EU Declaration of Conformity

according to MDR Regulation (EU) 2017/745

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

Manufacturer:	**DongguanQuandingMedicalSuppliesCo.,Ltd**
Address:	**3YongfaEastRoad,QishiTown,DongguanCity, Guangdong,China**
	+86-769-86004236
SRN in EUDAMED:	CN-MF-000030836

Manufacturer's authorised representative (EC Rep):
Wellkang Ltd (www.CE-marking.eu)
Enterprise Hub, NW Business Complex,
1 Beraghmore Rd., Derry, BT48 8SE, Northern Ireland.
(EC Rep's SRN in EUDAMED: XI-AR-000001836)

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

the medical device(s)	**Product Name**	Adhesive Electrodes (TENS Electrodes)
	Model/code/Ref, (for identification/traceability)	TE-A20、TE-A21、TE-A22、TE-A23、TE-A24; TE-B20、TE-B21、TE-B22、TE-B23、TE-B24; TE-C20、TE-C21、TE-C22、TE-C23、TE-C24; TE-D20、TE-D21、TE-D22、TE-D23、TE-D24; TE-E20、TE-E21、TE-E22、TE-E23、TE-E24.
Risk class	**Class I** Medical Device (non-sterile, without measuring function, reusable)	
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: 7 Jan.2023. Place: Dongguan, Guangdong, China

Signature (on behalf of the manufacturer)

Name of authorized signatory: Xiaoting Zhao

Position held in the company: GeneralManager