KONFORMITÄTSERKLÄRUNG
DECLARATION OF CONFORMITY

gemäß Anhang IX der Verordnung (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Hiermit erklären wir in alleiniger Verantwortung, dass die unten aufgeführten Produkte die relevanten allgemeinen Sicherheits- und Leistungsanforderungen gemäß der **Verordnung (EU) 2017/745** über Medizinprodukte erfüllen und mit folgenden Anforderungen konform sind

- der Liste der angewandten Normen
- Gemeinsame Spezifikationen: n/a
- Weitere Regularien: n/a

Diese Konformitätserklärung ist bis zum Ablauf des Zertifikates der Verordnung (EU) 2017/745 über Medizinprodukte (Registrierungsnummer: HZ 1123799-1) und nur zusammen mit den zugehörigen Chargenfreigabedokumenten gültig.

We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the ***Regulation (EU) 2017/745 on Medical Devices*** *and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Hersteller *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Deutschland */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
Einmalige Registrierungsnummer *Single Registration Number (SRN):*	DE-MF-000005414
Produktfamilie *Name of product family:*	Lot für edelmetallfreie Legierungen *Base-metal Brazing Materials*
Produktname und REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Zweckbestimmung *Intended purpose:*	Füllstoff, bestimmt zum Löten von dentalen Restaurationen wie Modellguss und Kombinationsarbeiten. *Filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Produktklassifizierung *Product class:*	Klasse IIa *Class IIa*
Basis UDI-DI *Basic UDI-DI:*	++EBGOMetal-braz-matUU
Benannte Stelle *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg Deutschland

	TÜV Rheinland LGA Products GmbH *Tillystrasse 2* *90431 Nuremberg* *Germany*
CE-Kennzeichen *Marked*	CE 0197

Andere anwendbaren EU-Sprachen sind in den folgenden Anhängen zu finden.
Other applicable EU-languages can be found in the following Annex.

Bremen, 12.05.2026

Ort, Datum
Place, Date

Steffen Böhm
Managing Director (CPO)

Dr. Ulrich Abend
Person Responsible for Regulatory Compliance – Technical Documentation

es / en

DECLARACIÓN DE CONFORMIDAD
DECLARATION OF CONFORMITY

según el anexo IX, Reglamento (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Por la presente declaramos, bajo nuestra única responsabilidad, que los productos que se detallan a continuación cumplen con los requisitos generales de seguridad y rendimiento relevantes según lo establecido del **Reglamento (UE) 2017/745 sobre los productos sanitarios**, y guardan conformidad con

- la lista de normas aplicadas
- especificaciones comunes: *n/a*
- reglamentos: n/a

Esta declaración de conformidad tiene validez hasta la fecha de vencimiento del certificado REGLAMENTO (UE) 2017/745 sobre los productos sanitarios (número de registro: HZ 1123799-1); y únicamente en combinación con el documento correspondiente de aprobación del lote.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Fabricante *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Alemania */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
número de registro único *Single Registration Number (SRN)*	DE-MF-000005414
Familia de productos *Name of product family*	Lot per leghe di metalli preziosi *Base-metal Brazing Materials*
nombre del producto y REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Finalidad prevista *Intended purpose*	Un material de relleno y está previsto para la soldadura indirecta de restauraciones dentales como colado sobre modelo y trabajos combinados. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Clasificación del producto *Product class*	Clase IIa *Class IIa*
UDI-DI básico *Basic UDI-DI*	++EBGOMetal-braz-matUU
organismo notificado *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Núremberg

	Alemania *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
marcado CE *Marked*	CE 0197

fr / en

DÉCLARACION DE CONFOMITÉ
DECLARATION OF CONFORMITY

conformément à l'annexe IX, réglement (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Nous déclarons par la présente, sous notre seule responsabilité, que les dispositifs listés ci-après satisfont aux exigences générales pertinentes en matière de sécurité et de performances conformément à du **Règlement (UE) 2017/745 relatif aux dispositifs médicaux** et sont conformes aux

