KULZER
MITSUI CHEMICALS GROUP

KONFORMITÄTSERKLÄRUNG / *DECLARATION OF CONFORMITY*

Name und Adresse der Firma / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Deutschland / *Germany* *SRN: DE-MF-000007705*

Wir erklären in alleiniger Verantwortung, dass / *We declare under our sole responsibility that*

das Medizinprodukt / *the medical device*	**GLUMA 2Bond**
Bezeichnung, Typ oder Modell, Chargen- oder Seriennummer, ev. Herkunft und Stückzahl / *Name, type or model, batch or serial number, possibly sources and number of items*	Artikelliste siehe Anhang / *List of Articles see Annex*
EMDN-Nummer / *EMDN-Code* GMDN-Nummer / *GMDN code* UMDNS-Nummer / *UMDNS code* Basis-UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
der Klasse / *of class*	IIa
nach Regel / *according to rule*	8-1 nach Anhang VIII der Medizinprodukte-Verordnung, 2017/745 / *according to Annex VIII of Medical Device Regulation 2017/745*

allen Anforderungen der Medizinprodukte-Verordnung 2017/745 entspricht, die anwendbar sind / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Angewandte harmonisierte Normen, nationale Normen oder andere normative Dokumente / *Applied harmonised standards, national standards or other normative documents*	Weitere angewandte Normen siehe Version 02 der Technischen Dokumentation von Product **GLUMA 2Bond** / *Further Applied standards see Technical Documentation of Product **GLUMA 2Bond**, Version 02*
Konformitätsbewertungsverfahren nach / *Conformity assessment procedure acc. to*	Medizinprodukte-Verordnung 2017/745 Anhang IX, Kapitel I, Abschnitt 2 und 3 and Kapitel III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Benannte Stelle / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Germany CE 0197
Registrierungsnr. / *Registration No.*:	HZ 1198082-1
Versionsnummer / *Version number*	01
Ersetzt Konformitätserklärung vom / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Ort, Datum / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Name und Funktion / *Name and function*

Diese Konformitätserklärung ist gültig für 2 Jahre in Verbindung mit den Freigabe-Dokumenten für die jeweilige Charge der produzierten Medizinprodukte / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Artikelliste / *List of Articles*
Anhang zur Konformitätserklärung / Annex to declaration of conformity

das Medizinprodukt / *for the medical device*	**GLUMA 2Bond**
Versionsnummer Artikelliste/ *Version number article list*	**V02**
Ersetzt Artikelliste vom / *Replaces article list from*	**V01, 28.03.2022**
Diese Artikelliste ist gültig für die Konformitätserklärung Version/ *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Ort, Datum / *Place, date*

i.V.
Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Name und Funktion / *Name and function*

ДЕКЛАРАЦИЯ ЗА СЪОТВЕТСТВИЕ / *DECLARATION OF CONFORMITY*

Име и адрес на фирмата / *Name and address of the company*	*Kulzer GmbH* *Leipziger Straße 2, 63450 Hanau* *Германия / Germany* *SRN: DE-MF-000007705*

Декларираме на наша собствена отговорност, че / *We declare under our sole responsibility that*

медицинското изделие / *the medical device*	**GLUMA 2Bond**
Наименование, тип или модел, партиден или сериен номер, евентуално произход и брой елементи / *Name, type or model, batch or serial number, possibly sources and number of items*	Списък с артикули, вижте Приложението / *List of Articles see Annex*
Код по EMDN / *EMDN-Code* Код по GMDN / *GMDN code* Код по UMDNS / *UMDNS code* Основна UDI-DI идентификация / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
от клас / *of class*	IIa
съгласно правило / *according to rule*	8-1 съгласно Приложение VIII от Регламента за медицинските изделия 2017/745 / *according to Annex VIII of Medical Device Regulation 2017/745*

отговаря на всички разпоредби на Регламента за медицинските изделия 2017/745, който се прилага за него. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Приложени хармонизирани стандарти, национални стандарти или други нормативни документи / *Applied harmonised standards, national standards or other normative documents*	Други приложени стандарти, вижте техническата документация на продукт **GLUMA 2Bond** Версия 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Процедура за оценка на съответствието съгласно / *Conformity assessment procedure acc. to*	Регламента за медицинските изделия 2017/745 Приложение IX, глава I, раздел 2 и 3 и глава III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Нотифициран орган / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Германия CE 0197
Регистрационен номер / *Registration number*:	HZ 1198082-1
Номер на версия / *Version number*	01
Заменя Декларация за съответствие от / *Replaces Declaration of Conformity from*	28.03.2022
Ханау, Aug 15, 2022 Място, дата / *Place, date*	i.V. от името на д-р Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Име и длъжност / *Name and function*

Тази Декларация за съответствие е валидна за 2 години във връзка с публикуваните документи за съответната партида произведени устройства / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Списък с артикули / *List of Articles*
Приложение / *Annex*: Декларация за съответствие / *Declaration of Conformity*

Медицинското изделие / *The medical device*	**GLUMA 2Bond**
Номер на версия / *Version number*	**V02**
Заменя Приложението от / *Replaces Annex from*	**V01, 28.03.2022**
Този списък със статии е валиден във връзка с декларацията за съответствие, версия / This This *article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Ханау, Nov 23, 2022

Място, дата / *Place, date*

от името на д-р Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Име и длъжност / *Name and function*

PROHLÁŠENÍ O SHODĚ / *DECLARATION OF CONFORMITY*

Název a adresa společnosti / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Německo / *Germany* *SRN: DE-MF-000007705*

