile	DD 1023663-1 NB 0197
SERVI lux S **Gauze lap sponges with X-ray thread, sterile** **Basic UDI-DI: 59079968M02010302-STG**	Class IIa	SERVI lux S Gauze lap sponges with X-ray thread, sterile	DD 1023663-1 NB 0197
SERVI lux S **Gauze lap sponges with X-ray chip, pre-washed, sterile**	Class IIa	SERVI lux S Gauze lap sponges with X-ray chip, pre-washed, sterile	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968M02010302-PE86			
SERVI lux S Gauze lap sponges with X-ray chip, pre-washed, sterile **Basic UDI-DI: 59079968M02010302-PS92**	Class IIa	SERVI lux S Gauze lap sponges with X-ray chip, pre-washed, sterile	DD 1023663-1 NB 0197
SERVI lux Gauze lap sponges with X-ray thread, non-sterile **Basic UDI-DI: 59079968M02010302-NT6**	Class IIa	SERVI lux Gauze lap sponges with X-ray thread, non-sterile	DD 1023663-1 NB 0197
SERVI lux Gauze lap sponges with X-ray chip, pre-washed, non-sterile **Basic UDI-DI: 59079968M02010302-PN8Q**	Class IIa	SERVI lux Gauze lap sponges with X-ray chip, pre-washed, non-sterile	DD 1023663-1 NB 0197
TUPFER lux S Gauze balls without X-ray thread, sterile **Basic UDI-DI: 59079968M02010501-ET5**	Class IIa	TUPFER lux S Gauze balls without X-ray thread, sterile	DD 1023663-1 NB 0197
TUPFER lux S Gauze balls without X-ray thread, sterile **Basic UDI-DI: 59079968M02010501-STZ**	Class IIa	TUPFER lux S Gauze balls without X-ray thread, sterile	DD 1023663-1 NB 0197
TUPFER lux S Gauze balls with X-ray thread, sterile **Basic UDI-DI: 59079968M02010502-ETA**	Class IIa	TUPFER lux S Gauze balls with X-ray thread, sterile	DD 1023663-1 NB 0197
TUPFER lux S Gauze balls with X-ray thread, sterile **Basic UDI-DI: 59079968M02010502-SU6**	Class IIa	TUPFER lux S Gauze balls with X-ray thread, sterile	DD 1023663-1 NB 0197
TUPFER lux Gauze balls without X-ray thread, non-sterile	Class IIa	TUPFER lux Gauze balls without X-ray thread, non-sterile	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968M02010501-NTP			
TUPFER lux Gauze balls with X-ray thread, non-sterile **Basic UDI-DI: 59079968M02010502-NTU**	Class IIa	TUPFER lux Gauze balls with X-ray thread, non-sterile	DD 1023663-1 NB 0197
SETON lux S Gauze rolls without X-ray thread, sterile **Basic UDI-DI: 59079968M02010701-SUP**	Class IIa	SETON lux S Gauze rolls without X-ray thread, sterile	DD 1023663-1 NB 0197
SETON lux S Gauze rolls with X-ray thread, sterile **Basic UDI-DI: 59079968M02010702-ETY**	Class IIa	SETON lux S Gauze rolls with X-ray thread, sterile	DD 1023663-1 NB 0197
SETON lux S Gauze rolls with X-ray thread, sterile **Basic UDI-DI: 59079968M02010702-SUU**	Class IIa	SETON lux S Gauze rolls with X-ray thread, sterile	DD 1023663-1 NB 0197
SETON lux Gauze rolls without X-ray thread, non-sterile **Basic UDI-DI: 59079968M02010701-NUD**	Class IIa	SETON lux Gauze rolls without X-ray thread, non-sterile	DD 1023663-1 NB 0197
SETON lux Gauze rolls with X-ray thread, non-sterile **Basic UDI-DI: 59079968M02010702-NUJ**	Class IIa	SETON lux Gauze rolls with X-ray thread, non-sterile	DD 1023663-1 NB 0197
NONVI lux S Non-woven swab, sterile **Basic UDI-DI: 59079968M02020101-ESA**	Class IIa	NONVI lux S Non-woven swabs, sterile	DD 1023663-1 NB 0197
NONVI lux S Non-woven swab, sterile **Basic UDI-DI: 59079968M02020101-ST6**	Class IIa	NONVI lux S Non-woven swabs, sterile	DD 1023663-1 NB 0197
NONVI lux S Non-woven swabs with X-ray thread, sterile	Class IIa	NONVI lux S Non-woven swabs with X-ray thread, sterile	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968M02020102-ESF			
NONVI lux S Non-woven swabs with X-ray thread, sterile **Basic UDI-DI: 59079968M02020102-STB**	Class IIa	NONVI lux S Non-woven swabs with X-ray thread, sterile	DD 1023663-1 NB 0197
NONVI lux Non-woven swabs, non-sterile **Basic UDI-DI: 59079968M02020101-NSU**	Class IIa	NONVI lux Non-woven swabs, non-sterile	DD 1023663-1 NB 0197
NONVI lux Non-woven swabs with X-ray thread, non-sterile **Basic UDI-DI: 59079968M02020102-NSZ**	Class IIa	NONVI lux Non-woven swabs with X-ray thread, non-sterile	DD 1023663-1 NB 0197
paraffiNET Paraffin gauze dressing, sterile **Basic UDI-DI: 59079968M020302DG**	Class IIa	paraffiNET Paraffin gauze dressing, sterile	DD 1023663-1 NB 0197
SANVIflon I.V. cannula SANVIflon S I.V. cannula without injection port SANVIflon premium I.V. cannula SANVIflon safe Safety I.V. cannula **Basic UDI-DI: 59079968C0101017F**	Class IIa	SANVIflon I.V. cannula SANVIflon S I.V. cannula without injection port SANVIflon premium I.V. cannula SANVIflon safe Safety I.V. cannula	DD 1023663-1 NB 0197
OXYGEN TUBING **Basic UDI-DI: 59079968R03010204LA**	Class IIa	OXYGEN TUBING	DD 1023663-1 NB 0197
NEBULIZER with T F/M adaptor, mouthpiece, corrugated tube and tubing **Basic UDI-DI: 59079968R030103-FMV5**	Class IIa	NEBULIZER with T F/M adaptor, mouthpiece, corrugated tube and tubing	DD 1023663-1 NB 0197
NEBULIZER mask with tubing **Basic UDI-DI: 59079968R030103-MMN**	Class IIa	NEBULIZER with mask and tubing	DD 1023663-1 NB 0197

