Certificate of CE-Registration

This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Global Med, Inc.
PO Box 340 155 North Murray Street
K8V 5R5 Trenton, Ontario
Canada

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

UMDNS Code	Description	Classification	Registration Number
10139	Anaesthesia Circuits & Accessories	IIa	DE/CA09/0760/G09/001-01
15003	Respiratory/Ventilator Circuits & Accessories	IIa	DE/CA09/0760/G09/003-01
16987	Continuous Positive Airway Pressure Units	IIa	DE/CA09/0760/G09/002-01

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 02 July 2018

Werner Sander
President