Prohlašujeme na svou výlučnou zodpovědnost, že / *We declare under our sole responsibility that*

zdravotnický prostředek / *the medical device*	**GLUMA 2Bond**
Název, typ nebo model, šarže nebo výrobní číslo, případně zdroje a počet kusů / *Name, type or model, batch or serial number, possibly sources and number of items*	Seznam položek je uveden v příloze / *List of Articles see Annex*
Kód EMDN / *EMDN-Code* Kód GMDN / *GMDN code* Kód UMDNS / *UMDNS code* Základní UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
třídy / *of class*	IIa
podle pravidla / *according to rule*	8-1 podle přílohy VIII k nařízení 2017/745 o zdravotnických prostředcích / *according to Annex VIII of Medical Device Regulation 2017/745*

splňuje všechna ustanovení nařízení 2017/745 o zdravotnických prostředcích, která se ho týkají. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Použité harmonizované normy, národní normy nebo jiné normativní dokumenty / *Applied harmonised standards, national standards or other normative documents*	Další použité normy najdete v technické dokumentaci k výrobku **GLUMA 2Bond**, verze 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Procedura posouzení shody podle / *Conformity assessment procedure acc. to*	nařízení 2017/745 o zdravotnických prostředcích, příloha IX, kapitola I, oddíl 2 a 3 a kapitola III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Notifikovaná osoba / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Německo CE 0197
Registrační číslo / *Registration number*:	HZ 1198082-1
Číslo verze / *Version number*	01
Nahrazuje Prohlášení o shodě ze dne / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH**
Místo, datum / *Place, date*	Jméno a funkce / *Name and function*

Toto prohlášení o shodě je platné po dobu 2 let ve spojení s příbalovými informacemi pro příslušnou šarži vyrobených zdravotnických prostředků. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices*.

Seznam položek / *List of Articles*
Příloha / *Annex*: Prohlášení o shodě / *Declaration of Conformity*

Zdravotnický prostředek / *The medical device*	**GLUMA 2Bond**
Číslo verze / *Version number*	**V02**
Nahrazuje přílohu ze dne / *Replaces Annex from*	**V01, 28.03.2022**
Tento seznam zboží platí pro verzi prohlášení o shodě / *This article list is valid for the declaration of conformity version:*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Místo, datum / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Jméno a funkce / *Name and function*

VASTAVUSDEKLARATSIOON / *DECLARATION OF CONFORMITY*

Ettevõtte nimi ja aadress / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Saksamaa / *Germany* *SRN: DE-MF-000007705*

Ettevõte kinnitab oma ainuvastusel, et / *We declare under our sole responsibility that*

meditsiiniseade / *the medical device*	**GLUMA 2Bond**
Nimi, tüüp või mudel, partii või seerianumber, võimalikud allikad ja osad / *Name, type or model, batch or serial number, possibly sources and number of items*	Artiklite loendit vt lisast / *List of Articles see Annex*
EMDN-kood / *EMDN-Code* GMDN-i kood / *GMDN code* UMDNS-i kood / *UMDNS code* Põhi-UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
klassil / *of class*	IIa
kooskõlas reegliga / *according to rule*	8-1 kooskõlas meditsiiniseadme määrusega 2017/745 lisaga VIII / *according to Annex VIII of Medical Device Regulation 2017/745*

vastab kõigile meditsiiniseadme määruse 2017/745 sätetele, mis seadmele kohaldub. /
meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.

Kehtivad harmoniseeritud standardid, riiklikud standardid ja muud normatiivdokumendid / *Applied harmonised standards, national standards or other normative documents*	Muid rakenduvaid standardeid vt tehnilistest dokumentidest tootele **GLUMA 2Bond**, Version 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Vastavushindamismenetlus vastavalt / *Conformity assessment procedure acc. to*	meditsiiniseadmete määruse 2017/745 IX. lisa, I. peatükk, 2. ja 3. jaotis ning III. peatükk *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Teavitatud asutus / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Saksamaa CE 0197
Registreerimisnumber / *Registration number*:	HZ 1198082-1
Versiooni number / *Version number*	01
Asendab vastavusdeklaratsiooni alates / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Koht, kuupäev / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Nimi ja funktsioon / *Name and function*

See vastavusdeklaratsioon kehtib 2 aastat koos toodetud seadmete vastava partii väljastusdokumentidega / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Artiklite loend / *List of Articles*
lisa / *Annex*: Vastavusdeklaratsioon / *Declaration of Conformity*

Meditsiiniseade / *The medical device*	**GLUMA 2Bond**
Versiooni number / *Version number*	**V02**
Asendab lisa alates / *Replaces Annex from*	**V01, 28.03.2022**
See artiklite loend kehtib vastavusdeklaratsiooni versiooni kohta / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Koht, kuupäev / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Nimi ja funktsioon / *Name and function*

DECLARACIÓN DE CONFORMIDAD / *DECLARATION OF CONFORMITY*

Nombre y dirección de la empresa / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Alemania / *Germany* *SRN: DE-MF-000007705*