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
OXYGEN MASK with tubing **Basic UDI-DI: 59079968R03010201L4**	Class IIa	OXYGEN MASK with tubing	DD 1023663-1 NB 0197
NON-REBREATHER MASK with tubing **Basic UDI-DI: 59079968R03010206LE**	Class IIa	NON-REBREATHER MASK with tubing	DD 1023663-1 NB 0197
VENTURI MASK with adjustable diluter and tubing **Basic UDI-DI: 59079968R03010202-AXK**	Class IIa	VENTURI MASK with adjustable diluter and tubing	DD 1023663-1 NB 0197
NASAL OXYGEN CANNULA for adults **NASAL OXYGEN CANNULA for children** **NASAL OXYGEN CANNULA for infants** **Basic UDI-DI: 59079968R03010203L8**	Class IIa	NASAL OXYGEN CANNULA for adults NASAL OXYGEN CANNULA for children NASAL OXYGEN CANNULA for infants	DD 1023663-1 NB 0197
SUCTION CATHETER **SUCTION CATHETER IDEAL** **SUCTION CATHETER with vacuum control** **SUCTION CATHETER with vacuum control IDEAL** **Basic UDI-DI: 59079968R0501QP**	Class IIa	SUCTION CATHETER SUCTION CATHETER IDEAL SUCTION CATHETER with vacuum control SUCTION CATHETER with vacuum control IDEAL	DD 1023663-1 NB 0197
Two-way Foley catheter with rubber valve (silicone-coated latex) **Basic UDI-DI: 59079968U010201-LRXH**	Class IIa	TWO-WAY FOLEY CATHETER with rubber valve	DD 1023663-1 NB 0197
Two-way Foley catheter with plastic valve (silicone-coated latex) **Basic UDI-DI: 59079968U010201-LPXD**	Class IIa	TWO-WAY FOLEY CATHETER with plastic valve	DD 1023663-1 NB 0197
Two-way Foley catheter with plastic valve (100% silicone, X-ray contrast)	Class IIa	TWO-WAY FOLEY CATHETER with plastic valve	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968U010201-SP6			
Three-way Foley catheter with plastic valve (silicone-coated latex) **Basic UDI-DI: 59079968U010201-3LUV**	Class IIa	THREE-WAY FOLEY CATHETER with plastic valve	DD 1023663-1 NB 0197
Three-way Foley catheter with plastic valve (100% silicone, X-ray contrast) **Basic UDI-DI: 59079968U010201-3SVB**	Class IIa	THREE-WAY FOLEY CATHETER with plastic valve	DD 1023663-1 NB 0197
Two-way Foley catheter with plastic valve, Tiemann tip (silicone-coated latex) **Basic UDI-DI: 59079968U0102R6**	Class IIa	TWO-WAY FOLEY CATHETER with plastic valve, Tiemann tip	DD 1023663-1 NB 0197
TIEMANN CATHETER **Basic UDI-DI: 59079968U010106HB**	Class IIa	TIEMANN CATHETER	DD 1023663-1 NB 0197
PEZZER CATHETER **Basic UDI-DI: 59079968U010107HD**	Class IIa	PEZZER CATHETER	DD 1023663-1 NB 0197
FEEDING TUBE **Basic UDI-DI: 59079968G02020101BU**	Class IIa	FEEDING TUBE	DD 1023663-1 NB 0197
STOMACH TUBE DUODENAL TUBE **Basic UDI-DI: 59079968G020201A3**	Class IIa	STOMACH TUBE DUODENAL TUBE	DD 1023663-1 NB 0197
SUCTION CANNULA with suction control SUCTION CANNULA without suction control **Basic UDI-DI: 59079968A06010184**	Class IIa	SUCTION CANNULA with suction control SUCTION CANNULA without suction control	DD 1023663-1 NB 0197
SUCTION CANNULA with suction control, ball tip SUCTION CANNULA without suction control, ball tip **Basic UDI-DI: 59079968A060101-BA2**	Class IIa	SUCTION CANNULA with suction control, ball tip SUCTION CANNULA without suction control, ball tip	DD 1023663-1 NB 0197

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**
SURGICAL SUCTION SET with suction control, Yankauer tip **SURGICAL SUCTION SET without suction control, Yankauer tip** **Basic UDI-DI: 59079968A060101039F**	Class IIa	SURGICAL SUCTION SET with suction control, Yankauer tip SURGICAL SUCTION SET without suction control, Yankauer tip	DD 1023663-1 NB 0197
SURGICAL SUCTION SET with suction control, funnel-funnel cut-to-fit, Yankauer tip **SURGICAL SUCTION SET without suction control, funnel-funnel cut-to-fit, Yankauer tip** **Basic UDI-DI: 59079968A06010103-FFUC**	Class IIa	SURGICAL SUCTION SET with suction control, funnel-funnel cut-to-fit, Yankauer tip SURGICAL SUCTION SET without suction control, funnel-funnel cut-to-fit, Yankauer tip	DD 1023663-1 NB 0197
SURGICAL SUCTION SET with suction control, funnel-funnel cut-to-fit, ball tip **SURGICAL SUCTION SET without suction control, funnel-funnel cut-to-fit, ball tip** **Basic UDI-DI: 59079968A06010103-FFB6J**	Class IIa	SURGICAL SUCTION SET with suction control, funnel-funnel cut-to-fit, ball tip SURGICAL SUCTION SET without suction control, funnel-funnel cut-to-fit, ball tip	DD 1023663-1 NB 0197
SUCTION TUBE funnel-funnel **Basic UDI-DI: 59079968A060304-FFG4**	Class IIa	SUCTION TUBE funnel-funnel	DD 1023663-1 NB 0197
SUCTION TUBE funnel-funnel cut-to-fit **Basic UDI-DI: 59079968A060304-FCFW**	Class IIa	SUCTION TUBE funnel-funnel cut-to-fit	DD 1023663-1 NB 0197
SUCTION TUBE funnel-Kapkon **Basic UDI-DI: 59079968A060304-FKGE**	Class IIa	SUCTION TUBE funnel-Kapkon	DD 1023663-1 NB 0197
easyWAY Three-way stopcock	Class IIa	easyWAY Three-way stopcock	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968A0703KA			
easyWAY L Three-way stopcock with extension **Basic UDI-DI: 59079968A0703-LA4**	Class IIa	easyWAY L Three-way stopcock with extension	DD 1023663-1 NB 0197
easyFLOW LINE Extension tube for infusion pump, phthalate-free **Basic UDI-DI: 59079968A03020178**	Class IIa	easyFLOW LINE Extension tube for infusion pump, phthalate-free	DD 1023663-1 NB 0197
easyFLOW LINE AMBER Extension tube for infusion pump, amber, phthalate-free **Basic UDI-DI: 59079968A030201-A8Q**	Class IIa	easyFLOW LINE AMBER Extension tube for infusion pump, amber, phthalate-free	DD 1023663-1 NB 0197
easyFLOW IS Infusion set easyFLOW IS ECO Infusion set **Basic UDI-DI: 59079968A03010103-PHT6H**	Class IIa	easyFLOW IS Infusion set easyFLOW IS ECO Infusion set	DD 1023663-1 NB 0197
easyFLOW IS Infusion set, phthalate-free easyFLOW IS ECO Infusion set, phthalate-free easyFLOW IS PREMIUM Infusion set, phthalate-free **Basic UDI-DI: 59079968A030101037U**	Class IIa	easyFLOW IS Infusion set, phthalate-free easyFLOW IS ECO Infusion set, phthalate-free easyFLOW IS PREMIUM Infusion set, phthalate-free	DD 1023663-1 NB 0197
easyFLOW IS SAFE Safety infusion set, phthalate-free easyFLOW IS SAFE PREMIUM Safety infusion set, phthalate-free **Basic UDI-DI: 59079968A03010103-SG2**	Class IIa	easyFLOW IS SAFE Safety infusion set, phthalate-free easyFLOW IS SAFE PREMIUM Safety infusion set, phthalate-free	DD 1023663-1 NB 0197
easyFLOW IS REG Infusion set with precision flow rate regulator, phthalate-free	Class IIa	easyFLOW IS REG Infusion set with precision flow rate	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968A03010103-RFY		regulator, phthalate-free	
easyFLOW IS AMBER Infusion set, amber, phthalate-free **Basic UDI-DI: 59079968A03010103-AEW**	Class IIa	easyFLOW IS AMBER Infusion set, amber, phthalate-free	DD 1023663-1 NB 0197
ENDOTRACHEAL TUBE UNCUFFED **Basic UDI-DI: 59079968R010301FQ**	Class IIa	ENDOTRACHEAL TUBE UNCUFFED	DD 1023663-1 NB 0197
ENDOTRACHEAL TUBE CUFFED **Basic UDI-DI: 59079968R010302FS**	Class IIa	ENDOTRACHEAL TUBE CUFFED	DD 1023663-1 NB 0197
REINFORCED ENDOTRACHEAL TUBE CUFFED WITH PRELOADED STYLET **Basic UDI-DI: 59079968R010302-RMF**	Class IIa	REINFORCED ENDOTRACHEAL TUBE CUFFED WITH PRELOADED STYLET	DD 1023663-1 NB 0197
BREATHING CIRCUIT FOR CHILDREN with bag, expandable, additional arm **BREATHING CIRCUIT FOR ADULTS with bag, expandable, additional arm** **Basic UDI-DI: 59079968R0201-BGG**	Class IIa	BREATHING CIRCUIT FOR CHILDREN with bag, expandable, additional arm BREATHING CIRCUIT FOR ADULTS with bag, expandable, additional arm	DD 1023663-1 NB 0197
BREATHING CIRCUIT FOR CHILDREN **BREATHING CIRCUIT FOR ADULTS** **Basic UDI-DI: 59079968R0201Q8**	Class IIa	BREATHING CIRCUIT FOR CHILDREN BREATHING CIRCUIT FOR ADULTS	DD 1023663-1 NB 0197
CATHETER MOUNT with double swivel elbow connector, smooth-bore **Basic UDI-DI: 59079968R020202-SMP**	Class IIa	CATHETER MOUNT with double swivel elbow connector, smooth-bore	DD 1023663-1 NB 0197
CATHETER MOUNT with double swivel elbow connector, expandable	Class IIa	CATHETER MOUNT with double swivel	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968R020202-ELT		elbow connector, expandable	
CATHETER MOUNT with double swivel elbow connector, corrugated **Basic UDI-DI: 59079968R020202-CLP**	Class IIa	CATHETER MOUNT with double swivel elbow connector, corrugated	DD 1023663-1 NB 0197
CATHETER MOUNT with straight connector, smooth-bore **Basic UDI-DI: 59079968R020201-SMJ**	Class IIa	CATHETER MOUNT with straight connector, smooth-bore	DD 1023663-1 NB 0197
CATHETER MOUNT with straight connector, corrugated **Basic UDI-DI: 59079968R020201-CLJ**	Class IIa	CATHETER MOUNT with straight connector, corrugated	DD 1023663-1 NB 0197
CATHETER MOUNT with straight connector, expandable **Basic UDI-DI: 59079968R020201-ELN**	Class IIa	CATHETER MOUNT with straight connector, expandable	DD 1023663-1 NB 0197
CATHETER MOUNT with elbow connector, smooth-bore **Basic UDI-DI: 59079968R0202-SHP**	Class IIa	CATHETER MOUNT with elbow connector, smooth-bore	DD 1023663-1 NB 0197
CATHETER MOUNT with elbow connector, corrugated **Basic UDI-DI: 59079968R0202-CGP**	Class IIa	CATHETER MOUNT with elbow connector, corrugated	DD 1023663-1 NB 0197
CATHETER MOUNT with elbow connector, expandable **Basic UDI-DI: 59079968R0202-EGT**	Class IIa	CATHETER MOUNT with elbow connector, expandable	DD 1023663-1 NB 0197
TRACHEOSTOMY TUBE cuffed **Basic UDI-DI: 59079968R010502G4**	Class IIa	TRACHEOSTOMY TUBE cuffed	DD 1023663-1 NB 0197
TRACHEOSTOMY TUBE uncuffed	Class IIa	TRACHEOSTOMY TUBE uncuffed	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968R010501G2			
LARYNGEAL MASK, PVC, disposable **Basic UDI-DI: 59079968R0102-PH6**	Class IIa	LARYNGEAL MASK, PVC, disposable	DD 1023663-1 NB 0197
LARYNGEAL MASK, silicone, disposable **Basic UDI-DI: 59079968R0102-SHC**	Class IIa	LARYNGEAL MASK, silicone, disposable	DD 1023663-1 NB 0197
AIR CUSHION ANAESTHETIC MASK **Basic UDI-DI: 59079968R030101-CLQ**	Class IIa	AIR CUSHION ANAESTHETIC MASK	DD 1023663-1 NB 0197
ANAESTHETIC MASK with open seal **Basic UDI-DI: 59079968R030101-OMG**	Class IIa	ANAESTHETIC MASK with open seal	DD 1023663-1 NB 0197
duoNEX **Single use syringe, 2-part** **Basic UDI-DI: 59079968A0201020101DK**	Class IIa	duoNEX Single use syringe, 2-part	DD 1023663-1 NB 0197
dicoNEX **Single use syringe, 3-part (luer)** **Basic UDI-DI: 59079968A0201020102DM**	Class IIa	dicoNEX Single use syringe, 3-part (luer)	DD 1023663-1 NB 0197
Apteczka ABC **Strzykawka 3-częściowa** **Basic UDI-DI: 59079968A0201020102DM**	Class IIa	Apteczka ABC Strzykawka 3-częściowa	DD 1023663-1 NB 0197
dicoNEX **Single use syringe, 3-part (luer lock)** **Basic UDI-DI: 59079968A0201020201DQ**	Class IIa	dicoNEX Single use syringe, 3-part (luer lock)	DD 1023663-1 NB 0197
dicoNEX **Single use amber syringe, 3-part (luer lock)** **Basic UDI-DI: 59079968A0201020201-AVY**	Class IIa	dicoNEX Single use amber syringe, 3-part (luer lock)	DD 1023663-1 NB 0197
dicoNEX	Class IIa	dicoNEX	DD 1023663-1 NB 0197

