EU DECLARATIONOF CONFORMITY

According to Art. 19 of Regulation (EU) 2017/745 on Medical Devices

Manufacturer:	Shanghai PAX Medical Instrument Co., Ltd. Building 3,No.438 Fulian Two Road, Baoshan District, Shanghai 201906, China
Trademark:	PANALEX
SRN:	CN-MF-000027540
EuropeanRepresentative:	MedPath GmbH Mies-van-der-Rohe-Strasse8 80807 Munich, Germany
DIMDI NO.:	DE/0000047823
SRN:	DE-AR-000000087
Product name:	Operating Table
Product Model/Type:	DS-II Series: DS-IIA、DS-IIC、DS-IIE、DS-IIF、DS-IIK、DS-IIM、DS-IIN、DS-IIS、CF280、CF380、CFT-C8、L7
Basic UDI:	697567880OT001SP
Classification acc. to MDR Ax. VIII:	Class I, rule 13
Applied Standard &Common Specification:	EN 60601-1:2006+AC:2010+A1:2013,EN 60601-2-46:2011, EN 60601-1-2:2015
Conformity assessment procedure:	Annex II + Annex III of Regulation (EU) 2017/745 MDR ;

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Sign for and behalf of the manufacturer:

General Manager

SHANGHAI PAX MEDICAL INSTRUMENT CO., LTD.

Date and Place:
09. 08. 2022 Shanghai, China