Declaramos bajo nuestra exclusiva responsabilidad que / *We declare under our sole responsibility that*

el producto sanitario / *the medical device*	**GLUMA 2Bond**
Nombre, tipo o modelo, lote o número de serie, posiblemente fuentes y número de elementos / *Name, type or model, batch or serial number, possibly sources and number of items*	Lista de artículos en el Anexo / *List of Articles see Annex*
Código EMDN / *EMDN-Code* Código GMDN / *GMDN code* Código UMDNS / *UMDNS code* UDI-DI básico / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
de la clase / *of class*	IIa
de acuerdo con la norma / *according to rule*	8-1 de acuerdo con el Anexo VIII del Reglamento sobre productos sanitarios 2017/745 / *according to Annex VIII of Medical Device Regulation 2017/745*

cumple todas las disposiciones del Reglamento sobre productos sanitarios 2017/745 que se le aplican. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Normas armonizadas, normas nacionales u otros documentos normativos que se aplican / *Applied harmonised standards, national standards or other normative documents*	Para otras normas aplicadas consulte la documentación técnica del producto **GLUMA 2Bond**, versión 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Procedimiento de evaluación de la conformidad de acuerdo con / *Conformity assessment procedure acc. to*	Reglamento sobre productos sanitarios 2017/745 Anexo IX, Capítulo I, Secciones 2 y 3 y Capítulo III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Organismo notificado / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Alemania CE 0197
Número de registro / *Registration number*:	HZ 1198082-1
Número de versión / *Version number*	01
Sustituye a la declaración de conformidad del / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Lugar, fecha / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Nombre y cargo / *Name and function*

La presente declaración de conformidad tendrá una validez de 2 años según la documentación emitida para el correspondiente lote de productos fabricados. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Lista de artículos / *List of Articles*
Anexo / *Annex*: Declaración de conformidad / *Declaration of Conformity*

El producto sanitario / *The medical device*	**GLUMA 2Bond**
Número de versión / *Version number*	**V02**
Sustituye al Anexo del / *Replaces Annex from*	**V01, 28.03.2022**
Esta lista de artículos es válida para la versión de la declaración de conformidad / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Lugar, fecha / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Nombre y cargo / *Name and function*

ΔΗΛΩΣΗ ΣΥΜΜΟΡΦΩΣΗΣ / *DECLARATION OF CONFORMITY*

Επωνυμία και διεύθυνση εταιρείας / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Γερμανία / *Germany* *SRN: DE-MF-000007705*

Δηλώνουμε με δική μας ευθύνη ότι / *We declare under our sole responsibility that*

το ιατροτεχνολογικό προϊόν / *the medical device*	**GLUMA 2Bond**
Επωνυμία, τύπος ή μοντέλο, παρτίδα ή αριθμός σειράς, πιθανές πηγές και αριθμός ειδών / *Name, type or model, batch or serial number, possibly sources and number of items*	Κατάλογος ειδών Παράρτημα / *List of Articles see Annex*
Κωδικός EMDN / *EMDN-Code* Κωδικός GMDN / *GMDN code* Κωδικός UMDNS / *UMDNS code* Βασικό UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
κλάσης / *of class*	IIa
σύμφωνα με τον κανόνα / *according to rule*	8-1 σύμφωνα με το Παράρτημα VIII του Κανονισμού 2017/745 για τα ιατροτεχνολογικά προϊόντα / *according to Annex VIII of Medical Device Regulation 2017/745*

πληροί όλες τις ισχύουσες διατάξεις του Κανονισμού 2017/745 για τα ιατροτεχνολογικά προϊόντα. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Εφαρμοζόμενα εναρμονισμένα πρότυπα, εθνικά πρότυπα ή άλλα κανονιστικά έγγραφα / *Applied harmonised standards, national standards or other normative documents*	Για περαιτέρω εφαρμοζόμενα πρότυπα βλ. την τεχνική τεκμηρίωση του προϊόντος **GLUMA 2Bond**, έκδοση 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version* 2
Διαδικασία αξιολόγησης συμμόρφωσης σύμφωνα με / *Conformity assessment procedure acc. to*	Κανονισμός 2017/745 για τα ιατροτεχνολογικά προϊόντα, Παράρτημα IX, Κεφάλαιο I, Τμήμα 2 και 3, και Κεφάλαιο III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Κοινοποιημένος οργανισμός / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Γερμανία CE 0197
Αριθμός καταχώρησης / *Registration number*:	HZ 1198082-1
Αριθμός έκδοσης / *Version number*	01
Αντικαθιστά τη δήλωση συμμόρφωσης από / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Τόπος, ημερομηνία / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Ονοματεπώνυμο και τίτλος / *Name and function*

Αυτή η δήλωση συμμόρφωσης ισχύει για 2 χρόνια σε σχέση με τα έγγραφα κυκλοφορίας για την αντίστοιχη παρτίδα των παραγόμενων προϊόντων. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Κατάλογος ειδών / *List of Articles*
Παράρτημα / *Annex*: Δήλωση συμμόρφωσης / *Declaration of Conformity*

Το ιατροτεχνολογικό προϊόν / *The medical device*	**GLUMA 2Bond**
Αριθμός έκδοσης / *Version number*	**V02**
Αντικαθιστά το Παράρτημα από / *Replaces Annex from*	**V01, 28.03.2022**
Αυτός ο κατάλογος προϊόντων ισχύει για την έκδοση δήλωσης συμμόρφωσης / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Τόπος, ημερομηνία / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Ονοματεπώνυμο και τίτλος / *Name and function*

IZJAVA O USKLAĐENOSTI / *DECLARATION OF CONFORMITY*

Naziv i adresa tvrtke / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Njemačka / *Germany* *SRN: DE-MF-000007705*

Izjavljujemo pod punom odgovornošću da / *We declare under our sole responsibility that*

medicinski proizvod / *the medical device*	**GLUMA 2Bond**
Naziv, tip ili model, broj serije, po mogućnosti izvori i broj stavki / *Name, type or model, batch or serial number, possibly sources and number of items*	Popis artikala, pogledajte Dodatak / *List of Articles see Annex*
šifra EMDN / *EMDN-Code* šifra GMDN / *GMDN code* šifra UMDNS / *UMDNS code* osnovni UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
klase / *of class*	IIa
u skladu s pravilom / *according to rule*	8-1 u skladu s Dodatkom VIII Uredbe 2017/745 o medicinskim proizvodima / *according to Annex VIII of Medical Device Regulation 2017/745*

ispunjava sve odredbe Uredbe 2017/745 o medicinskim proizvodima koje se na njega odnose. /
meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.

