DECLARATION OF CONFORMITY

Manufacturer:	Reinvent Dental Products Unit 5 16673 148th Avenue, Spring Lake MI 49456 USA
SRN:	US-MF-000004250
EU Authorised Representative:	Eurolink Europe Compliance Limited 25 Herbert Place Dublin D02 AY86 Republic of Ireland
SRN:	IE-AR-000002852

We the undersigned certify and declare under our sole responsibility that the following apparatus

Conforming Apparatus:	NiTin Forceps
Device Name and UDI-DI Numbers:	See Page 2 below
Risk Class:	Class I - MDR Annex VIII Rule 1
Health Products Regulatory Authority Registration Reference:	MDROEO202003251341_8
Technical Document Reference No:	None
Common Specifications	EN ISO 14971:2019 Medical Devices – Application of Risk Management to Medical Devices Medical Devices Regulations – Annex I, General Requirements

conforms to the requirements of REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices

Signed: ***Date:*** 9 December 2021

Issued: Dublin, Ireland

Michael Morris
Reinvent Dental Products

Device Name	Device Model	Device UDI-DI
Band Forceps	NT-DF6-8	00810038094649
Ring Placement Forceps	NT-MRDF-100	00810038094694