- normes appliquées listées,
- spécifications communes : n/a
- règlements : n/a

Cette déclaration de conformité est valable jusqu'à expiration du Certificat RÈGLEMENT (EU) 2017/745 relatif aux dispositifs médicaux (n° d'enregistrement: HZ 1123799-1) et uniquement avec le document correspondant de libération du lot.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Fabricant *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Allemagne */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
numéro d'enregistrement unique *Single Registration Number (SRN)*	DE-MF-000005414
Famille de Produits *Name of product family*	Matériau d'apport pour alliages non précieux *Base-metal Brazing Materials*
nom du produit et REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Destination *Intended purpose*	Une charge destinée au brasage de restaurations dentaires comme les châssis et les travaux combinés. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Classe de produit *Product class*	Classe IIa *Class IIa*
IUD-ID de base *Basic UDI-DI*	++EBGOMetal-braz-matUU
Organisme notifié	TÜV Rheinland LGA Products GmbH

Notified Body	Tillystrasse 2 90431 Nürnberg Allemagne *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
CE-marque *Marked*	CE 0197

DICHIARAZIONE DI CONFORMITÀ
DECLARATION OF CONFORMITY

secondo dall'allegato IX, Regolamento (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Con la presente dichiariamo, sotto la nostra esclusiva responsabilità, che i dispositivi elencati di seguito rispettano i pertinenti requisiti generali di sicurezza e prestazione come previsto del **Regolamento (UE) 2017/745 relativo ai Dispositivi Medici** e sono conformi

- all'elenco delle norme applicate
- alle specifiche comuni: n/a
- ai regolamenti: n/a

La presente dichiarazione di conformità è valida fino alla scadenza del Certificato REGOLAMENTO (UE) 2017/745 relativo ai Dispositivi Medici (Numero di Registrazione: HZ 1123799-1) e soltanto in combinazione con il relativo certificato di rilascio del lotto.

We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the ***Regulation (EU) 2017/745 on Medical Devices*** *and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Fabbricante *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Germania */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
numero di registrazione unico *Single Registration Number (SRN)*	DE-MF-000005414
Famiglia di prodotti *Name of product family*	Lot per leghe di metalli preziosi *Base-metal Brazing Materials*
nome del prodotto e REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Destinazione d'uso *Intended purpose*	Un riempitivo indicato per la brasatura di restauri dentali come scheletrati e lavori combinati. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
classificazione del prodotto *Product class*	Classe IIa *Class IIa*
UDI-DI di base *Basic UDI-DI*	++EBGOMetal-braz-matUU
organismo notificato *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2

	90431 Norimberga Germania *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
marchio CE *Marked*	CE 0197

ATBILSTĪBAS DEKLARĀCIJAS
DECLARATION OF CONFORMITY

Saskaņā ar IX pielikumā, regula (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Ar šo mēs, uzņemoties pilnu atbildību, deklarējam, ka iepriekš minētie izstrādājumi atbilst attiecīgajām vispārējām drošuma un veiktspējas prasībām saskaņā ar **Regulas (ES) 2017/745 par medicīnas ierīcēm** atbilst

- uzskaitītajiem attiecināmajiem standartiem
- kopīgās specifikācijas: n/a
- regulas: n/a

Šī atbilstības deklarācija ir derīga līdz sertifikāta REGULA (ES) 2017/745 par medicīnas ierīcēm (reģistrācijas Nr.: HZ 1123799-1) derīguma beigām un tikai kopā ar saistītās partijas laidiena dokumentu.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

ražotāji *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Vācija */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
vienotu reģistrācijas numuru (VRN) *Single Registration Number (SRN)*	DE-MF-000005414
Produktu grupa *Name of product family*	Lodējamais materiāls cēlmetālus nesaturošiem sakausē Jumiem *Base-metal Brazing Materials*
Produkta nosaukums un REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Paredzētais nolūks *Intended purpose*	Ir pildviela, kas paredzēta tādu dentālo restaurāciju lodēša nai kā modeļlējums un kombinētie darbi. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Produktu klasifikācija *Product class*	Klase IIa *Class IIa*
pamata UDI-DI *Basic UDI-DI*	++EBGOMetal-braz-matUU
paziņotajai struktūrai *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2