Primijenjene usklađene norme, državne norme ili drugi normativni dokumenti / *Applied harmonised standards, national standards or other normative documents*	Druge primijenjene norme, pogledajte Tehničku dokumentaciju za proizvod **GLUMA 2Bond**, verzija 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Postupak procjene usklađenosti prema / *Conformity assessment procedure acc. to*	Prilog IX Uredbi 2017/745 o medicinskim proizvodima, Poglavlje I, Odjeljak 2 i 3 te Poglavlje III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Obaviješteno tijelo / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Njemačka CE 0197
Registracijski broj / *Registration number*:	HZ 1198082-1
Broj verzije / *Version number*	01
Zamjenjuje Izjavu o usklađenosti od / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Mjesto, datum / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Ime i funkcija / *Name and function*

Ova Izjava o usklađenosti valjana je 2 godine u odnosu na dokumente o izdanju za odgovarajuće serije proizvedenih umedicinskih proizvoda. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Popis artikala / *List of Articles*
Dodatak / *Annex*: Izjava o usklađenosti / *Declaration of Conformity*

Medicinski proizvod / *The medical device*	**GLUMA 2Bond**
Broj verzije / *Version number*	**V02**
Zamjenjuje Dodatak od / *Replaces Annex from*	**V01, 28.03.2022**
Ovaj popis artikala valjan je za verziju izjave u sukladnosti / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	*Article number*	*Article name*
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Mjesto, datum / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Ime i funkcija / *Name and function*

MEGFELELŐSÉGI NYILATKOZAT / *DECLARATION OF CONFORMITY*

A vállalat neve és címe / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Németország / *Germany* *SRN: DE-MF-000007705*

Kizárólagos felelősségünkre kijelentjük, hogy / *We declare under our sole responsibility that*

az orvostechnikai eszköz / *the medical device*	**GLUMA 2Bond**
Név, típus vagy modell, tétel vagy sorozatszám, esetleg források és tételek száma / *Name, type or model, batch or serial number, possibly sources and number of items*	A cikkek listáját lásd a mellékletben / *List of Articles see Annex*
EMDN kód / *EMDN-Code* GMDN kód / *GMDN code* UMDNS kód / *UMDNS code* Alapvető UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
osztálya / *of class*	IIa
a következő szabály szerint / *according to rule*	8-1 az orvostechnikai eszközökről szóló 2017/745 rendelet VIII. melléklete szerint / *according to Annex VIII of Medical Device Regulation 2017/745*

megfelel az orvostechnikai eszközökről szóló, 2017/745 rendelet valamennyi rá vonatkozó rendelkezésének. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Alkalmazott harmonizált szabványok, nemzeti szabványok vagy más normatív dokumentumok / *Applied harmonised standards, national standards or other normative documents*	További alkalmazott szabványokat lásd a műszaki dokumentációban, termék: **GLUMA 2Bond**, 2 verzió *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Megfelelőségértékelési eljárás a következő szerint /*Conformity assessment procedure acc. to*	Az orvostechnikai eszközökről szóló, 2017/745 rendelet IX. függeléke, az I. fejezet 2. és 3. szakasza, és a III. fejezet *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Bejelentett szervezet / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Németország CE 0197
Regisztrációs szám / *Registration number*:	HZ 1198082-1
Verziószám / *Version number*	01
Felváltja a megfelelőségi nyilatkozatot ettől / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Hely, dátum / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Név és funkció / *Name and function*

Ez a megfelelőségi nyilatkozat 2 évig érvényes a gyártott eszközök adott tételére vonatkozó kibocsátási dokumentumokkal együtt. / This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.

Cikkek listája / *List of Articles*
Melléklet / *Annex*: Megfelelőségi nyilatkozat / *Declaration of Conformity*

Az orvostechnikai eszköz / *The medical device*	**GLUMA 2Bond**
Verziószám / *Version number*	**V02**
Felváltja a mellékletet ettől / *Replaces Annex from*	**V01, 28.03.2022**
Ez a tétellista a megfelelőségi nyilatkozat következő verziójához érvényes / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Hely, dátum / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Név és funkció / *Name and function*

ATITIKTIES DEKLARACIJA / *DECLARATION OF CONFORMITY*

Bendrovės pavadinimas ir adresas / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Vokietija / *Germany* *SRN: DE-MF-000007705*

Prisiimdami visą atsakomybę pareiškiame, kad / *We declare under our sole responsibility that*

medicinos prietaisas / *the medical device*	**GLUMA 2Bond**
Pavadinimas, tipas arba modelis, partija arba serijos numeris, galimi šaltiniai ir elementų skaičius / *Name, type or model, batch or serial number, possibly sources and number of items*	Prekių sąrašo ieškokite Priede / *List of Articles see Annex*
EMDN kodas / *EMDN-Code* GMDN kodas / *GMDN code* UMDNS kodas / *UMDNS code* Pagrindinis UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
klasės / *of class*	IIa
pagal taisyklę / *according to rule*	8-1 Pagal Medicinos prietaisų reglamento 2017/745 VIII priedą / *according to Annex VIII of Medical Device Regulation 2017/745*

atitinka visas jam taikomas Medicinos prietaisų reglamento 2017/745 sąlygas. /
meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.

