EC Declaration of Conformity on Medical Devices

We, the manufacturer, declare in sole responsibility that the product mentioned below is in conformity with the respective regulations of the following guideline.

Manufacturer:	WEINMANN Emergency Medical Technology GmbH + Co. KG Frohbösestraße 12, 22525 Hamburg GERMANY
Product description:	**Ventilator**
Product name / Model:	**MEDUVENT Standard**
Guideline:	Medical Device Directive 93/42/EEC (according to Annex II excluding (4))
Classification:	IIb
Accessories:	2 m reusable breathing circuit 2 m disposable breathing circuit MEDUtrigger for 2 m breathing circuit for manual breath initiation Hygiene filter
Marking:	TÜV-Rheinland LGA Products GmbH, Tillystraße 2, 90431 Nürnberg, Germany CE 0197

This EC declaration of conformity is valid from the date of signature until a revised EC declaration of conformity is issued as a result of a new directive certificate being issued.

Hamburg, 10/12/2020

André Schulte
Chief Executive Officer

WM 74550c 12/2020 EN