globalcare 中山市全康医疗科技有限公司	EC Declaration of Conformity	F-QA-037 Rev.01 20160107

MANUFACTURER:	Globalcare Medical Technology CO., LTD. 7th Building, 39 Middle Industrial Main Road, European Industrial Zone, Xiaolan Town, 528415 Zhongshan City, Guangdong Province, PEOPLE'S REPUBLIC OF CHINA
Product Category:	Aerosoltherapy Nebulizers
Product Code:	95118
Product Description:	MEDEL FAMILY PLUS
Classification - Annex IX:	Class IIa, Rule 11
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; including, at 21 March 2010, the amendments by Council Directive 2007/47/EEC.
All supporting documentation is retained at the premises of the manufacturer.
Globalcare Medical Technology CO., LTD. is exclusively responsible for this EC Declaration of Conformity.

Standards Applied: see attached list

Notified Body:	TÜV SÜD Product service GmbH Ridlerstr 65, D-80339 München, Germany
Identification Number	0123
(EC) Certificate(s):	G2 088855 0010 Rev.00
EC REP European Representative:	Donawa Lifescience Consulting Srl Piazza Albania, 10 00153 Rome Italy

Place, Date of Declaration:	Zhongshan, 2019-10-24
Signature:	Name: Lambert Zhao Position: General Manager