	90431 Nirnberga Vācija *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
CE- marķējums *Marked*	CE 0197

ATITIKTIES DEKLARACIJO
DECLARATION OF CONFORMITY

Pagal IX priedą IX, reglamentai (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Savo atsakomybe pareiškiame, kad toliau išvardyti gaminiai tenkina atitinkamus bendrosios saugos ir veikimo reikalavimus, numatytus **Medicinos prietaisų reglamento (ES) 2017/745** ir atitinka

- išvardytus taikomus standartus
- bendrosios specifikacijos: n/a
- reglamentai: n/a

Ši atitikties deklaracija galioja iki Medicinos prietaisų reglamente (ES) 2017/745 numatyto sertifikato (registracijos Nr.: HZ 1123799-1) galiojimo pabaigos ir tik kartu su atitinkamos partijos išleidimo dokumentu.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Gamintojų *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Vokietija */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
unikalųjį registracijos numerį *Single Registration Number (SRN)*	DE-MF-000005414
produktų grupė *Name of product family*	Lydmetalis lydiniams, kuriuose nėra tauriųjų metalų *Base-metal Brazing Materials*
produkto pavadinimas ir REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Numatytoji paskirtis *Intended purpose*	Yra užpildas, skirtas lituoti dantų restauracijas, tokias kaip modelių liejinius, ir kombinuotiems darbams. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Produkto klasifikacija *Product class*	Klasė IIa *Class IIa*
Bazinis UDI-DI *Basic UDI-DI*	++EBGOMetal-braz-matUU
įsisteigusi notifikuotoji *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Niurnbergas Vokietija

	TÜV Rheinland LGA Products GmbH *Tillystrasse 2* *90431 Nuremberg* *Germany*
CE ženklu *Marked*	CE 0197

MEGFELELŐSÉGI NYILATKOZATOT
DECLARATION OF CONFORMITY

a IX. melléklet szerint, Rendeletek (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Ezúton kizárólagos felelősségünk tudatában kijelentjük, hogy az alábbiakban felsorolt termékek betartják **az orvostechnikai eszközökről szóló (EU) 2017/745** mellékletében meghatározott biztonságosságra és teljesítőképességre vonatkozó általános követelményeknek, valamint

- az alkalmazott szabványok listájának
- általános műszaki adatok: n/a
- előírások: n/a

Ez a megfelelőségi nyilatkozat az orvostechnikai eszközökről szóló (EU) 2017/745 RENDELET (regisztrációs szám: HZ 1123799-1) tanúsítványának lejáratáig érvényes, és csak a kapcsolódó gyártási tétel felszabadítási dokumentummal együtt.

We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the ***Regulation (EU) 2017/745 on Medical Devices*** *and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Gyártó *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Németország/*Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
egyedi regisztrációs számot *Single Registration Number (SRN)*	DE-MF-000005414
Termékcsalád *Name of product family*	Forrasz nemesfémmentes ötvözetekhez *Base-metal Brazing Materials*
terméknév és REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Rendeltetés *Intended purpose*	A forrasz olyan töltőanyag, amelyet fogászati restaurációk, mint például modellöntvények és kombinált munkák forrasztására szántak. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Termékosztályozás *Product class*	Osztály IIa *Class IIa*
alapvető UDI-DI-nek *Basic UDI-DI*	++EBGOMetal-braz-matUU
bejelentett szervezet	TÜV Rheinland LGA Products GmbH

Notified Body	Tillystrasse 2 90431 Nürnberg Németország *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
CE-jelölés *Marked*	CE 0197