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**
Single use catheter syringe, 3-part **Basic UDI-DI: 59079968A020102037G**		Single use catheter syringe, 3-part	
dicoNEX MN Single use syringe, 3-piece with mounted needle (luer) **dicoNEX SN Single use syringe, 3-piece, with needle alongside the syringe (luer)** **Basic UDI-DI: 59079968A0201020102-IWA**	Class IIa	dicoNEX MN Single use syringe, 3-piece with mounted needle (luer) dicoNEX SN Single use syringe, 3-piece, with needle alongside the syringe (luer)	DD 1023663-1 NB 0197
dicoNEX MN Single use syringe, 3-piece with mounted needle (luer lock) **dicoNEX SN Single use syringe, 3-piece, with needle alongside the syringe (luer lock)** **Basic UDI-DI: 59079968A0201020201-IWG**	Class IIa	dicoNEX MN Single use syringe, 3-piece with mounted needle (luer lock) dicoNEX SN Single use syringe, 3-piece, with needle alongside the syringe (luer lock)	DD 1023663-1 NB 0197
dicoNEX MN Single use amber syringe, 3-piece with mounted needle (luer lock) **dicoNEX SN Single use amber syringe, 3-piece, with needle alongside the syringe (luer lock)** **Basic UDI-DI: 59079968A0201020201-IA9D**	Class IIa	dicoNEX MN Single use amber syringe, 3-piece with mounted needle (luer lock) dicoNEX SN Single use amber syringe, 3-piece, with needle alongside the syringe (luer lock)	DD 1023663-1 NB 0197
dicoSULIN Insulin syringe **Basic UDI-DI: 59079968A02010672**	Class IIa	dicoSULIN Insulin syringe	DD 1023663-1 NB 0197
dicoTUBER Tuberculin syringe **Basic UDI-DI: 59079968A02010978**	Class IIa	dicoTUBER Tuberculin syringe	DD 1023663-1 NB 0197
dispoFINE Injection needle	Class IIa	dispoFINE Injection needle	DD 1023663-1 NB 0197

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**
Basic UDI-DI: 59079968A0101010102CK			
dispoGUARD Safety injection needle **Basic UDI-DI: 59079968A0101010101CH**	Class IIa	dispoGUARD Safety injection needle	DD 1023663-1 NB 0197
dispoSULIN Insulin pen needle **Basic UDI-DI: 59079968A010101026Q**	Class IIa	dispoSULIN Insulin pen needle	DD 1023663-1 NB 0197
easyFLOW TS Transfusion set **Basic UDI-DI: 59079968A03010102-PHT66**	Class IIa	easyFLOW TS Transfusion set	DD 1023663-1 NB 0197
easyFLOW TS Transfusion set, phthalate-free easyFLOW TS PREMIUM Transfusion set, phthalate-free **Basic UDI-DI: 59079968A030101027S**	Class IIa	easyFLOW TS Transfusion set, phthalate-free easyFLOW TS PREMIUM Transfusion set, phthalate-free	DD 1023663-1 NB 0197
NEEDLE FREE VALVE blue **Basic UDI-DI: 59079968A0705KE**	Class IIa	NEEDLE FREE VALVE blue	DD 1023663-1 NB 0197
NEEDLE FREE VALVE transparent **Basic UDI-DI: 59079968A07050295**	Class IIa	NEEDLE FREE VALVE transparent	DD 1023663-1 NB 0197
NEEDLE FREE VALVE transparent with extension line, single NEEDLE FREE VALVE transparent with extension line, double NEEDLE FREE VALVE transparent with extension line, triple NEEDLE FREE VALVE transparent with extension line, quadruple **Basic UDI-DI: 59079968A070502-LCQ**	Class IIa	NEEDLE FREE VALVE transparent with extension line, single NEEDLE FREE VALVE transparent with extension line, double NEEDLE FREE VALVE transparent with extension line, triple NEEDLE FREE VALVE transparent with extension line, quadruple	DD 1023663-1 NB 0197