Taikomi harmonizuotieji standartai, nacionaliniai standartai ar kiti normatyviniai dokumentai / *Applied harmonised standards, national standards or other normative documents*	atkuriamosiomis medžiagomis Kitus taikomus standartus žr. produkto **GLUMA 2Bond**, techninėje dokumentacijoje, 2 versijoje *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
]Atitikties patvirtinimo procedūra pagal / *Conformity assessment procedure acc. to*	Medicinos priemonių reglamento 2017/745 IX priedas, I skyrius, 2 ir 3 straipsniai bei III skyrius *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Notifikuotoji įstaiga / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Vokietija CE 0197
Registracijos numeris / *Registration number*:	HZ 1198082-1
Versijos numeris / *Version number*	01
Pakeičia atitikties deklaraciją nuo / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH**
Vieta, data / *Place, date*	Vardas, pavardė ir pareigos / *Name and function*

Ši atitikties deklaracija galioja 2 metus kartu su atitinkamos pagamintų priemonių partijos pateikimo į rinką dokumentais. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Prekių sąrašas / *List of Articles*
Priedas / *Annex*: Atitikties deklaracija / *Declaration of Conformity*

Medicinos prietaisas / *The medical device*	**GLUMA 2Bond**
Versijos numeris / *Version number*	**V02**
Pakeičia Priedą nuo / *Replaces Annex from*	**V01, 28.03.2022**
Šis straipsnių sąrašas tinka atitikties deklaracijai, kurios versija yra / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Vieta, data / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Vardas, pavardė ir pareigos / *Name and function*

ATBILSTĪBAS DEKLARĀCIJA / *DECLARATION OF CONFORMITY*

Uzņēmuma nosaukums un adrese / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Vācija / *Germany* *SRN: DE-MF-000007705*

Mēs vienīgi uz savu atbildību deklarējam, ka / *We declare under our sole responsibility that*

medicīniskā ierīce / *the medical device*	**GLUMA 2Bond**
Preču nosaukums, tips vai modelis, partijas vai sērijas numurs, iespējamie avoti un skaits / *Name, type or model, batch or serial number, possibly sources and number of items*	Preču sarakstu skatiet pielikumā / *List of Articles see Annex*
EMDN kods / *EMDN-Code* GMDN kods / *GMDN code* UMDNS kods / *UMDNS code* Pamata UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
klase / *of class*	IIa
saskaņā ar noteikumu / *according to rule*	8-1 saskaņā ar Medicīnisko ierīču regulas 2017/745 VIII pielikumu / *according to Annex VIII of Medical Device Regulation 2017/745*

atbilst visiem tiem Medicīnisko ierīču regulas 2017/745 noteikumiem, kas uz to attiecas. /
meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.

Izmantotie saskaņotie standarti, nacionālie standarti vai citi normatīvie dokumenti / *Applied harmonised standards, national standards or other normative documents*	Citus izmantotos standartus skatiet tehniskajā dokumentācijā par produktu **GLUMA 2Bond**, 2. versija *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Atbilstības novērtēšanas procedūra saskaņā ar / *Conformity assessment procedure acc. to*	Medicīnisko ierīču regulas 2017/745 IX pielikumu, I nodaļu, 2. un 3. iedaļu un III nodaļu *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Pilnvarotā iestāde / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg/Vācija CE 0197
Reģistrācijas numurs / *Registration number*:	HZ 1198082-1
Versijas numurs / *Version number*	01
Aizstāj atbilstības deklarāciju, kas datēta ar / *Replaces Declaration of Conformity from*	28.03.2022
Hānava Aug 15, 2022 Vieta, datums / *Place, date*	i.V. v.i. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Vārds, uzvārds un amats / *Name and function*

Šis paziņojums par atbilstību ir derīgs 2 gadus saistībā ar ražoto ierīču atbilstošās partijas izlaišanas dokumentiem. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Preču saraksts / *List of Articles*
Pielikums / *Annex*: Atbilstības deklarācija / *Declaration of Conformity*

Medicīniskā ierīce / *The medical device*	**GLUMA 2Bond**
Versijas numurs / *Version number*	**V02**
Aizstāj pielikumu, kas datēts ar / *Replaces Annex from*	**V01, 28.03.2022**
Šis izstrādājumu saraksts ir derīgs šādai atbilstības deklarācijas versijai / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hānava Nov 23, 2022

Vieta, datums / *Place, date*

v.i. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Vārds, uzvārds un amats / *Name and function*

DEKLARACJA ZGODNOŚCI / *DECLARATION OF CONFORMITY*

Nazwa i adres firmy / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Niemcy / *Germany* *SRN: DE-MF-000007705*

