



wöhlk
CONTACTLINSEN

EG – KONFORMITÄTSERKLÄRUNG

EC DECLARATION OF CONFORMITY · DÉCLARATION CE DE CONFORMITÉ ·
DICHIARAZIONE CE DI CONFORMITÀ · ЕС – СЕРТИФИКАТ СООТВЕТСТВИЯ

Name und Adresse des Herstellers: /
Name and address of the manufacturer: /
Nom et adresse du fabricant: /
Nome e indirizzo del fabbricante: /
Название и адрес производителя:

Wöhlk Contactlinsen GmbH
Bürgermeister-Schade-Str. 12 - 16
24232 Schönkirchen
Germany

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che /
мы заявляем о полной ответственности, что

das hergestellte Medizinprodukt: /
the manufactured medical device: /
le produit de dispositif médical: /
el producto de dispositivo medico: /
то произведённый медицинский продукт:

wöhlk AS

wöhlk AS RT

wöhlk AS BTK

wöhlk AS BT

wöhlk AS VPT

EGO_as

EGO_as RT

EGO_as BTK

EGO_as BT

EGO_as VPT

wöhlk BIFO AS

wöhlk BIFO AS RT

wöhlk BIFO AS BTK

wöhlk BIFO AS BT

wöhlk BIFO AS VPT

EGO_bifo as

EGO_bifo as RT

EGO_bifo as BTK

EGO_bifo as BT

EGO_bifo as VPT

wöhlk MULTI AS

wöhlk MULTI AS RT

wöhlk MULTI AS BTK

wöhlk MULTI AS BT

wöhlk MULTI AS VPT

EGO_multi as

EGO_multi as RT

EGO_multi as BTK

EGO_multi as BT



wöhlk
CONTACTLINSEN

EGO_multi as VPT

Diagnoselinse wöhlk AS

Diagnoselinse wöhlk AS RT

Diagnoselinse wöhlk AS BTK

Diagnoselinse wöhlk AS BT

Diagnoselinse wöhlk AS VPT

Diagnoselinse EGO_as

Diagnoselinse EGO_as RT

Diagnoselinse EGO_as BTK

Diagnoselinse EGO_as BT

Diagnoselinse EGO_as VPT

Diagnoselinse wöhlk BIFO AS

Diagnoselinse wöhlk BIFO AS RT

Diagnoselinse wöhlk BIFO AS BTK

Diagnoselinse wöhlk BIFO AS BT

Diagnoselinse wöhlk BIFO AS VPT

Diagnoselinse EGO_bifo as

Diagnoselinse EGO_bifo as RT

Diagnoselinse EGO_bifo as BTK

Diagnoselinse EGO_bifo as BT

Diagnoselinse EGO_bifo as VPT

Diagnoselinse wöhlk MULTI AS

Diagnoselinse wöhlk MULTI AS RT

Diagnoselinse wöhlk MULTI AS BTK

Diagnoselinse wöhlk MULTI AS BT

Diagnoselinse wöhlk MULTI AS VPT

Diagnoselinse EGO_multi as

Diagnoselinse EGO_multi as RT

Diagnoselinse EGO_multi as BTK

Diagnoselinse EGO_multi as BT

Diagnoselinse EGO_multi as VPT

der Klasse: / of class: /
de la classe: / di classe: / класса:

II a

nach Anhang IX der Richtlinie 93/42/EWG / according to annex IX of directive 93/42/EEC / selon l'annexe IX de la directive 93/42/CEE / secondo l'allegato IX della direttiva 93/42/CEE / по дополнению IX Закона 93/42/EEC

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 93/42/EWG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“.

/meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

/remplit toutes les exigences de la directive sur les dispositifs médicaux 93/42/CEE et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l'inspection finale» du produit.

/soddisfa tutte le disposizioni della direttiva 93/42/CEE e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il "rapporto di ispezione finale" del prodotto. /

соответствует требованиям законодательства медицинской продукции 93/42/EWG и их применении. заявление действительно в соответствии с принадлежащим к продукту Контрольному протоколу.

Konformitätsbewertungsverfahren: /
Conformity assessment procedure: /
Procédure d'évaluation de la conformité: /
Procedura di valutazione della conformità:
процедура оценки соответствия:

**Richtlinie 93/42/EWG Anhang II, ohne Abschnitt 4
Directive 93/42/EEC Annex II, excluding Section 4
Directive 93/42/CEE Annexe II, hors section 4
Direttiva 93/42/CEE senza Allegato II, sezione 4
руководство 93/42/EWG Приложение 2, без
подразделения 4**

Zertifikat-Registrier-Nr.:/ Certificate-Registration
no:/ Certificat n°d'enregistrement:/ Certificado
Numero di registrazione:/ Сертификат
Регистрационный номер:

HD 1566826-1

Gültig bis:/ Valid until:/ Valable jusqu'au:/
Valido fino al:/ действует до:

2024/05/26

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:/
Регистрационный орган:

**TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Germany
CE 0197**

Schönkirchen, 2022/01/04
Ort, Datum / Place, date / Lieu, date /
Luogo, data / Место, дата



Dr. Dirk Lauscher / Managing Director