EU Declaration of Conformity

according to MDR Regulation (EU) 2017/745

*for **Class I** Medical Device (non-sterile, without measuring function, non-reusable)*

Manufacturer:	Shenzhen Denco Medical Co., Ltd
Address:	Room 301, No.8, 1st road of Xiawei Industrial Zone Zhangkengjing Community, Guanhu Street, Longhua District Shenzhen, 518110, China
SRN in EUDAMED:	CN-MF-000014271

Manufacturer's authorised representative (EC Rep):
Share info GmbH
Heerdter Lohweg 83, 40549 Düsseldorf, Germany
EC Rep's **SRN** in EUDAMED: DE-AR-000005132**.**

We, the manufacturer, declare under the sole responsibility of the manufacturer that

the medical device(s)	Product Name	Model/code/Ref, (for identification/traceability)
	Dental Rulers	Endo Block, Gauge for Gutta Percha, Z-Gauge Ruler, Multi-function file holder, Soporte y Regla, Endo Ruler, Endo Ruler with slider, Endo Ruler with cleaner
Risk class	**Class I** Medical Device (non-sterile, without measuring function)	
covered by the present declaration is/are in conformity with the MDR- Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices and, if applicable, with any other relevant Union legislation.		
Notified Body (name & number), conformity assessment procedure, & **Certificate** no.	**NOT applicable**	
Basic UDI-DI	**NOT available at present**	
Common specification (CS)	**NOT applicable**	

Signed on: 21 March 2024. Place: Shenzhen, China

Signature (on behalf of the manufacturer) _____

SHENZHEN DENCO MEDICAL CO., LTD 深圳市德科医疗器械有限公司 中国 深圳 SHENZHEN CHINA

Name of authorized signatory: WU GUO JUN
Position held in the company: GENERANAL MANAGER