Niniejszym deklarujemy pod rygorem odpowiedzialności, że /
We declare under our sole responsibility that

wyrób medyczny / *the medical device*	**GLUMA 2Bond**
Nazwa, typ lub model, numer partii lub serii, ewentualnie źródła i liczba elementów / *Name, type or model, batch or serial number, possibly sources and number of items*	Wykaz wyrobów znajduje się w załączniku / *List of Articles see Annex*
Kod wyrobu wg EMDN / *EMDN-Code* Kod wyrobu wg GMDN / *GMDN code* Kod wyrobu wg UMDNS / *UMDNS code* Kod Basic UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
klasy / *of class*	IIa
zgodnie z regułą / *according to rule*	8-1 zgodnie z załącznikiem VIII do Rozporządzenia 2017/745 w sprawie wyrobów medycznych / *according to Annex VIII of Medical Device Regulation 2017/745*

spełnia wszystkie przepisy Rozporządzenia 2017/745 w sprawie wyrobów medycznych, które go dotyczą. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Zastosowane normy zharmonizowane, normy krajowe lub inne dokumenty normatywne / *Applied harmonised standards, national standards or other normative documents*	Pozostałe stosowane normy znajdują się w dokumentacji technicznej produktu **GLUMA 2Bond**, wersja 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond**, *Version 2*
Procedura oceny zgodności wg. / *Conformity assessment procedure acc. to*	Rozporządzenie 2017/745 w sprawie wyrobów medycznych, załącznik IX, rozdział I, sekcja 2 i 3 oraz rozdział III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Jednostka notyfikowana / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg/Niemcy CE 0197
Numer rejestracyjny / *Registration number*:	HZ 1198082-1
Numer wersji / *Version number*	01
Zastępuje Deklarację zgodności z / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Miejscowość, data / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Imię i nazwisko, stanowisko / *Name and function*

Niniejsze deklaracja zgodności jest ważna przez 2 lata w połączeniu z dokumentami zwolnienia odpowiedniej partii wyprodukowanych wyrobów / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices*

Wykaz wyrobów / *List of Articles*
Załącznik / *Annex*: Deklaracja zgodności / *Declaration of Conformity*

Wyrób medyczny / *The medical device*	**GLUMA 2Bond**
Numer wersji / *Version number*	**V02**
Zastępuje załącznik z dnia / *Replaces Annex from*	**V01, 28.03.2022**
Poniższa lista artykułów obowiązuje dla następujących wersji deklaracji zgodności / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Miejscowość, data / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Imię i nazwisko, stanowisko / *Name and function*

DECLARAÇÃO DE CONFORMIDADE / *DECLARATION OF CONFORMITY*

Nome e morada da empresa / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Alemanha / *Germany* *SRN: DE-MF-000007705*

Declaramos, sob nossa exclusiva responsabilidade, que / *We declare under our sole responsibility that*

o dispositivo médico / *the medical device*	**GLUMA 2Bond**
Nome, tipo ou modelo, número de lote ou de série, possivelmente origem e quantidade de itens / *Name, type or model, batch or serial number, possibly sources and number of items*	Lista de artigos, ver Anexo / *List of Articles see Annex*
Código EMDN / *EMDN-Code* Código GMDN / *GMDN code* Código UMDNS / *UMDNS code* UDI-DI básico / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
da classe / *of class*	IIa
em conformidade com o regulamento / *according to rule*	8-1 em conformidade com o Anexo VIII do Regulamento 2017/745 relativo aos Dispositivos Médicos / *according to Annex VIII of Medical Device Regulation 2017/745*

cumpre todas as disposições aplicáveis do Regulamento 2017/745 relativo aos Dispositivos Médicos. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Normas harmonizadas aplicadas, normas nacionais ou outros documentos normativos / *Applied harmonised standards, national standards or other normative documents*	Outras normas aplicadas, ver Documentação técnica do produto **GLUMA 2Bond**, Versão 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Procedimento de avaliação da conformidade de acordo com / *Conformity assessment procedure acc. to*	Anexo IX do Regulamento 2017/745 relativo aos Dispositivos Médicos, Capítulo I, secção 2 e 3 e Capítulo III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Organismo notificado / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Alemanha CE 0197
Número de registo / *Registration number*:	HZ 1198082-1
Número de versão / *Version number*	01
Substitui a Declaração de Conformidade de / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Local, data / *Place, date*	p.p. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Nome e função / *Name and function*

A presente declaração de conformidade é válida durante 2 anos em associação aos documentos do respetivo lote de dispositivos produzidos. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Listă de articole / *List of Articles*
Anexă / *Annex*: Declarație de conformitate / *Declaration of Conformity*

Dispozitivul medical / *The medical device*	**GLUMA 2Bond**
Număr versiune / *Version number*	**V02**
Înlocuiește Anexa de la / *Replaces Annex from*	**V01, 28.03.2022**
Această listă de articole este valabilă pentru declarația de conformitate versiunea / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Loc, dată / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Nume și funcție / *Name and function*

IZJAVA O SKLADNOSTI / *DECLARATION OF CONFORMITY*

Ime in naslov podjetja / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Nemčija / *Germany* *SRN: DE-MF-000007705*

Z izključno odgovornostjo izjavljamo, da / *We declare under our sole responsibility that*

medicinski pripomoček / *the medical device*	**GLUMA 2Bond**
Ime, vrsta ali model, številka šarže ali serijska številka, po možnosti izvor in število izdelkov / *Name, type or model, batch or serial number, possibly sources and number of items*	Seznam artiklov je na voljo v Prilogi / *List of Articles see Annex*
Koda EMDN / *EMDN-Code* Koda GMDN / *GMDN code* Koda UMDNS / *UMDNS code* Osnovni UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
razreda / *of class*	IIa
v skladu s členom / *according to rule*	8-1, v skladu s Prilogo VIII Uredbe o medicinskih pripomočkih 2017/745 / *according to Annex VIII of Medical Device Regulation 2017/745*

izpolnjuje vse določbe Uredbe o medicinskih pripomočkih 2017/745, ki veljajo zanj. /
meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.