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
safeCARE Surgical gloves, latex, powdered **Basic UDI-DI: 59079968T01010101-RYM**	Class IIa	safeCARE Surgical gloves, latex, powdered	DD 1023663-1 NB 0197
safeCARE PF Surgical gloves powder free, sterile **Basic UDI-DI: 59079968T01010102-RYS**	Class IIa	safeCARE PF Surgical gloves powder free, sterile	DD 1023663-1 NB 0197
safeCARE basic Surgical gloves latex, powdered **Basic UDI-DI: 59079968T01010101-RYM**	Class IIa	safeCARE basic Surgical gloves latex, powdered	DD 1023663-1 NB 0197
safeCARE basic PF Surgical gloves latex, powder-free **Basic UDI-DI: 59079968T01010102-RYS**	Class IIa	safeCARE basic PF Surgical gloves latex, powder-free	DD 1023663-1 NB 0197
safeCARE premium Surgical gloves latex, powder-free **safeCARE UG Surgical gloves latex, powder-free** **safeCARE micro Surgical gloves latex, powder-free** **safeCARE ortho Surgical gloves latex, powder-free** **safeCARE dual Surgical gloves latex, powder-free** **Basic UDI-DI: 59079968T01010102-RYS**	Class IIa	safeCARE premium Surgical gloves latex, powder-free safeCARE UG Surgical gloves latex, powder-free safeCARE micro Surgical gloves latex, powder-free safeCARE ortho Surgical gloves latex, powder-free safeCARE dual Surgical gloves latex, powder-free	DD 1023663-1 NB 0197
safeCARE synthetic Surgical gloves neoprene, powder-free **safeCARE synthetic UG Surgical gloves neoprene, powder-free** **Basic UDI-DI: 59079968T010102-NRWL**	Class IIa	safeCARE synthetic Surgical gloves neoprene, powder-free safeCARE synthetic UG Surgical gloves neoprene, powder-free	DD 1023663-1 NB 0197
safeCARE fusion Surgical gloves polyisoprene, powder-free	Class IIa	safeCARE fusion Surgical gloves polyisoprene, powder-free	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968T010102-PRWS			
safeCARE virtuo Surgical gloves flexylon, powder-free **safeCARE virtuo UG Surgical gloves flexylon, powder-free** **safeCARE pro protect Surgical gloves flexylon, powder-free** **Basic UDI-DI: 59079968T010102-FRVU**	Class IIa	safeCARE virtuo Surgical gloves flexylon, powder-free safeCARE virtuo UG Surgical gloves flexylon, powder-free safeCARE pro protect Surgical gloves flexylon, powder-free	DD 1023663-1 NB 0197
safeLANCE Pressure-activated safety lancet **Basic UDI-DI: 59079968V0104RM**	Class IIa	safeLANCE Pressure-activated safety lancet	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-SETA**	Class IIa	deltaset Urinary bladder catheterization kit	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-SHTG**	Class IIa	deltaset Urinary bladder catheterization kit	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-CERS**	Class IIa	deltaset Dialysis kit	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-CHRY**	Class IIa	deltaset Dialysis kit	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-WETN**	Class IIa	deltaset Dressing change kit	DD 1023663-1 NB 0197
deltaset Procedure kit O **Basic UDI-DI: 59079968V0599-WHTU**	Class IIa	deltaset Dressing change kit	DD 1023663-1 NB 0197
deltaset Procedure kit O	Class IIa	deltaset Suture application kit	DD 1023663-1 NB 0197

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**
Basic UDI-DI: 59079968V0599-RET7		deltaset Suture removal kit	
deltaset Procedure kit O Basic UDI-DI: 59079968V0599-RHTD	Class IIa	deltaset Suture application kit deltaset Suture removal kit	DD 1023663-1 NB 0197
deltaset Procedure kit O Basic UDI-DI: 59079968V0599-IESC	Class IIa	deltaset Anasthesia kit deltaset Puncture kit	DD 1023663-1 NB 0197
deltaset Procedure kit O Basic UDI-DI: 59079968V0599-IHSJ	Class IIa	deltaset Anasthesia kit deltaset Puncture kit	DD 1023663-1 NB 0197
elastopor STERIL Non-woven dressing with absorbent pad, self-adhesive, sterile elastoKIDS STERIL Non-woven dressing with absorbent pad, self-adhesive, sterile Basic UDI-DI: 59079968T0305RB	Class I devices placed on the market in sterile condition	elastopor STERIL Non-woven dressing with absorbent pad, self-adhesive, sterile elastoKIDS STERIL Non-woven dressing with absorbent pad, self-adhesive, sterile	DD 1023663-1 NB 0197
elastoDERM PAD Foil dressing, with absorbent pad, self-adhesive, sterile Basic UDI-DI: 59079968M040101-FHU	Class I devices placed on the market in sterile condition	elastoDERM PAD Foil dressing, with absorbent pad, self-adhesive, sterile	DD 1023663-1 NB 0197
elastoSTRIP Wound closure strips, sterile Basic UDI-DI: 59079968M040499FL	Class I devices placed on the market in sterile condition	elastoSTRIP Wound closure strips, sterile	DD 1023663-1 NB 0197
UMBILICAL CORD CLAMP, sterile Basic UDI-DI: 59079968V0202RN	Class I devices placed on the market in sterile condition	UMBILICAL CORD CLAMP, sterile	DD 1023663-1 NB 0197
ALPHAtex Procedure gown NORMAL, sterile	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown NORMAL, sterile	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968T0205R6			
ALPHAtex Procedure gown NORMAL-P **Basic UDI-DI: 59079968T0205R6**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown NORMAL-P, sterile	HD 1023663-1 NB 0197
ALPHAtex Surgical gown STANDARD, sterile **ALPHAtex Surgical gown COMFORT sterile** **Basic UDI-DI: 59079968T020401HA**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown STANDARD, sterile ALPHAtex Surgical gown COMFORT sterile	DD 1023663-1 NB 0197
ALPHAtex Surgical gown CLASSIC-P **ALPHAtex Surgical gown STANDARD-P** **ALPHAtex Surgical gown COMFORT-P** **Basic UDI-DI: 59079968T020401HA**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown CLASSIC-P, sterile ALPHAtex Surgical gown STANDARD-P, sterile ALPHAtex Surgical gown COMFORT-P sterile	HD 1023663-1 NB 0197
ALPHAtex Surgical gown STANDARD PLUS, with impermeable parts, sterile **ALPHAtex Surgical gown COMFORT PLUS, with impermeable parts, sterile** **ALPHAtex Surgical gown EXTRA SAFE, with impermeable parts, sterile** **Basic UDI-DI: 59079968T020402HC**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown STANDARD PLUS, with impermeable parts, sterile ALPHAtex Surgical gown COMFORT PLUS, with impermeable parts, sterile ALPHAtex Surgical gown EXTRA SAFE, with impermeable parts, sterile	DD 1023663-1 NB 0197
ALPHAtex Surgical gown STANDARD PLUS-P, with impermeable parts **ALPHAtex Surgical gown COMFORT PLUS-P, with impermeable parts** **ALPHAtex Surgical gown EXTRA SAFE-P, with impermeable parts** **Basic UDI-DI: 59079968T020402HC**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical gown STANDARD PLUS-P, with impermeable parts, sterile ALPHAtex Surgical gown COMFORT PLUS-P, with impermeable parts, sterile ALPHAtex Surgical gown EXTRA SAFE-P, with	HD 1023663-1 NB 0197

