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	Pulse oximeter JPD-500A, JPD-500B, JPD-500C, JPD-500D, JPD-510D, JPD-500E, JPD-510E, JPD-500F, JPD-501F, JPD-502F, JPD-500G, JPD-501G, JPD-502G, JPD-500H, JPD-500I, JPD-510I
of class	Class IIa, Rule 10 According to the classification principle in Annex VIII of Regulation (EU) 2017/745
SRN	CN-MF-000014343
Basic UDI-DI	69517405SP01TL

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 ISO 20417:2021 / EN ISO 14971:2019+ A11:2021 EN ISO 10993-1:2020/ EN ISO 10993-5:2009 / EN ISO 10993-10:2023 EN 60601-1:2006+A1:2013+A2:2021 EN 60601-1-2:2015+A1:2021 EN ISO 80601-2-61:2019 EN 60601-1-11:2015+A1：2021 EN 60601-1-6:2010+A1:2015+A2:2021/ EN 62366-1:2015+A1:2020 EN 62304:2006+A1:2015 EN ISO 13485:2016+A11:2021
Conformity assessment procedure	This declaration of conformity is based on the Regulation (EU) 2017/745, Annex IX **(I, III and TDA in Section 4).**
Notified Body (if consulted)	SGS FIMKO OY NB 0598
European Representative	MedPath GmbH Mies-van-der-Rohe-Strasse 8 80807 Munich, Germany
Authorized Representative's SRN	DE-AR-000000087
CE certificate No.	FI24/1008005

2024-03-06 Shenzhen, China

Manufacturer: Shenzhen Jumper Medical Equipment Co., Ltd.

Name and function

PRRC