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9747 AN Groningen
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V.A.T.no: NL008483280B01
E-mail: ask@lode.nl

Manufacturer's Declaration

in relation to Regulation (EU) 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices, in particular with respect to

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

Manufacturer name	Lode B.V.
Manufacturer address and contact details	Zernikepark 16 9747 AN Groningen
Single Registration Number (SRN) (if available)	NL-MF-000001493

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

Notified body name (if applicable)	☒ See attached schedule
Notified body number (if applicable)	☒ See attached schedule
Directive Certificate number(s) to which this confirmation is made (if applicable)	☒ See attached schedule

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

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Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	☒ 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 *and/or*[2]
- the listed **device(s)** in the attached schedule and we as their manufacturer are in compliance with the conditions listed in Article 120.3c of the MDR for continued placing on the market and putting into service,

namely by fulfilling the following conditions:

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

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covering the listed device(s) was/were issued after 25 May 2017, was/were valid on 26 May 2021 and have not been withdrawn afterwards.

  Choose applicable statements:

  ☐ Expired *before* 20 March 2023:

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

  ☐ A Competent Authority has granted a derogation from the applicable conformity assessment procedure in accordance with Article 59(1) MDR (may be provided upon request), or

  ☐ A Competent Authority has required the manufacturer, in accordance with Article 97(1) MDR, to carry out the applicable conformity assessment procedure (may be provided upon request)

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

  ☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be

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

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in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

☒ Expired/expires *after* 20 March 2023:

Choose one applicable statement:

☒ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➢ **Upclassified devices**

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

Choose one applicable statement:

☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitutes and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➢ **Quality Management System (QMS)**

Choose one applicable statement:

☐ A QMS in accordance with Article 10(9) MDR will be put in place by no later than 26 May 2024.

☐ A QMS in accordance with Article 10(9) MDR is in place.

☒ A notified body has issued the attached certificate for the MDR-compliant QMS.

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

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

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Signed for and on behalf of the manufacturer:

Full Company Name	Lode B.V.
Location & Date	Groningen, 07-09-2026
Signature, Print Name, Title	K. Brouwer, Managing Director Lode B.V.
Contact Details	+31 (0) 50 571 2811 kbrouwer@lode.nl

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Schedule of Devices

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

ERGOMETERS		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Corival cpet	960900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	960900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival rehab	960901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival rehab with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	960901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival pediatric	960903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival pediatric with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	960903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival OEM	960912	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival OEM with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	960912	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival Recumbent cpet	969900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Product name	**Article number**					
Corival Recumbent cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	969900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival Recumbent rehab	969901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Corival Recumbent rehab with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	969901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Excalibur cpet	965900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Excalibur cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	965900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	925901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport PFM	925910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport PFM with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	925910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport 2	965911	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport 2 with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	965911	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	

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Product name	**Article number**					
Brachumera Sport 2 PFM	965921	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Brachumera Sport 2 PFM with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	965921	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet	967900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab	967901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet with static wall fixation	967902	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet with static wall fixation with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967902	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet with el. adj.wall fixation	967903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio cpet with el. adj.wall fixation with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Product name	**Article number**					
Angio cpet with automatic stand	967904	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio imaging with automatic stand with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967904	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio imaging with fix.set for instr.rail	967905	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio imaging with fix.set for instr.rail with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967905	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with static wall fixation	967922	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with static wall fixation with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967922	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with el. adj. wall fixation	967923	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with el. adj. wall fixation with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967923	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with automatic stand	967924	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with automatic stand with Blood pressure	967924	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Product name	**Article number**					
measurement and / or with SpO2 (blood oxygen saturation)						
Angio rehab with fix.set for instr.rail	967925	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with fix.set for instr.rail with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967925	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with bed carrier	967927	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio rehab with bed carrier with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967927	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

STRESS TABLES (in combination with Angio imaging)		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Angio Imaging with stress support radiology	967930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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STRESS TABLES (in combination with Angio imaging)		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Angio Imaging with stress support radiology	967940	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound	967931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound	967941	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967941	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology	967930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology	967940	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology with Blood pressure measurement and /	967940	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Product name	**Article number**					
or with SpO2 (blood oxygen saturation)						
Angio Imaging with stress support radiology ultrasound	967931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound	967941	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Angio Imaging with stress support radiology ultrasound with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	967941	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

MRI ergometers		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
MRI Pedal	937901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
MRI Push-pull	937902	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
MRI Up-down	937903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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MRI ergometers		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
MRI Dorsal ankle flex	937904	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

Software		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Lode Ergometry Manager (LEM 10)						
Main Software	955900	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A
Module Wingate plus	955915	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A
Module Export	955920	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A
Module Astrand	955916	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A
Module Architect	955921	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A
Lode Cardiac Rehabilitation Manager (LCRM)	950902	2110393CE02	26 May 2024	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Treadmills		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Flow cpet	938910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Flow cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Flow rehab	938911	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Flow rehab with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938911	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
(Valiant /) OEM2	939910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
(Valiant /) OEM2 with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	939910	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 family						
Valiant 2 cpet	938900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 cpet with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 rehab	938901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 rehab with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Treadmills		**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
Product name	**Article number**					
Valiant 2 sport	938902	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 sport with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938902	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 rehab XL	938903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 rehab XL with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938903	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 pediatric	938904	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 pediatric with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938904	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 cpet XL	938905	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 cpet XL with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938905	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 sport XL	938906	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Valiant 2 sport XL with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	938906	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A

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Product name	**Article number**					
Katana Family						
Katana Sport 400 V	934900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport 400 V with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	934900	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport 200 V	934901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport 200 V with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	934901	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport XL 400 V	934930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport XL 400 V with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	934930	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport XL 200 V	934931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A
Katana Sport XL 200 V with Blood pressure measurement and / or with SpO2 (blood oxygen saturation)	934931	2110393CE01	1 December 2023	DEKRA Certification B.V. 0344	31-12-2028	N/A