

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*
- 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	Wöhlk Contactlinsen GmbH
Manufacturer address and contact details	Wöhlk Contactlinsen GmbH Bürgermeister-Schade-Str. 12 - 16 24232 Schönkirchen Germany
Single Registration Number (SRN) (if available)	DE-MF-000022424

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

Notified body name (if applicable)	<input checked="" type="checkbox"/> See attached schedule
Notified body number (if applicable)	<input checked="" type="checkbox"/> See attached schedule
Directive Certificate number to which this confirmation is made (if applicable)	<input checked="" type="checkbox"/> See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	<input checked="" type="checkbox"/> See attached schedule
End date of extended validity/transition period	<input checked="" type="checkbox"/> See attached schedule



We, as the manufacturer declare under our sole responsibility:

- for the above listed **Directive Certificate** the conditions for the legal extension of validity as required in Article 120.2 of the MDR are met *and/or*
- 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Certificate covering the listed device was issued after 25 May 2017, was valid on 26 May 2021 and have not been withdrawn afterwards.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 been made by us to a notified body no later than 26 May 2024 for the devices listed in the attached schedule and a signed written agreement is 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)**

- 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.

➤ **Devices as listed in the attached schedule**

- The devices continue to comply with the AIMDD or MDD.
- There are no significant changes in the design and intended purpose.
- The devices 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:

Wöhlk Contactlinsen GmbH

Schönkirchen, 2024-05-24

Dirk Reschat, Head of Quality Management, Regulatory Affairs and PRRC

dirk.reschat@woehlk.com

Schedule of Devices

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

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
CONTACT life spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ZEISS CD 30 Compatic spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Eigenmarke Vitafilcon A spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Eigenmarke bio 30 spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Eigenmarke Premium spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk Eigenmarke Advance spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ContaView Premium spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT life toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ZEISS CD 30 Compatic toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Eigenmarke Vitafilcon A toric 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Eigenmarke bio 30 toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk Eigenmarke Premium toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Eigenmarke Advance toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ContaView premium toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ZEISS Contact Day 1 spheric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ZEISS Contact Day 1 toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
ZEISS Contact Day 1 multifocal	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk Weflex 55	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Weflex 55 toric	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Weflex 55 toric Advance	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk Geaflex 70	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual SH	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual SH TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
CONTACT individual SH TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual SH TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi SH	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi SH TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi SH TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi SH TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_soft SH	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft SH TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft SH TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft SH TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi SH	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi SH TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_soft multi SH TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi SH TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual BIO	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual BIO TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual BIO TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual BIO TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
CONTACT individual multi BIO	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi BIO TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi BIO TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
CONTACT individual multi BIO TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft BIO	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft BIO TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_soft BIO TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft BIO TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi BIO	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi BIO TP	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi BIO TD	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_soft multi BIO TDS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse Wöhlk S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI FEN	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI FEN RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI FEN BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI FEN VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi FEN	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi FEN RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_multi FEN BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi FEN VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk AS 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_as 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_mas BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_mas BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk KE	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk KE RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk KE BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk KE BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk KE VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_ke	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_ke RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_ke BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_ke BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_ke VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk BIFO S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk BIFO S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk BIFO S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk BIFO S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse Wöhlk BIFO S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse Wöhlk bifo S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk bifo S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk bifo S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk bifo S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse Wöhlk bifo S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_bifo s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk BIFO AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_bifo as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk BIFO MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk BIFO AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk BIFO AS 2VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_bifo as2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_bifo as2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk MULTI S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_multi s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk MULTI S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk MULTI AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_multi as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinsewöhlk MULTI MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi mas BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse EGO_multi mas BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse wöhlk MULTI AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Diagnoselinse wöhlk MULTI AS 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Diagnoselinse EGO_multi as 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI FEN	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI FEN RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI FEN BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI FEN VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi FEN	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi FEN RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_multi FEN BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi FEN VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk AS 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_as 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_mas BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_mas BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk KE	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk KE RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk KE BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk KE BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk KE VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_ke	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_ke RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_ke BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_ke BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_ke VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
Wöhlk bifo S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
Wöhlk bifo S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk BIFO AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk BIFO AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO AS 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk BIFO MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk BIFO MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_bifo s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_bifo s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_bifo as 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo as 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_bifo mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo mas BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo mas BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_bifo mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 2K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTli S 2K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk MULTI S 2K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 2K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTli S 2K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 3K	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 3K RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 3K BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk MULTI S 3K BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI S 3K VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

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wöhlk MULTI AS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI MAS	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI MAS RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI MAS BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI MAS BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI MAS VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

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wöhlk MULTI AS 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk MULTI AS 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 2k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

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EGO_multi s 2k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 2k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 2k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 2k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 3k	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 3k RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_multi s 3k BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 3k BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi s 3k VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_multi as BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi mas	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi mas RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi mas BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi mas BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
EGO_multi mas VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as 2	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as 2 RT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as 2 BT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as 2 BTK	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_multi as 2 VPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A

Identification of the devices (e.g., device name, family/group name device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made	Original expiry date as indicated on the Directive Certificate prior to the extension of the validity	Notified Body name and number that issued the Directive Certificate	Notified Body name and number where the MDR application was lodged/contract signed	End date of extended validity / transition period	Substitute Device(s)
wöhlk OKE	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_oke	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
wöhlk OKE RPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A
EGO_oke RPT	HD 1566826-1	2024-05-26	TÜV Rheinland LGA Products GmbH CE 0197	TÜV Rheinland LGA Products GmbH CE 0197	2028-12-31	N/A