Uveljavljeni usklajeni standardi, nacionalni standardi ali drugi normativni dokumenti / *Applied harmonised standards, national standards or other normative documents*	Drugi uveljavljeni standardi so na voljo v Tehnični dokumentaciji izdelka **GLUMA 2Bond**, različica 2 *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version* 2
Postopek ugotavljanja skladnosti v skladu z / *Conformity assessment procedure acc. to*	Uredbo o medicinskih pripomočkih 2017/745, Priloga IX, poglavje I, oddelka 2 in 3, poglavje III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Priglašeni organ / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg/Nemčija CE 0197
Registrska številka / *Registration number*:	HZ 1198082-1
Številka različice / *Version number*	01
Nadomešča Izjavo o skladnosti z dne / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Kraj, datum / *Place, date*	i.V. zastopnica Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Ime in položaj / *Name and function*

Ta izjava o skladnosti je veljavna 2 leti v povezavi z dokumenti o izdaji za zadevne serije proizvedenih pripomočkov. / *This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.*

Seznam artiklov / *List of Articles*
Priloga / *Annex*: Izjava o skladnosti / *Declaration of Conformity*

Medicinski pripomoček / *The medical device*	**GLUMA 2Bond**
Številka različice / *Version number*	**V02**
Nadomešča Prilogo z dne / *Replaces Annex from*	**V01, 28.03.2022**
Ta seznam izdelkov velja za naslednjo različico izjave o skladnosti / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Kraj, datum / *Place, date*

i.V. Matthias Hartmann

zastopnica Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Ime in položaj / *Name and function*

VYHLÁSENIE O ZHODE / *DECLARATION OF CONFORMITY*

Názov a adresa spoločnosti / *Name and address of the company*	**Kulzer GmbH** Leipziger Straße 2, 63450 Hanau Nemecko / *Germany* *SRN: DE-MF-000007705*

Vyhlasujeme na svoju výlučnú zodpovednosť, že / *We declare under our sole responsibility that*

zdravotnícka pomôcka / *the medical device*	**GLUMA 2Bond**
Názov, typ alebo model, číslo šarže alebo sériové číslo, prípadne zdroje a počet kusov / *Name, type or model, batch or serial number, possibly sources and number of items*	Zoznam položiek je uvedený v prílohe / *List of Articles see Annex*
Kód EMDN / *EMDN-Code* Kód GMDN / *GMDN code* Kód UMDNS / *UMDNS code* Základné identifikačné číslo UDI-DI / *Basic UDI-DI*	Q010101 34782 10-034 ++J0141103BA01010104bL8
triedy / *of class*	IIa
podľa pravidla / *according to rule*	8-1 podľa prílohy VIII k nariadeniu 2017/745 o zdravotníckych pomôckach / *according to Annex VIII of Medical Device Regulation 2017/745*

spĺňa všetky ustanovenia nariadenia 2017/745 o zdravotníckych pomôckach, ktoré sa na ňu vzťahujú. / *meets all the provisions of the Medical Device Regulation 2017/745 which apply to it.*

Použité harmonizované normy, národné normy alebo iné normatívne dokumenty / *Applied harmonised standards, national standards or other normative documents*	Ďalšie použité normy nájdete v technickej dokumentácii verzie 2 k produktu **GLUMA 2Bond** *Further Applied standards see Technical Documentation of Product* **GLUMA 2Bond***, Version 2*
Postup posúdenia zhody podľa / *Conformity assessment procedure acc. to*	prílohy IX k nariadeniu 2017/745 o zdravotníckych pomôckach, kapitola I, časť 2 a 3 a kapitola III *Medical Device Regulation 2017/745 Annex IX, Chapter I, Section 2 and 3 and Chapter III*
Notifikovaný orgán / *Notified Body*	TÜV Rheinland LGA Products GmbH Tillystrasse 2 90431 Nürnberg / Nemecko CE 0197
Registračné číslo / *Registration number*:	HZ 1198082-1
Číslo verzie / *Version number*	01
Nahrádza vyhlásenie o zhode z / *Replaces Declaration of Conformity from*	28.03.2022
Hanau, Aug 15, 2022 Miesto, dátum / *Place, date*	i.V. Dr. Matthias Hartmann Head of Global Quality, Regulatory & Scientific Services **Kulzer GmbH** Meno a funkcia / *Name and function*

Toto vyhlásenie o zhode je platné 2 roky v súvislosti s dokumentmi o uvoľnení príslušnej šarže vyrobených pomôcok / This statement of conformity is valid for 2 years in connection with the release documents for the respective batch of produced devices.

Zoznam položiek / *List of Articles*
Príloha / *Annex*: Vyhlásenie o zhode / *Declaration of Conformity*

Zdravotnícka pomôcka / *The medical device*	**GLUMA 2Bond**
Číslo verzie / *Version number*	**V02**
Nahrádza prílohu z / *Replaces Annex from*	**V01, 28.03.2022**
Tento zoznam tovaru je platný pre vyhlásenie o zhode, verzia / *This article list is valid for the declaration of conformity version*	**V01**

UDI-DI	***Article number***	***Article name***
+J014660409720	66040972	GLUMA 2Bond Bottle Refill
+J014660409710	66040971	GLUMA 2Bond Bottle Value Pack
+J014660961100	66096110	PROC GLUMA 2BOND KIT 1X4ML

Hanau, Nov 23, 2022

Miesto, dátum / *Place, date*

i.V. Dr. Matthias Hartmann
Head of Global Quality, Regulatory & Scientific Services
Kulzer GmbH