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DECLARAÇÃO DE CONFORMIDADE
DECLARATION OF CONFORMITY

De acordo com o anexo IX, Regulamentos (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Declaramos, sob nossa exclusiva responsabilidade, que os produtos abaixo indicados cumprem os requisitos gerais de segurança e desempenho relevantes, de acordo com do **Regulamento (UE) 2017/745 relativo aos Dispositivos Médicos** e estão em conformidade com

- a lista de normas aplicadas
- especificações comuns: n/a
- regulamentos: n/a

Esta declaração de conformidade é válida até ao termo de validade do Certificado REGULAMENTO (UE) 2017/745 relativo aos Dispositivos Médicos (N.º de registo: HZ 1123799-1) e apenas em conjunto com o documento correspondente de libertação do lote.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Fabricante *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Alemanha */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
número único de registo *Single Registration Number (SRN)*	DE-MF-000005414
Família de produtos *Name of product family*	Solda para ligas sem metais nobres *Base-metal Brazing Materials*
nome de produto y REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Finalidade *Intended purpose*	Uma massa de enchimento, destinada à soldadura de restaurações dentárias, como fundição de modelos e trabalhos de combinação. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
Classe *Product class*	Classe IIa *Class IIa*
UDI-DI básico *Basic UDI-DI*	++EBGOMetal-braz-matUU
organismo notificado *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2

	90431 Nuremberga Alemanha *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
marcação CE *Marked*	CE 0197

IZJAVO O SKLADNOSTI
DECLARATION OF CONFORMITY

V skladu s Prilogo IX, Uredbe (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

S tem na izključno lastno odgovornost izjavljamo, da spodaj navedeni izdelki izpolnjujejo veljavne splošne zahteve glede varnosti in učinkovitosti v skladu s **Uredbe (EU) 2017/745 o medicinskih pripomočkih** in so v skladu

- s seznamom uporabljenih standardov
- s skupnimi specifikacijami: n/a
- s predpisi: n/a

Ta izjava o skladnosti je veljavna do poteka certifikata o UREDBI (EU) 2017/745 o medicinskih pripomočkih (registrska št.: HZ 1123799-1) in samo skupaj z ustreznim dokumentom o sprostitvi serije.

*We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the **Regulation (EU) 2017/745 on Medical Devices** and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Proizvajalcev *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Nemčija */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
enotno registrsko številko *Single Registration Number (SRN):*	DE-MF-000005414
družina izdelkov *Name of product family*	Spajka za zlitine brez plemenitih kovin *Base-metal Brazing Materials*
ime izdelka in REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Predvidena uporaba *Intended purpose*	Je polnilo, namenjeno spajkanju zobnih restavracij, kot so delne proteze z ulito kovinsko bazo in kombinirana dela. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
razvrstitev izdelka *Product class*	Razred IIa *Class IIa*
Osnovni UDI-DI *Basic UDI-DI*	++EBGOMetal-braz-matUU
priglašenega organa *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg

	Nemčija *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
oznaka CE *Marked*	CE 0197

δήλωση συμμόρφωσης ΕΕ
DECLARATION OF CONFORMITY

σύμφωνα με το παράρτημα IX, ΚΑΝΟΝΙΣΜΟΙ (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

Με το παρόν δηλώνουμε με αποκλειστική μας ευθύνη ότι τα προϊόντα που αναφέρονται παρακάτω συμμορφώνονται τις σχετικές γενικές απαιτήσεις ασφάλειας και επιδόσεων σύμφωνα με του **Κανονισμού (ΕΕ) 2017/745 για τα ιατροτεχνολογικά προϊόντα** και είναι σύμφωνες με

- τον κατάλογο των εφαρμοζόμενων προτύπων
- κοινές προδιαγραφές: n/a
- τους κανονισμούς: n/a

Η παρούσα δήλωση συμμόρφωσης ισχύει μέχρι τη λήξη του Πιστοποιητικού σύμφωνα με τον ΚΑΝΟΝΙΣΜΟ (ΕΕ) 2017/745 για τα ιατροτεχνολογικά προϊόντα (Αριθ. πρωτοκόλλου: HZ 1123799-1) και μόνο σε συνδυασμό με το σχετικό έγγραφο αποδέσμευσης παρτίδων.

