

# 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

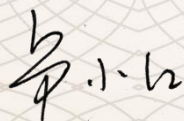
**Basic UDI-DI:** See attachment

**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

