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

MANUFACTURER:	Shenzhen Yimi Life Technology Co. ,Ltd 305, Building A, Tengbo Industrial Park, Changshangjiang Street, Longbei Village, Pingshan District, 518118, Shenzhen, PEOPLE'S REPUBLIC OF CHINA
EUROPEAN REPRESENTATIVE:	SHARE INFO CONSULTANT SERVICE LLC REPR SENTANZB RO
PRODUCT/MODEL:	Pulse Oximeter/ YM101,YM102,YM103,YM201,YM301
The accessories are used together with the product:	*Not applicable*
UMDNS/GMDN [name/code]:	Oximeter / 12853
CLASSIFICATION:	Class IIb, Rule 10 According To Annex IX OF the MDD
CONFORMITY ASSESSMENT ROUTE:	Annex II EXCLUDING (4)

WE, SHENZHEN YIMI LIFE TECHNOLOGY CO. LTD., HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCT(S) MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 INCLUDING AMENDMENTS BY DERECTIVE 2007/47/EC.

ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

SHENZHEN YIMI LIFE TECHNOLOGY CO. LTD IS RESPONSIBLE FOT THIS DECLARATION OF CONFORMITY

STANDARDS APPLIED:	
NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 M NCHEN, GERMANY
IDENTIFICATION NUMBER:	0123
(EC) CERTIFICATE(S):	NO.G1 104553 0001 REV.00 VALID UNTIL: 2024-05-26
START OF CE-MARKING:	2020-03-20
PLACE, DATE OF ISSUE:	SHENZHEN,
SIGNATURE:	YiYAo NAME YIYAO GENERAL MANAGER