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
		impermeable parts, sterile	
ALPHAtex Surgical drape, sterile **ALPHAtex 2-layer surgical drape, with cellulose layer, sterile** **ALPHAtex 2-layer surgical drape, sterile** **ALPHAtex 2-layer surgical drape with adhesive edge, sterile** **ALPHAtex 2-layer surgical drape with central fenestration, sterile** **ALPHAtex 2-layer surgical drape with central adhesive fenestration, sterile** **ALPHAtex 2-layer surgical drape with central, adjustable adhesive fenestration, sterile** **ALPHAtex 2-layer epidural surgical drape with adhesive fenestration, sterile** **ALPHAtex 3-layer surgical drape, sterile** **ALPHAtex 3-layer surgical drape with adhesive edge, sterile** **ALPHAtex 3-layer surgical drape with central fenestration, sterile** **ALPHAtex 3-layer surgical drape with central adhesive fenestration, sterile** **ALPHAtex Reinforced surgical drape with adhesive fenestration, sterile** **Basic UDI-DI: 59079968T0201QW**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical drape, sterile ALPHAtex 2-layer surgical drape, with cellulose layer, sterile ALPHAtex 2-layer surgical drape, sterile ALPHAtex 2-layer surgical drape with adhesive edge, sterile ALPHAtex 2-layer surgical drape with central fenestration, sterile ALPHAtex 2-layer surgical drape with central adhesive fenestration, sterile ALPHAtex 2-layer surgical drape with central, adjustable adhesive fenestration, sterile ALPHAtex 2-layer epidural surgical drape with adhesive fenestration, sterile ALPHAtex 3-layer surgical drape, sterile ALPHAtex 3-layer surgical drape with adhesive edge, sterile ALPHAtex 3-layer surgical drape with central fenestration, sterile ALPHAtex 3-layer surgical drape with central adhesive fenestration, sterile ALPHAtex Reinforced surgical drape with adhesive fenestration, sterile	DD 1023663-1 NB 0197

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
ALPHAtex Surgical drape **ALPHAtex 2-layer surgical drape, with cellulose layer** **ALPHAtex 2-layer surgical drape** **ALPHAtex 2-layer surgical drape with adhesive edge** **ALPHAtex 2-layer surgical drape with central fenestration** **ALPHAtex 2-layer surgical drape with central adhesive fenestration** **ALPHAtex 3-layer surgical drape** **ALPHAtex 3-layer surgical drape with adhesive edge** **ALPHAtex 3-layer surgical drape with central fenestration** **ALPHAtex 3-layer surgical drape with central adhesive fenestration** **Basic UDI-DI: 59079968T0201QW**	Class I devices placed on the market in sterile condition	ALPHAtex Surgical drape, sterile ALPHAtex 2-layer surgical drape, with cellulose layer, sterile ALPHAtex 2-layer surgical drape, sterile ALPHAtex 2-layer surgical drape with adhesive edge, sterile ALPHAtex 2-layer surgical drape with central fenestration, sterile ALPHAtex 2-layer epidural surgical drape with adhesive fenestration, sterile ALPHAtex 3-layer surgical drape, sterile ALPHAtex 3-layer surgical drape with adhesive edge, sterile ALPHAtex 3-layer surgical drape with central fenestration, sterile ALPHAtex 3-layer surgical drape with central adhesive fenestration, sterile	HD 1023663-1 NB 0197
ALPHAtex Instrument table cover, sterile **Basic UDI-DI: 59079968T030101-INJ**	Class I devices placed on the market in sterile condition	ALPHAtex Instrument table cover, sterile	DD 1023663-1 NB 0197
ALPHAtex Reinforced Mayo stand cover, sterile **ALPHAtex Reinforced Mayo stand cover, red, sterile** **Basic UDI-DI: 59079968T030101-MNS**	Class I devices placed on the market in sterile condition	ALPHAtex Reinforced Mayo stand cover, sterile ALPHAtex Reinforced Mayo stand cover, red, sterile	DD 1023663-1 NB 0197

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
ALPHAtex Armboard cover, sterile **ALPHAtex Surgical pocket for syringes, sterile** **Basic UDI-DI: 59079968T030101-NNU**	Class I devices placed on the market in sterile condition	ALPHAtex Armboard cover, sterile ALPHAtex Surgical pocket for syringes, sterile	DD 1023663-1 NB 0197
ALPHAtex Camera cables cover, sterile **ALPHAtex Circular banded cover for medical devices, sterile** **ALPHAtex Square banded cover for medical devices, sterile** **ALPHAtex C-arm cover set, sterile** **ALPHAtex Lamp handle cover, sterile** **ALPHAtex Ultrasound cover kits, sterile** **Basic UDI-DI: 59079968T030101-FNC**	Class I devices placed on the market in sterile condition	ALPHAtex Camera cables cover, sterile ALPHAtex Circular banded cover for medical devices, sterile ALPHAtex Square banded cover for medical devices, sterile ALPHAtex C-arm cover set, sterile ALPHAtex Lamp handle cover, sterile ALPHAtex Ultrasound cover kits, sterile	DD 1023663-1 NB 0197
ALPHAtex Absorbent drape, sterile **ALPHAtex Absorbent drape for newborn, sterile** **Basic UDI-DI: 59079968T020199-SRU**	Class I devices placed on the market in sterile condition	ALPHAtex Absorbent drape, sterile ALPHAtex Absorbent drape for newborn, sterile	DD 1023663-1 NB 0197
ALPHAtex Absorbent drape, sterile **ALPHAtex Absorbent drape for newborn, sterile** **Basic UDI-DI: 59079968T020199-SRU**	Class I devices placed on the market in sterile condition	ALPHAtex Absorbent drape, sterile ALPHAtex Absorbent drape for newborn, sterile	HD 1023663-1 NB 0197
ALPHAtex Limb cover, sterile **ALPHAtex Head Turban drape, sterile** **ALPHAtex Under buttocks drape, sterile** **Basic UDI-DI: 59079968T020102GV**	Class I devices placed on the market in sterile condition	ALPHAtex Limb cover, sterile ALPHAtex Head Turban drape, sterile ALPHAtex Under buttocks drape, sterile	DD 1023663-1 NB 0197
ALPHAtex Adhesive pouch, one-chamber, sterile	Class I devices placed on the	ALPHAtex Adhesive pouch,	DD 1023663-1 NB 0197

