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	eyelike GmbH
Manufacturer address and contact details	Ringstraße 12 63543 Neuberg, Hessen, DE +49 (0) 6185 / 570 j.giehl@eyelike-kontaktlinsen.de
Single Registration Number (SRN) (if available)	DE-MF-000008175

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

Notified body name (if applicable)	DNV MedCert □ See attached schedule
Notified body number (if applicable)	0482 □ See attached schedule
Directive Certificate number(s) to which this confirmation is made (if applicable)	3924DE410200505 □ See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	2024-05-27 □ See attached schedule
End date of extended validity/transition period	2028-12-31 □ 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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Handelsregister Hanau HRB 91414 Ust-IdNr.DE241933083; St.Nr.:035 232 50757; Geschäftsführerin: Maria Giehl
Bank: VR Bank Main-Kinzig-Büdingen eG IBAN:DE52506616390002221500; BIC:GENODEF1LSR

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 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.

[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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☒ 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.

Signed for and on behalf of the manufacturer:

Company:

Eyelike GmbH
Ringstraße 12
63543 Neuberg
Tel.: +49 (0) 6185-570
info@eyelike-kontaktlinsen.de

21.05.2024,

Datum Johannes Giehl, CEO

eyelike GmbH Ringstraße 12 63543 Neuberg Telefon: (+49) 6185-570 Fax: (+49) 6185-894828 E-Mail: info@eyelike-kontaktlinsen.de
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Schedule of Devices

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

Identification of the device(s) (e.g., device name, family/group name device model or catalogue number)		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 that issued the Directive Certificate (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)
Device Name:	Catalogue (EYE) Number / EAN	eyelike CE certificate	YYYY-MM-DD	NB - Number	NB - Number	YYYY-MM-DD	non applicable
Visiomax Monatslinse, Dioptrie: -0,75	EYE 2173 / 401035559 1074	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -1	EYE 1100 / 401035558 2119	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -1,25	EYE 1125 / 401035558 2133	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -1,5	EYE 1150 / 401035558 2157	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -1,75	EYE 1175 / 401035558 2171	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -2	EYE 1200 / 401035558 2195	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -2,25	EYE 1225 / 401035558 2218	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -2,5	EYE 1250 / 401035558 2232	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -2,75	EYE 1275 / 401035558 2256	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -3	EYE 1300 / 401035558 2270	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -3,25	EYE 1325 / 401035558 2294	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -3,5	EYE 1350 / 401035558 2317	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -3,75	EYE 1375 / 401035558 2331	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -4	EYE 1400 / 401035558 2355	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -4,25	EYE 1425 / 401035558 2379	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -4,5	EYE 1450 / 401035558 2393	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -4,75	EYE 1475 / 401035558 2416	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -5	EYE 1500 / 401035558 2430	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -5,25	EYE 1525 / 401035558 2454	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -5,5	EYE 1550 / 401035558 2478	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -5,75	EYE 1575 / 401035558 2492	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A

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Visiomax Monatslinse, Dioptrie: -6	EYE 1600 / 401035558 2515	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Monatslinse, Dioptrie: -6,5	EYE 1650 / 401035558 2539	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -1	EYE 2126 / 401035558 2553	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -1,25	EYE 2127 / 401035558 2577	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -1,5	EYE 2128 / 401035558 2591	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -1,75	EYE 2129 / 401035558 2614	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -2	EYE 2130 / 401035558 2638	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -2,25	EYE 2131 / 401035558 2652	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -2,5	EYE 2132 / 401035558 2676	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -2,75	EYE 2133 / 401035558 2690	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -3	EYE 2134 / 401035558 2713	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -3,25	EYE 2135 / 401035558 2737	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -3,5	EYE 2136 / 401035558 2751	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -3,75	EYE 2137 / 401035558 2775	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -4	EYE 2138 / 401035558 2799	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -4,25	EYE 2139 / 401035558 2812	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -4,5	EYE 2140 / 401035558 2836	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
Visiomax Silikon-Hydrogel Monatskontaktlinse, Dioptrie: -4,75	EYE 2141 / 401035558 2850	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -0,75	EYE 2191 / 405817252 8934	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -1,00	EYE 2192 / 405817252 8958	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -1,25	EYE 2193 / 405817252 8972	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -1,50	EYE 2194 / 405817252 8996	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -1,75	EYE 2195 / 405817252 9016	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -2,00	EYE 2196 / 405817252 9030	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -2,25	EYE 2197 / 405817252 9054	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -2,50	EYE 2198 / 405817252 9078	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -2,75	EYE 2199 / 405817252 9092	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -3,00	EYE 2201 / 405817252 9115	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -3,25	EYE 2202 / 405817252 9139	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -3,50	EYE 2203 / 405817252 9153	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -3,75	EYE 2204 / 405817252 9177	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -4,00	EYE 2205 / 405817252 9191	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A

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VISIOMAX Tageskontaktlinsen, Dioptrie -4,25	EYE 2206 / 405817252 9214	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -4,50	EYE 2207 / 405817252 9238	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -4,75	EYE 2208 / 405817252 9252	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -5,00	EYE 2209 / 405817252 9276	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -5,25	EYE 2230 / 405817252 9290	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -5,50	EYE 2231 / 405817252 9306	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -5,75	EYE 2232 / 405817252 9313	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -6,00	EYE 2233 / 405817252 9320	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Tageskontaktlinsen, Dioptrie -6,50	EYE 2234 / 405817252 9337	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kombi Lösung, 360 ml	EYE 2178 / 405817233 1701	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISOMAX Kombi Lösung Ampullen, 15 x 10 ml	EYE 2171 / 405817203 0079	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kombilösung Hyaluron, 360 ml	EYE 2170 / 405817202 9974	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISOMAX Kombilösung Sensitive 360 ml	EYE 2142 / 405817233 1800	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISOMAX Kombilösung SH mit Augentrost und Kamille, 360 ml	EYE 2143 / 405817203 0031	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kombilösung Super, 60 ml	EYE 2172 / 405817203 0055	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kombilösung Super, 360 ml	EYE 2168 / 405817202 9950	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Peroxidlösung Komfort Super, 360 ml	EYE 5329 / 401035579 5243	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kochsalz Lösung, 360 ml	EYE 2181 / 405817233 1725	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Kochsalz Lösung, 100 ml	EYE 2190 / 405817252 8910	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Hartlinsenreiniger, 30 ml	EYE 2186 / 405817233 1909	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Hartlinsen Aufbewahrung, 240 ml	EYE 2187 / 405817233 1923	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Augentropfen Hyaluron, 0,1 %, 15 ml	EYE 2183 / 405817233 1787	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Augentropfen Hyaluron 0,2 %, 15 x 0,4 ml	EYE 2184 / 405817233 1824	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Augentropfen Hyaluron 0,3 %, 10ml	EYE 2185 / 405817233 1848	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Augentropfen mit Augentrost und Kamille, 15 x 0,4 ml	EYE 2144 / 405817208 0500	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A
VISIOMAX Augentropfen Aloe vera, 10ml	EYE 2242 / 406644725 1012	3924DE410200505	2024-05-27	DNV MedCert - 0482	DNV MedCert - 0482	2028-12-31	N/A

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