

## IZJAVA EU O SKLADNOSTI EU DECLARATION OF CONFORMITY

(v skladu s prilogo IV Uredbe o medicinskih pripomočkih (EU) 2017/745)  
(in accordance with Annex IV of Medical Device Regulation (EU) 2017/745)

Podjetje/ Company: **INTERDENT® d.o.o.**  
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SRN: **SI-MF-000004584**

Izjavljamo, da smo kot proizvajalec izključno odgovorni za izdajo izjave EU o skladnosti. /  
*We declare, that as a manufacturer, we are solely responsible for issuing the EU declaration of conformity.*

Sledeči proizvodi, razvrščeni v razred I (pravilo 5) po prilogi VIII MDR,  
*Following Class I Products (rule 5) according to Annex VIII of the MDR,*

<b>GENERICNO IME / GENERIC NAME</b>	<b>DENTALNI MATRIČNI TRAKOVI / DENTAL MATRIX BAND</b>
<b>TRGOVSKO IME / TRADE NAME</b>	TRAČNE MATRICE / <i>RIBBON MATRIX</i> IVORY MATRICE / <i>IVORY MATRIX</i> POLIESTERSKI TRAKOVI / <i>POLYESTER MATRIX</i>
<b>GMDN</b>	<b>38786</b>
<b>EMDN</b>	<b>Q01010401</b>
<b>OSNOVNI UDI-DI / BASIC UDI-DI</b>	++D058RIPMATRIX16R

ustrezajo splošnim zahtevam glede varnosti in učinkovitosti Uredbe o medicinskih pripomočkih (EU) 2017/745.

*comply with general safety and performance requirements of the Medical Devices Regulation (EU) 2017/745.*

### HARMONIZIRANI IN OSTALI STANDARDI / *HARMONISED AND OTHER STANDARDS:*

EN ISO 13485:2016+A11:2021 Medicinski pripomočki – Sistem vodenja kakovosti – Zahteve za zakonodajne namene / *Medical devices – Quality management systems – Requirements for regulatory purpose*

EN ISO 9001:2015- Sistem vodenja kakovosti – zahteve / *Quality management system – Requirements*

EN ISO 14971:2019/A11:2021 Medicinski pripomočki- Uporaba obvladovanja tveganja pri medicinskih pripomočkih / *Medical devices - Application of risk management to medical devices*

EN ISO 10993-1:2020 Biološko vrednotenje medicinskih pripomočkov – 1. del: Ocena in preskusi znotraj ocene tveganja / *Biological Evaluation of Medical Devices- Part 1: Evaluation and testing within a risk management process*

EN ISO 15223-1:2021 Medicinski pripomočki – Simboli za označevanje medicinskih pripomočkov - 1. Del: Splošne zahteve / *Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements*

EN ISO 7405:2018 Zobozdravstvo – Ocena biokompatibilnosti medicinskih pripomočkov v zobozdravstvu / *Dentistry – Evaluation of biocompatibility of medical devices used in dentistry*

EN ISO 10993-18:2020 Biološko ovrednotenje medicinskih pripomočkov - 18. del: Kemična opredelitev lastnosti materialov za medicinske pripomočke znotraj procesov obvladovanja tveganja / *Biological evaluation of medical devices – Part 18: Chemical characterization of materials*

ISO/TS 10993-19:2020 Biološko ovrednotenje medicinskih pripomočkov - 19. del: Fizikalno-kemijska, morfološka in topografska karakterizacija materialov / *Biological evaluation of medical devices – Part 19: Physico-chemical, morphological and topographical characterization of materials*

EN 62366-1:2015/A1:2020 Medicinski pripomočki – Del 1: Uporaba inženiringa uporabnosti medicinskih pripomočkov / *Medical devices – Part 1: Application of usability engineering to medical devices*  
ISO/TR 20416:2020 Medicinski pripomočki - Nadzor proizvajalcev po dajanju v promet / *Medical devices - Post-market surveillance for manufacturers*  
EN ISO 20417:2021 Informacije proizvajalca za medicinske pripomočke / *Information supplied by the manufacturer of medical devices*  
EN 1641:2009 Zobozdravstvo - Medicinski pripomočki za zobozdravstvo - Materiali / *Dentistry – Part 1: Medical devices for dentistry - Materials*

ANSI/HIBC 2.6 standard 2016: Standard označevanja dobaviteljev zdravstvene industrije za varnost pacientov in edinstveno identifikacijo naprave (UDI) / *The health industry supplier labeling standard for patient safety & unique device identification (UDI)*

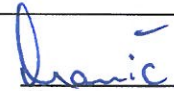
Veljavnost izjave o skladnosti je vezana na spremembo medicinskega pripomočka. / *The validity of declaration of conformity is linked to a change in medical device.*

Celje, 10.05.2023

Place, Date

Anja Mavrič, B.Sc.

Responsible person for MD and technical files



Signature:

Verzija / Version: 3