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
ALPHAtex Adhesive pouch, two-chamber, sterile **ALPHAtex Arthroscopy fluid collection pouch, cone-shaped, with outlet tip, sterile** **ALPHAtex Fluid collection pouch, cone-shaped, with outlet tip, sterile** **Basic UDI-DI: 59079968T020199-PRN**	market in sterile condition	one-chamber, sterile ALPHAtex Adhesive pouch, two-chamber, sterile ALPHAtex Arthroscopy fluid collection pouch, cone-shaped, with outlet tip, sterile ALPHAtex Fluid collection pouch, cone-shaped, with outlet tip, sterile	
ALPHAtex Non-woven surgical tape, adhesive, sterile **ALPHAtex Velcro surgical tape, sterile** **Basic UDI-DI: 59079968T020199-TRW**	Class I devices placed on the market in sterile condition	ALPHAtex Non-woven surgical tape, adhesive, sterile ALPHAtex Velcro surgical tape, sterile	DD 1023663-1 NB 0197
ALPHAtex Abdominal drape, sterile **ALPHAtex Abdo-Perineal drape, sterile** **ALPHAtex Angiography drape, sterile** **ALPHAtex Cardiology drape, sterile** **ALPHAtex Cardiac drape, sterile** **ALPHAtex C-section drape, sterile** **ALPHAtex Delivery drape, sterile** **ALPHAtex Extremity drape, sterile** **ALPHAtex Gynaecology drape, sterile** **ALPHAtex Laparoscopy drape, sterile** **ALPHAtex Ophthalmic drape, sterile** **ALPHAtex Orthopaedic drape, sterile** **ALPHAtex Shoulder drape, sterile** **ALPHAtex Vertical isolation drape, sterile**	Class I devices placed on the market in sterile condition	ALPHAtex Abdominal drape, sterile ALPHAtex Abdo-Perineal drape, sterile ALPHAtex Angiography drape, sterile ALPHAtex Cardiology drape, sterile ALPHAtex Cardiac drape, sterile ALPHAtex C-section drape, sterile ALPHAtex Delivery drape, sterile ALPHAtex Extremity drape, sterile ALPHAtex Gynaecology drape, sterile ALPHAtex Laparoscopy drape, sterile	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968T0202QY		ALPHAtex Ophthalmic drape, sterile ALPHAtex Orthopaedic drape, sterile ALPHAtex Shoulder drape, sterile ALPHAtex Vertical isolation drape, sterile	
ALPHAtex Abdominal drape, sterile **ALPHAtex Abdo-Perineal drape, sterile** **ALPHAtex Angiography drape, sterile** **ALPHAtex Cardiology drape, sterile** **ALPHAtex Cardiac drape, sterile** **ALPHAtex C-section drape, sterile** **ALPHAtex Delivery drape, sterile** **ALPHAtex Extremity drape, sterile** **ALPHAtex Gynaecology drape, sterile** **ALPHAtex Laparoscopy drape, sterile** **ALPHAtex Ophthalmic drape, sterile** **ALPHAtex Orthopaedic drape, sterile** **ALPHAtex Shoulder drape, sterile** **ALPHAtex Vertical isolation drape, sterile** **Basic UDI-DI: 59079968T0202QY**	Class I devices placed on the market in sterile condition	ALPHAtex Abdominal drape (from 1 to 100) ALPHAtex Abdo-Perineal drape (from 1 to 100) ALPHAtex Angiography drape (from 1 to 100) ALPHAtex Cardiology drape (from 1 to 100) ALPHAtex Cardiac drape (from 1 to 100) ALPHAtex C-section drape (from 1 to 100) ALPHAtex Delivery drape (from 1 to 100) ALPHAtex Extremity drape (from 1 to 100) ALPHAtex Gynaecology drape (from 1 to 100) ALPHAtex Laparoscopy drape (from 1 to 100) ALPHAtex Ophthalmic drape (from 1 to 100) ALPHAtex Orthopaedic drape (from 1 to 100) ALPHAtex Shoulder drape (from 1 to 100)	HD 1023663-1 NB 0197

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
		ALPHAtex Vertical isolation drape (from 1 to 100)	
ALPHAtex Abdominal set, sterile **ALPHAtex Abdo-Perineal set, sterile** **ALPHAtex Ablation set, sterile** **ALPHAtex Angiography set, sterile** **ALPHAtex Arthroscopy set, sterile** **ALPHAtex Basic set, sterile** **ALPHAtex Cardiology set, sterile** **ALPHAtex Cardiac set, sterile** **ALPHAtex Craniotomy set, sterile** **ALPHAtex C-section set, sterile** **ALPHAtex Cystoscopy set, sterile** **ALPHAtex Delivery set, sterile** **ALPHAtex Dental set, sterile** **ALPHAtex Dynamic hip screw set, sterile** **ALPHAtex Extremity set, sterile** **ALPHAtex Gynaecology set, sterile** **ALPHAtex Hip set, sterile** **ALPHAtex Laparoscopy set, sterile** **ALPHAtex Laryngology set, sterile** **ALPHAtex Ophthalmic set, sterile** **ALPHAtex Otolaryngology set, sterile** **ALPHAtex Pediatric set, sterile** **ALPHAtex Percutaneous lithotripsy set, sterile** **ALPHAtex Shoulder set, sterile** **ALPHAtex Spine set, sterile**	Class I devices placed on the market in sterile condition	ALPHAtex Abdominal set, sterile ALPHAtex Abdo-Perineal set, sterile ALPHAtex Ablation set, sterile ALPHAtex Angiography set, sterile ALPHAtex Arthroscopy set, sterile ALPHAtex Basic set, sterile ALPHAtex Cardiology set, sterile ALPHAtex Cardiac set, sterile ALPHAtex Craniotomy set, sterile ALPHAtex C-section set, sterile ALPHAtex Cystoscopy set, sterile ALPHAtex Delivery set, sterile ALPHAtex Dental set, sterile ALPHAtex Dynamic hip screw set, sterile ALPHAtex Extremity set, sterile ALPHAtex Gynaecology set, sterile ALPHAtex Hip set, sterile ALPHAtex Laparoscopy set, sterile ALPHAtex Laryngology set, sterile	DD 1023663-1 NB 0197

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
ALPHAtex Thyroid set, sterile **ALPHAtex TUR set, sterile** **ALPHAtex Universal set, sterile** **ALPHAtex Uro/gynaecology set, sterile** **ALPHAtex Varicose vein set, sterile** **ALPHAtex Vertical isolation set, sterile** **Basic UDI-DI: 59079968T0202QY**		ALPHAtex Ophthalmic set, sterile ALPHAtex Otolaryngology set, sterile ALPHAtex Pediatric set, sterile ALPHAtex Percutaneous lithotripsy set, sterile ALPHAtex Shoulder set, sterile ALPHAtex Spine set, sterile ALPHAtex Thyroid set, sterile ALPHAtex TUR set, sterile ALPHAtex Universal set, sterile ALPHAtex Uro/gynaecology set, sterile ALPHAtex Varicose vein set, sterile ALPHAtex Vertical isolation set, sterile	
ALPHAtex Abdominal set, sterile **ALPHAtex Abdo-Perineal set, sterile** **ALPHAtex Ablation set, sterile** **ALPHAtex Angiography set, sterile** **ALPHAtex Arthroscopy set, sterile** **ALPHAtex Basic set, sterile** **ALPHAtex Cardiology set, sterile** **ALPHAtex Cardiac set, sterile** **ALPHAtex Craniotomy set, sterile** **ALPHAtex C-section set, sterile**	Class I devices placed on the market in sterile condition	"ALPHAtex Abdominal set (from 1 to 200) ALPHAtex Abdo-Perineal set (from 1 to 200) ALPHAtex Ablation set (from 1 to 200) ALPHAtex Angiography set (from 1 to 200) ALPHAtex Arthroscopy set (from 1 to 200) ALPHAtex Basic set (from 1 to 200) ALPHAtex Cardiology set (from 1 to 200)	HD 1023663-1 NB 0197

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
ALPHAtex Cystoscopy set, sterile **ALPHAtex Delivery set, sterile** **ALPHAtex Dental set, sterile** **ALPHAtex Dynamic hip screw set, sterile** **ALPHAtex Extremity set, sterile** **ALPHAtex Gynaecology set, sterile** **ALPHAtex Hip set, sterile** **ALPHAtex Laparoscopy set, sterile** **ALPHAtex Laryngology set, sterile** **ALPHAtex Ophthalmic set, sterile** **ALPHAtex Otolaryngology set, sterile** **ALPHAtex Pediatric set, sterile** **ALPHAtex Percutaneous lithotripsy set, sterile** **ALPHAtex Shoulder set, sterile** **ALPHAtex Spine set, sterile** **ALPHAtex Thyroid set, sterile** **ALPHAtex TUR set, sterile** **ALPHAtex Universal set, sterile** **ALPHAtex Uro/gynaecology set, sterile** **ALPHAtex Varicose vein set, sterile** **ALPHAtex Vertical isolation set, sterile** **Basic UDI-DI: 59079968T0202QY**		ALPHAtex Cardiac set (from 1 to 200) ALPHAtex Craniotomy set (from 1 to 200) ALPHAtex C-section set (from 1 to 200) ALPHAtex Cystoscopy set (from 1 to 200) ALPHAtex Delivery set (from 1 to 200) ALPHAtex Dental set (from 1 to 200) ALPHAtex Dynamic hip screw set (from 1 to 200) ALPHAtex Extremity set (from 1 to 200) ALPHAtex Gynaecology set (from 1 to 200) ALPHAtex Hip set (from 1 to 200) ALPHAtex Laparoscopy set (from 1 to 200) ALPHAtex Laryngology set (from 1 to 200) ALPHAtex Ophthalmic set (from 1 to 200) ALPHAtex Otolaryngology set (from 1 to 200) ALPHAtex Pediatric set (from 1 to 200) ALPHAtex Percutaneous lithotripsy set (from 1 to 200) ALPHAtex Shoulder set (from 1 to 200) ALPHAtex Spine set (from 1 to 200) ALPHAtex Thyroid set (from 1 to 200)	

