DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES

MANUFACTURER:	GUANGDONG KINDLY MEDICAL DEVICE GROUP CO., LTD ROOM101, BUILDING A, NO.288 EAST AIRPORT ROAD, SANZAO TOWN, JINWAN DISTRICT, ZHUHAI, GUANGDONG, PEOPLE'S REPUBLIC OF CHINA
EUROPEAN REPRESENTATIVE:	SHANGHAI INTERNATIONAL HOLDING CORP. GMBH (EUROPE) EIFFESTRASSE 80,20537 HAMBURG GERMANY
MEDICAL DEVICE:	STERILE SYRINGES FOR SINGLE USE WITHOUT NEEDLE: 1ml 2ml 2.5ml 3ml 5ml 6ml 10ml 20ml 25ml 30ml 35ml 50ml 60ml
CLASSIFICATION - *ANNEX IX*:	CLASS I*, RULE1
CONFORMITY ASSESSMENT ROUTE:	ANNEX V

WE, THE MANUFACTURER, HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 CONCERNING MEDICAL DEVICES;
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY
IDENTIFICATION NUMBER	CE 0123
(EC) CERTIFICATE(S):	NO.G2S 075321 0008 REV.02
START OF CE-MARKING:	N/A
PLACE, DATE OF DECLARATION:	ZHUHAI 2020-05-28
SIGNATURE:	LiuZedong
	POSITION: QUALITY MANAGER

We, as the manufacturer, are exclusively responsible for the declaration of conformity.