EC Declaration of Conformity

The company

imtmedical ag

Gewerbestrasse 8
9470 Buchs SG
Switzerland

declares on sole responsibility that the following medical products of **class IIb**

Description	***Product name***	***Type(s)***
Ventilator	bellavista	1000
Ventilator	bellavista	1000e
Ventilator	bellavista	1000 neo

comply with the essential requirements (Annex I) of Directive 93/42/EEC as amended by Annex II of Directive 2007/47/EC.

Notified Body	DEKRA Certification GmbH Handwerkstrasse 15 D-70565 Stuttgart Identification no.: 0124		
Directive/Standard	*Date of Certification*	*Valid until*	*Certificate No.*
Annex II, Directive 93/42/EEC	2018-10-27	2023-10-26	50747-16-04
EN ISO 13485:2016	2018-10-27	2021-10-26	50747-14-00

The conformity of the products with the applicable provisions of Annex II excluding section (4) of Directive 93/42/EEC is declared by affixing the CE marking

CE **0124**

imtmedical ag

Buchs, October 27, 2018

Ramin Amiri, Regulatory Affairs- & Quality Manager