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
		ALPHAtex TUR set (from 1 to 200) ALPHAtex Universal set (from 1 to 200) ALPHAtex Uro/gynaecology set (from 1 to 200) ALPHAtex Varicose vein set (from 1 to 200) ALPHAtex Vertical isolation set (from 1 to 200)"	
elastoLUMENAL S Eye pad, superabsorbent, multi-layer, sterile **Basic UDI-DI: 59079968M040301-SKC**	Class I devices placed on the market in sterile condition	elastoLUMENAL S Eye pad, superabsorbent, multi-layer, sterile	DD 1023663-1 NB 0197
elastopor EYE Eye dressing, non-woven, with absorbent pad, self-adhesive, sterile elastoKIDS EYE Eye dressing, non-woven, with absorbent pad, self-adhesive, sterile **Basic UDI-DI: 59079968M0403NX**	Class I devices placed on the market in sterile condition	lastopor EYE Eye dressing, non-woven, with absorbent pad, self-adhesive, sterile elastoKIDS EYE Eye dressing, non-woven, with absorbent pad, self-adhesive, sterile	DD 1023663-1 NB 0197
COMBI STOPPER **Basic UDI-DI: 59079968C01018085**	Class I devices placed on the market in sterile condition	COMBI STOPPER	DD 1023663-1 NB 0197
LUER LOCK STOPPER **Basic UDI-DI: 59079968C01018085**	Class I devices placed on the market in sterile condition	LUER LOCK STOPPER	DD 1023663-1 NB 0197
elastopor STERIL D Non-woven dressing with absorbent pad, with incision and O-hole, self-adhesive, sterile elastoKIDS STERIL D Non-woven dressing, with absorbent pad, with	Class I devices placed on the market in sterile condition	elastopor STERIL D Non-woven dressing with absorbent pad, with incision and O-hole, self-adhesive, sterile elastopor STERIL D Non-woven	DD 1023663-1 NB 0197

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
incision and X-hole, self-adhesive, sterile **Basic UDI-DI: 59079968M04010201-DTG**		dressing, with absorbent pad, with incision and X-hole, self-adhesive, sterile elastoKIDS STERIL D Non-woven dressing, with absorbent pad, with incision and X-hole, self-adhesive, sterile	
NONVI lux S Non-woven swab, with O-incision, sterile **NONVI lux S Non-woven swab, with Y-incision, sterile** **Basic UDI-DI: 59079968M04010201-NU4**	Class I devices placed on the market in sterile condition	NONVI lux S Non-woven swab, with O-incision, sterile NONVI lux S Non-woven swab, with Y-incision, sterile	DD 1023663-1 NB 0197
elastopor IV IV cannula dressing, non-woven, selfadhesive, sterile **elastoKIDS IV IV cannula dressing, non-woven, selfadhesive, sterile** **Basic UDI-DI: 59079968M04010201H2**	Class I devices placed on the market in sterile condition	elastopor IV IV cannula dressing, non-woven, selfadhesive, sterile elastoKIDS IV IV cannula dressing, non-woven, selfadhesive, sterile	DD 1023663-1 NB 0197
elastoDERM F-IV IV cannula dressing, foil, with frame, with U-incision, self-adhesive, sterile **elastoDERM IV IV cannula dressing, foil, selfadhesive, sterile** **elastoDERM Foil dressing, self-adhesive, sterile** **elastoDERM F Foil dressing, with frame, selfadhesive, sterile** **elastoDERM C Foil dressing with a pocket to secure catheter, sterile** **Basic UDI-DI: 59079968M04010202H4**	Class I devices placed on the market in sterile condition	elastoDERM F-IV IV cannula dressing, foil, with frame, with U-incision, self-adhesive, sterile elastoDERM IV IV cannula dressing, foil, selfadhesive, sterile elastoDERM Foil dressing, self-adhesive, sterile elastoDERM F Foil dressing, with frame, selfadhesive, sterile elastoDERM C Foil dressing with a pocket to secure catheter, sterile	DD 1023663-1 NB 0197

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**
MULTIabsorb S ABD pad, non-woven and cellulose, sterile **Basic UDI-DI: 59079968M040201-SJZ**	Class I devices placed on the market in sterile condition	MULTIabsorb S ABD pad, non-woven and cellulose, sterile	DD 1023663-1 NB 0197
VAGINAL SPECULUM **Basic UDI-DI: 59079968U089006MJ**	Class I devices placed on the market in sterile condition	VAGINAL SPECULUM	DD 1023663-1 NB 0197
URINE BAG **Basic UDI-DI: 59079968A0603038J**	Class I devices placed on the market in sterile condition	URINE BAG	DD 1023663-1 NB 0197
URINE BAG with sample port, sterile **URINE BAG with sample port 2W, sterile** **Basic UDI-DI: 59079968A060303-PBW**	Class I devices placed on the market in sterile condition	URINE BAG with sample port, sterile URINE BAG with sample port 2W, sterile	DD 1023663-1 NB 0197
SAMPLES TAKING URINE BAG for boys, with sponge **SAMPLES TAKING URINE BAG for boys, without sponge** **SAMPLES TAKING URINE BAG for girls, with sponge** **SAMPLES TAKING URINE BAG for girls, without sponge** **Apteczka ABC Woreczek do pobierania próbek moczu dla dla chłopców z gąbką** **Apteczka ABC Woreczek do pobierania próbek moczu dla dziewczynek z gąbką** **Basic UDI-DI: 59079968A06030301AB**	Class I devices placed on the market in sterile condition	SAMPLES TAKING URINE BAG for boys, with sponge SAMPLES TAKING URINE BAG for boys, without sponge SAMPLES TAKING URINE BAG for girls, with sponge SAMPLES TAKING URINE BAG for girls, without sponge Apteczka ABC Woreczek do pobierania próbek moczu dla dla chłopców z gąbką Apteczka ABC Woreczek do pobierania próbek moczu dla dziewczynek z gąbką	DD 1023663-1 NB 0197
ENEMA BAG sterile **Basic UDI-DI: 59079968G020301-SDY**	Class I devices placed on the market in sterile condition	ENEMA BAG sterile	DD 1023663-1 NB 0197

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**
WOODEN TONGUE DEPRESSOR Sterile **Basic UDI-DI: 59079968V9001-SNM**	Class I devices placed on the market in sterile condition	WOODEN TONGUE DEPRESSOR sterile	DD 1023663-1 NB 0197
NELATON CATHETER NELATON CATHETER transparent **Basic UDI-DI: 59079968U010105H9**	Class I devices placed on the market in sterile condition	NELATON CATHETER NELATON CATHETER transparent	DD 1023663-1 NB 0197
GUEDEL AIRWAY **Basic UDI-DI: 59079968R010102FG**	Class I devices placed on the market in sterile condition	GUEDEL AIRWAY	DD 1023663-1 NB 0197
ENDOTRACHEAL TUBE HOLDER, vertical fixation ENDOTRACHEAL TUBE HOLDER, horizontal fixation **Basic UDI-DI: 59079968R010380-SNX**	Class I devices placed on the market in sterile condition	ENDOTRACHEAL TUBE HOLDER, vertical fixation ENDOTRACHEAL TUBE HOLDER, horizontal fixation	DD 1023663-1 NB 0197
INTUBATION STYLET TRACHEAL TUBE INTRODUCER CURVED **Basic UDI-DI: 59079968R010380-PNR**	Class I devices placed on the market in sterile condition	INTUBATION STYLET TRACHEAL TUBE INTRODUCER CURVED	DD 1023663-1 NB 0197
dicoSPIKE Withdrawal cannula with bacteria filter dicoSPIKE Withdrawal cannula with bacteria and particle filter dicoSPIKE Chemo Withdrawal cannula with bacteria and particle filter **Basic UDI-DI: 59079968A0704KC**	Class I devices placed on the market in sterile condition	dicoSPIKE Withdrawal cannula with bacteria filter dicoSPIKE Withdrawal cannula with bacteria and particle filter dicoSPIKE Chemo Withdrawal cannula with bacteria and particle filter	DD 1023663-1 NB 0197
elastoBAND BASIC S Knltted supporting bandage, sterile **Basic UDI-DI: 59079968M030301-SJT**	Class I devices placed on the market in sterile condition	elastoBAND BASIC S Knltted supporting bandage, sterile	HD 1023663-1 NB 0197
elastoBAND FLEX S Elastic bandage, sterile	Class I devices placed on the	elastoBAND FLEX S	DD 1023663-1 NB 0197

