EU DECLARATION OF CONFORMITY

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

Manufacturer:	Shenzhen Besdata Technology Co., Ltd 2 / F, Building C, Hongyi Industrial Park, No.4010 Banxuegang Road, Longgang District, Shenzhen, Guangdong, 518129, China
Trademark:	BESDATA
SRN:	Not available yet
European Representative:	MedPath GmbH Mies-van-der-Rohe-Strasse 8 80807 Munich, Germany
SRN:	DE-AR-000000087
Trade name:	Video Laryngoscope
Product name:	Video Laryngoscope
Model(s):	BD-DF，BD-DP，BD-VS，BD-M，BD-MK
Basic UDI-DI:	See attachment
Classification acc. to MDR Ax. VIII:	Class I
Applied Standard & Common Specification:	EN ISO 14971:2019 EN ISO 15223-2016 EN 60601-1:2006 /A1:2013 EN 60601-1-2:2015 IEC60601-2-18:2009
Conformity assessment procedure:	Annex II + Annex III of 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.

Lu Xiaohong,
General Manager

Shenzhen, 11. 12. 2020

Basic UDI-DI			Model
GS1 Company Prefix	Internal number	Check character pair	
697135535	6971355350062	Z4	BD-DF
697135535	6971355350079	ZM	BD-DP
697135535	6971355350123	YX	BD-VS
697135535	6971355350109	Z5	BD-M
697135535	6971355350116	Z2	BD-MK