We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the ***Regulation (EU) 2017/745 on Medical Devices*** *and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Κατασκευαστές *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Γερμανία */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
ενιαίο αριθμό καταχώρισης *Single Registration Number (SRN)*	DE-MF-000005414
οικογένεια προϊόντω *Name of product family*	Συγκολλητικό κράμα για κράματα χωρίς ευγενή μέταλλα *Base-metal Brazing Materials*
όνομα προϊόντος και REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Προβλεπόμενη χρήση *Intended purpose*	Το συγκολλητικό κράμα κοβαλτίου-χρωμίου αποτελεί πληρωτικό που προορίζεται για την ετερογενή συγκόλληση οδοντικών αποκαταστάσεων, όπως χυτών προπλα σμάτων και εργασιών συνδυασμού. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
ταξινόμηση προϊόντος *Product class*	Κατηγορία IIa *Class IIa*
βασικό UDI-DI *Basic UDI-DI*	++EBGOMetal-braz-matUU
κοινοποιημένο οργανισμό	TÜV Rheinland LGA Products GmbH

Notified Body	Tillystrasse 2 90431 Νυρεμβέργη Γερμανία *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
σήμανση CE *Marked*	CE 0197

ЕС декларацията за съответствие
DECLARATION OF CONFORMITY

съгласно приложение IX, РЕГЛАМЕНТИ (EU) 2017/745
according to annex IX, Regulation (EU) 2017/745

С настоящото декларираме под пълна отговорност, че продуктите, изброени по-долу, отговарят на съответните общи изисквания за безопасност и действие съгласно от **Регламент (ЕС) 2017/745 за медицинските изделия** и са в съответствие със

- списъка с прилагани стандарти
- общи спецификации: n/a
- регламенти: n/a

Настоящата декларация за съответствие е валидна до изтичане на срока на валидност на сертификата по РЕГЛАМЕНТ (ЕС) 2017/745 за медицинските изделия (регистрационен номер: HZ 1123799-1) и само заедно със съответния документ за освобождаване на партидата.

We herewith declare under sole responsibility that the products listed below are in conformity with the relevant general safety and performance requirements according to the ***Regulation (EU) 2017/745 on Medical Devices*** *and is in conformity with*

- *the list of applied standards*
- *Common specifications: n/a*
- *Regulations: n/a*

This declaration of conformity is valid until expiration of the Certificate REGULATION (EU) 2017/745 on Medical Devices (Registration No.: HZ 1123799-1) and only together with the related batch release document.

Производителит *Manufacturer*	BEGO Bremer Goldschlägerei Wilh. Herbst GmbH & Co. KG Wilhelm-Herbst-Str. 1 28359 Bremen, Германия */Germany* T. +49 421 2028-0 F. +49 421 2028-100 www.bego.com
единен регистрационен номер (EPH) *Single Registration Number (SRN)*	DE-MF-000005414
продуктова група *Name of product family*	Припой за сплави, несъдържащи благородни метали *Base-metal Brazing Materials*
наименование на продукта и REF *Name of products and REF*	Wiron®-Lot 52625 Wirobond®-Lot 52622 Kobalt-Chrom-Lot 52520
Предназначение *Intended purpose*	е пълнител, предназначен за запояване на дентални възстановявания, като моделно лети и комбинирани протези. *A filler material and intended for brazing of dental restorations such as partial dentures and combination works.*
класификация на продукта *Product class*	Клас IIa *Class IIa*
Базовият UDI-DI *Basic UDI-DI*	++EBGOMetal-braz-matUU
нотифицирания орган	TÜV Rheinland LGA Products GmbH

Notified Body	Tillystrasse 2 90431 Нюрнберг Германия *TÜV Rheinland LGA Products GmbH* *Tillystrasse 2* *90431 Nuremberg* *Germany*
маркировка CE *Marked*	CE 0197