Meno a funkcia / *Name and function*

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Abschlusszertifikat

Umschlag-ID: AF0798C3625C4BC79475276EECC0E471
Status: Abgeschlossen
Betreff: Mit DocuSign abschließen: Gluma 2Bond Annex komplett.pdf
Quellumschlag:
Dokumentenseiten: 14
Signaturen: 14
Umschlagersteller:
Zertifikatsseiten: 4
Initialen: 0
Anette Stadtfeld
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Hanau, Hessen 63450
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Anette.Stadtfeld@kulzer-dental.com
IP-Adresse: 89.246.249.146

Eintragsverfolgung

Status: Original
23.11.2022 11:42:09
Inhaber: Anette Stadtfeld
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Unterzeichnerereignisse	Signatur	Zeitstempel
Matthias Hartmann matthias.hartmann@kulzer-dental.com PRRC Sicherheitsstufe: E-Mail, Kontoauthentifizierung (keine)	 Signaturübernahme: Hochgeladenes Signaturbild Mit IP-Adresse: 77.179.118.94	Gesendet: 23.11.2022 11:44:23 Eingesehen: 23.11.2022 15:41:44 Signiert: 23.11.2022 15:42:09

Vereinbarung bezüglich elektronischer Unterlagen und Signaturen:
Akzeptiert: 23.11.2022 15:41:44
ID: 3039ef38-ff89-4cda-8943-8ab19e8972cf

Vor-Ort-Unterzeichner – Ereignisse	Signatur	Zeitstempel

Bearbeiterversandereignisse	Status	Zeitstempel

Beauftragtenzustellereignisse	Status	Zeitstempel

Vermittlerversandereignisse	Status	Zeitstempel

Zertifizierter Versand - Ereignisse	Status	Zeitstempel

Kopienereignisse	Status	Zeitstempel

Zeugen-Ereignisse	Signatur	Zeitstempel

Notarereignisse	Signatur	Zeitstempel

Umschlagereignisse – Überblick	Status	Zeitstempel
Umschlag gesendet	Hash-codiert/verschlüsselt	23.11.2022 11:44:23
Zertifiziert zugestellt	Sicherheitsprüfung ausgeführt	23.11.2022 15:41:44
Signiervorgang abgeschlossen	Sicherheitsprüfung ausgeführt	23.11.2022 15:42:09
Abgeschlossen	Sicherheitsprüfung ausgeführt	23.11.2022 15:42:09

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Quellumschlag:
Dokumentenseiten: 48
Signaturen: 48
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Initialen: 0
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Unterzeichnerereignisse	Signatur	Zeitstempel
Matthias Hartmann matthias.hartmann@kulzer-dental.com PRRC Sicherheitsstufe: E-Mail, Kontoauthentifizierung (keine)	 Signaturübernahme: Hochgeladenes Signaturbild Mit IP-Adresse: 77.185.159.19	Gesendet: 28.03.2022 08:39:50 Eingesehen: 28.03.2022 09:55:51 Signiert: 28.03.2022 09:56:49

Vereinbarung bezüglich elektronischer Unterlagen und Signaturen:
Akzeptiert: 28.03.2022 09:55:51
ID: 589f77c6-d7b8-4a1e-8720-e31c5b86fc6f

Vor-Ort-Unterzeichner – Ereignisse	Signatur	Zeitstempel

Bearbeiterversandereignisse	Status	Zeitstempel

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Zertifizierter Versand - Ereignisse	Status	Zeitstempel

Kopienereignisse	Status	Zeitstempel

Zeugen-Ereignisse	Signatur	Zeitstempel

Notarereignisse	Signatur	Zeitstempel

Umschlagereignisse – Überblick	Status	Zeitstempel
Umschlag gesendet	Hash-codiert/verschlüsselt	28.03.2022 08:39:50
Zertifiziert zugestellt	Sicherheitsprüfung ausgeführt	28.03.2022 09:55:51
Signiervorgang abgeschlossen	Sicherheitsprüfung ausgeführt	28.03.2022 09:56:49
Abgeschlossen	Sicherheitsprüfung ausgeführt	28.03.2022 09:56:49

Zahlungen	Status	Zeitstempel

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Zertifikatsseiten: 4
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Matthias Hartmann matthias.hartmann@kulzer-dental.com PRRC Sicherheitsstufe: E-Mail, Kontoauthentifizierung (keine)	 Signaturübernahme: Hochgeladenes Signaturbild Mit IP-Adresse: 77.190.156.252	Gesendet: 15.08.2022 11:17:28 Eingesehen: 15.08.2022 12:31:50 Signiert: 15.08.2022 12:32:22

Vereinbarung bezüglich elektronischer Unterlagen und Signaturen:
Akzeptiert: 15.08.2022 12:31:50
ID: 96f40bbd-7093-481d-b07f-6a82a82d0224

Vor-Ort-Unterzeichner – Ereignisse	Signatur	Zeitstempel

Bearbeiterversandereignisse	Status	Zeitstempel

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Zertifizierter Versand - Ereignisse	Status	Zeitstempel

Kopienereignisse	Status	Zeitstempel

Zeugen-Ereignisse	Signatur	Zeitstempel

Notarereignisse	Signatur	Zeitstempel

Umschlagereignisse – Überblick	Status	Zeitstempel
Umschlag gesendet	Hash-codiert/verschlüsselt	15.08.2022 11:17:28
Zertifiziert zugestellt	Sicherheitsprüfung ausgeführt	15.08.2022 12:31:50
Signiervorgang abgeschlossen	Sicherheitsprüfung ausgeführt	15.08.2022 12:32:22
Abgeschlossen	Sicherheitsprüfung ausgeführt	15.08.2022 12:32:22

Zahlungen	Status	Zeitstempel

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