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
Basic UDI-DI: 59079968M030402-SKB	market in sterile condition	Elastic bandage, sterile	
elastoFILM Incise film, self-adhesive, sterile **elastoFILM M Incise film, self-adhesive, sterile** **Basic UDI-DI: 59079968T020101GT**	Class I devices placed on the market in sterile condition	elastoFILM Incise film, self-adhesive, sterile elastoFILM M Incise film, self-adhesive, sterile	DD 1023663-1 NB 0197
CERVICAL BRUSH standard **CERVICAL BRUSH special** **Basic UDI-DI: 59079968U090303L7**	Class I devices placed on the market in sterile condition	CERVICAL BRUSH standard CERVICAL BRUSH special	DD 1023663-1 NB 0197
omegapack Surgical set B **Basic UDI-DI: 59079968V0599-EP2**	Class IIb excluding Class IIb implantable non-WET	omegapack Surgical set B Orthopedic surgery set B Universal set B Surgical set B Orthopedic surgery set B Universal set B	HD 1023663-1 NB 0197
omegapack Surgical set B **Basic UDI-DI: 59079968V0599-KPE**	Class IIb excluding Class IIb implantable non-WET	omegapack Surgical set B Orthopedic surgery set B Universal set B Surgical set B Orthopedic surgery set B Universal set B	HD 1023663-1 NB 0197
omegapack Surgical set **Basic UDI-DI: 59079968V0599-ANS**	Class IIa	omegapack Surgical set Angiography set C-section set Laparoscopy set Gynecological surgery set Cardiac surgery set Neurosurgical set Orthopedic surgery set Otolaryngologic surgery set Urologic surgery set Delivery set Dressing set	HD 1023663-1 NB 0197

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
		Universal set	
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-WQ6**	Class IIa	deltaset Central venous access kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-IPA**	Class IIa	deltaset Universal kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-CNW**	Class IIa	deltaset Dialysis kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-NPL**	Class IIa	deltaset Universal kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-OPN**	Class IIa	deltaset Universal kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-FP4**	Class IIa	deltaset Urinary bladder catheterization kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-RPU**	Class IIa	deltaset Sewing kit Suture removal kit Dressing change kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-SPW**	Class IIa	deltaset Anasthesia kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-NIT3**	Class I devices placed on the market in sterile condition	deltaset Universal kit I	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-UITQ**	Class I devices placed on the market in sterile condition	deltaset Universal kit I	HD 1023663-1 NB 0197

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
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-BIRX**	Class I devices placed on the market in sterile condition	deltaset Dialysis kit	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-MISY**	Class I devices placed on the market in sterile condition	deltaset Universal kit I	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-DIS5**	Class I devices placed on the market in sterile condition	deltaset Operating field disinfection kit I	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-ZIU7**	Class I devices placed on the market in sterile condition	deltaset Suture removal kit I	HD 1023663-1 NB 0197
deltaset Procedure kit **Basic UDI-DI: 59079968V0599-PIT9**	Class I devices placed on the market in sterile condition	deltaset Protective kit I Hygiene kit I Neonatal kit I	HD 1023663-1 NB 0197

Table 2: Devices covered by this letter and for which the NB is <u>NOT</u> 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
none			

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2024/05/15	ZARYS_CL607_2024-05-15	Initial issue

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 93/42/EEC on Medical Devices (MDD) (Directive Certificates)

Manufacturer name	ZARYS International Group sp. z o.o. sp.k.
Manufacturer address and contact details	ul. Pod Borem 18, 41-808 Zabrze, Poland
Single Registration Number (SRN)	PL-MF-000000410

Notified body name and address	TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg, Germany
Notified body number	0197
Directive Certificate number(s) to which this confirmation is made	See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	See attached schedule
End date of extended validity/transition period	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

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.

✓ Expired/expires *after* 20 March 2023:

✓ 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.

- **Quality Management System (QMS)**
  - ✓ A QMS in accordance with Article 10(9) MDR is in place.

- **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: ZARYS International Group spółka z ograniczoną odpowiedzialnością spółka komandytowa

Location & Date: Zabrze, 1.03.2024

Name, Title: Joanna Skiba-Klyta, Quality and Regulatory Affairs Department Coordinator/ Management Board's Representative for Integrated Management Systems

Contact Details: qm@zarys.pl

PEŁNOMOCNIK ZARZĄDU DS. ZINTEGROWANEGO

Schedule of Devices

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

Identification of the device(s) (e.g., device name, family/group name device model or catalogue number)	**Device Classification**	**Directive Certificate number(s) to which this confirmation is made**	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity**	**End date of extended validity / transition period**
Sterile and non-sterile cutting gauze	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Non-sterile dressing gauze	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile and non-sterile gauze swabs (with or without X-ray thread)	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile and non-sterile gauze lap sponges (with X-ray thread/ with X-ray chip)	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile and non-sterile gauze balls (with or without X-ray thread)	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile and non-sterile gauze rolls (with or without X-ray thread)	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile and non-sterile non-woven swabs (with or without X-ray thread)	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile paraffin gauze dressings	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile three-way stopcocks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile transfusion sets for single use	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile infusion sets for single use	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile extension tubes for infusion pump	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile endotracheal tubes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile tracheostomy tubes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile breathing circuits	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile catheter mounts	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Non-sterile anaesthetic masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile laryngeal masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile oxygen masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile Venturi masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile non-rebreath masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile nebulizer masks	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile nasal oxygen cannulas	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile nebulizers	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile oxygen tubing	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile suction catheters	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile feeding tubes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile stomach and duodenal tubes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile urology catheters	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile surgical suction sets	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile surgical suction cannulas	Class IIa	DD 1023663-1	26 May 2024	31 December 2028

Sterile syringes for single use	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile insulin syringes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile tuberculin syringes	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile hypodermic needles	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile insulin pen needles	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile blood lancets	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile IV cannulas	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile needle free valves	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile surgical gloves	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Sterile procedure kits	Class IIa	DD 1023663-1	26 May 2024	31 December 2028
Elastic bandages	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Adhesive cannula fixation dressings	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Adhesive wound dressings	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Eye pads	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Incise films	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Transparent film dressings	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Absorbent wound dressings	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Surgical gowns	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Surgical drapes	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Sets of surgical drapes	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Fluid collection pouches	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Nelaton catheters	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Vaginal speculums	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Cervical brushes	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Urine bags	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Enema bags	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Tongue depressors	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Oropharyngeal airways	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Intubation stylets	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Endotracheal tube holders	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Suction tubes	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Withdrawal cannulas	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Cannula stoppers	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Umbilical cord clamps	Class I sterile	DD 1023663-1	26 May 2024	31 December 2028
Sterile surgical sets	Class IIa	HD 1023663-1	26 May 2024	31 December 2028
Sterile procedure kits	Class IIa	HD 1023663-1	26 May 2024	31 December 2028
Sterile surgical sets	Class IIb	HD 1023663-1	26 May 2024	31 December 2028
Surgical gowns	Class I sterile	HD 1023663-1	26 May 2024	31 December 2028
Surgical drapes	Class I sterile	HD 1023663-1	26 May 2024	31 December 2028
Sets of surgical drapes	Class I sterile	HD 1023663-1	26 May 2024	31 December 2028
Procedure kits	Class I sterile	HD 1023663-1	26 May 2024	31 December 2028
Knitted bandages	Class I sterile	HD 1023663-1	26 May 2024	31 December 2028