EC DECLARATION OF CONFORMITY

Name and address of the firm	**Shenzhen Jumper Medical Equipment Co., Ltd.** **D Building, No. 71, Xintian Road, Fuyong Street,** **Baoan,Shenzhen, Guangdong,China** **Post Code:518103**

We declare under our sole responsibility that

the medical device	**TENS Therapy Device** **JPD-ES200, JPD-ES210, JPD-ES220**
	Name, type or model, batch or serial number, possibly sources and number of items
of class	**Class IIa, Rule 9**
	According to the classification principle in Annex VIII of Regulation (EU) 2017/745
SRN	**CN-MF-000014343**
Basic UDI-DI	**69517405ES02R3**

meets all the provisions of the Regulation (EU) 2017/745 which apply to it.

Applied harmonised standards, national standards or other normative documents	EN ISO 15223-1:2021 / EN 20417:2021 / EN ISO 14971:2019+A11:2021
	EN ISO 10993-1:2020 / EN ISO 10993-5:2009 / EN ISO 10993-10:2013
	EN 60601-1:2006+A1:2013+A12:2014
	EN 60601-1-2:2015
	EN 60601-2-10:2012 +A1:2016
	EN 60601-1-11:2015
	EN 60601-1-6:2010+A1:2015
	EN 62304:2006+A1:2015
	EN ISO 13485:2016
Conformity assessment procedure	**This declaration of conformity is based on the Regulation (EU) 2017/745, Annex IX Section 4, Chapters I and III.**
Notified Body (if consulted)	**SGS FIMKO OY** **NB 0598**
European Representative	**MedPath GmbH** **Mies-van-der-Rohe-Strasse 8** **80807 Munich, Germany**
CE certificate No.	**FI24/1008005**

2024-03-04 SHENZHEN

Manufacturer: Shenzhen Jumper Medical Equipment Co., Ltd.

Name and function
Tiffany Zhang, Regulatory Affairs Manager