bredent GmbH & Co.KG Weissenhomer Str. 2 D-89250 Senden



September 12, 2023

## Extension of the validity of the MDD certificate and the MDD declarations of conformity

With the confirmation letter in the appendix, the notified body (mdc medical device certification GmbH) confirms, that the validity of the MDD certificate (according to 93/42/EEC Annex II (without 4)) is extended until May 26, 2024.

As the validity of the MDD declarations of conformity is linked to the validity of the MDD certificate, the existing MDD declarations of conformity issued in 2021 will also remain valid until the expiry of the MDD certificate.

Senden, September 12, 2023

Olaf Glück, general manager

Appendix:

- D10202\_C\_SX\_2023\_08\_22\_Confirmation\_letter\_sig
- Zertifikat bredent Richtlinie 93\_42\_EWG Anhang II englisch

bredent GmbH & Co, KG Sitz: 89250 Senden Registergericht: Amtsgericht Memmingen HRA 11358 Pers. haft. Gesellschafterin: bredent Betelligungs- und Verwallungs-GmbH Sitz: 89250 Senden Registergericht: Amtsgericht Memmingen HRB 12912 Geschäftsführer: Peter Brehm Nils Brehm Brigitte Brehm Gerald Micko Olaf Glück Ust-Ident-Nummer: DE251590726 Steuemummer: 151/153/60105 Commerzbank Ulm BIC: COBADEFF630 IBAN: DE82 6304 0053 0920 0296 00

Hypovereinsbank Ulm BIC: HYVEDEMM461 IBAN: DE40 6302 0066 0002 7594 03



mdc medical device certification GmbH · Kriegerstr. 6 · 70191 Stuttgart, Germany

bredent GmbH & Co. KG Herrn Zeiger Weißenhorner Straße 2 89250 Senden Deutschland

Your signs, your letter dated

Our signs, our letter dated

Phone number +49 711 253597-289 Date 2023-08-22

#### Preliminary Confirmation Letter – validity of MDD certificates

Sehr geehrter Herr Zeiger,

mdc medical device certification GmbH (Kriegerstr. 6, 70191 Stuttgart, Germany) has for the manufacturer:

bredent GmbH & Co. KG Weißenhorner Straße 2 89250 Senden Deutschland

issued the following certificate in accordance with Directive 93/42/EEC:

Certificate	Certificate registration No.	Certification	Date of issue	Expiry date
25193	D1020200071	Directive 93/42/EEC, Annex II,	2021-01-01	2023-09-21
		excluding 4		
25784	D1020200072	Annex V, Section 3 of the EC	2021-03-04	2023-09-21
		Directive 93/42/EEC		

These certificates were valid as of 20 March 2023 and have not been withdrawn afterwards. In accordance with the requirements of Art. 120 (2), second subparagraph, first sentence of Regulation (EU) 2017/745 on Medical Devices (MDR), last amended by Regulation (EU) 2023/607 the above mentioned certificates shall remain valid until the date set out in Art 120 (3a) MDR.

This confirmation requires that the manufacturer complies with the requirements laid out in Art. 120 (3c) and (3d) MDR and will undergo further surveillance according to the rules of mdc medical device certification GmbH.

According to the requirements of Art. 120 (3a) MDR, the prerequisite for placing the concerned devices on the market is fulfilled until at least 26 May 2024, unless otherwise specified by the authorities or this letter is replaced by a contrary notification by mdc.

Kind regards,

mdc medical device certification GmbH

i. A. Katja Karich (Team Management)



ID: 10727– 002/04.2023 mdc medical device certification GmbH

Kriegerstraße 6 70191 Stuttgart, Germany 
 Phone
 +49 711 253597-0

 Fax
 +49 711 253597-10

 e-Mail
 mdc@mdc-ce.de

 Web
 www.mdc-ce.de

General Manager Chairman of the board Registered office of the company USt.IDNr DE812169576 Harald Rentschler Dirk Palige Stuttgart AG Stuttgart HRB 21357

# **EC** Certificate

#### mdc medical device certification GmbH

Notified Body 0483 herewith certifies that

### bredent GmbH & Co. KG Weißenhorner Straße 2 89250 Senden Germany

for the scope

dental products and products for epithetics (see attachment)

has introduced and applies a

## **Quality System**

for the design, manufacture and final inspection.

The mdc audit has proven that this quality system meets all requirements according to

## Annex II – excluding Section 4 of the Council Directive 93/42/EEC

of 14 June 1993 concerning medical devices.

The surveillance will be held as specified in Annex II, Section 5.

Valid from Valid until Registration no. Report no. Stuttgart 2021-01-01 2023-09-21 D1020200071 P20-01828-190536 2020-11-03

Head of Certification Body





mdc medical device certification GmbH Kriegerstraße 6 D-70191 Stuttgart, Germany Phone: +49-(0)711-253597-0 Fax: +49-(0)711-253597-10 Internet: http://www.mdc-ce.de

Attachment of the certificate				
No. D1020200071	Date 2020-11-03	Page 1 of 1		

Product category	Product	Class	Product code
	alloys, dental	lla	10-077
	elastomers, silicone rubber	lla	11-394
	screws	lla	16-055
	dental porcelains	lla	16-187
	dental veneer	lla	16-336
	adhesives, denture	lla	16-388
	dental, hand instrument, other	lla	16-667
	restorative materials, dental, crown and bridge	lla	16-723
	restorative materials, dental, crown and bridge	llb	16-723
dental products	denture base resins	lla	16-728
	denture base resins, heat-cured	lla	16-729
	composite restorative material kits, dental, heat-cured	lla	16-735
	dental precision attachment	lla	17-113
	dental precision attachment	llb	17-113
	teeth, dental prosthesis	lla	17-114
	denture reliners, soft	lla	17-610
	prostheses, dental, fixed	fla	17-841
	dentures, partial	lla	17-845
products for epithetics	prostheses, maxillofacial	lla	13-162



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Head of Certification Body

