EU DECLARATION OF CONFORMITY

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

Name and address of the manufacturer:	Ningbo Tianbo First Aid Product Co., Ltd. Yushantou Village, DongqiaoTown, 315000 Haishu Ningbo, Zhejiang, China
SRN:	CN-MF-000016721
Name and address European Representative:	Prolinx GmbH Brehmstr. 56, 40239, Duesseldorf, Germany
SRN:	DE-AR-000005129
Importer:	XD Connects B.V Lange Kleiweg 6-28,2288 GK Rijswijk, The Ietherlands
Product Name:	**FIRST AID KIT**
UMDNS/GMDN/EMDN:	UMDNS-Code: 11-723
Classification acc. to MDR:	I, Ax. VIII rule 4
Basic UDI-DI:	**697471270TBJJB1NQ**
Intended Purpose:	The product is intended to use for family first aid and wound treatment.
Conformity assessment procedure:	Annex IX of Regulation EU 2017/745(MDR)
CE mark:	CE

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.

For and on behalf of
NINGBO TIANBO FIRST AID PRODUCT CO., LTD.
HAORONG QIANG
Authorized Signature(s)

NINGBO 2026.05.25