EC Declaration of Conformity

in accordance with EC Directive 93/42/EEC concerning medical devices.

Technologie Institut Medizin GmbH (TIM),
August - Thyssen Str. 30, 56070 Koblenz, Germany

herewith declares under own responsibility as manufacturer that the Design, Construction and Version of the following Medical Device meet the Requirements of the European Directive 93/42/EEC.

Product identification:	**MC-MC ISO** **MC-MC-SEVO** **MC-MC-DES**
Medical Device Trade Name:	**MIRUS Controller M**
Medical Device Class:	**Class IIb according to MDD 93/42/EEC rule 11 Annex IX**
Conformity Assessment Procedure:	**According to Annex II of MDD 93/42/EEC** **Reg.-No. 44 232 117873** Issued by TÜV NORD CERT GmbH Langemarckstr. 20, 45141 Essen Notified body registered under # 0044 This Declaration of Conformity is valid until 2024-01-20.
UDMNS Classification:	**10-144** (vaporizer for volatile anaesthetic agents)

The Quality Management System has been approved in accordance with ISO 13485: 2016 by the Notified Body TÜV NORD CERT GmbH Langemarckstr. 20, 45141 Essen, Reg.No. 4422110117873.

Any modification to the Product not authorized by TIM GmbH will invalidate this declaration.

Koblenz, 2019-01-20

Prof. Ing. Thomas P. Kriesmer
General Manager

Technologie Institut Medizin GmbH (TIM). Executive Partner Prof. Dipl.-Ing. Thomas P. Kriesmer
District Court Koblenz. HRB 20959 • Tax number: 2967605551 • VAT-ID Nr. DE 198702204
August-Thyssen-Str. 30 • D-56070 Koblenz • Germany. Tel. +49 261 899 689 00 • Fax +49 261 899 689 09
Commerzbank AG • BIC COBADEFF570 • IBAN DE66 5704 0044 0109 9126 00
info@tim-gmbh.de • www.tim-gmbh.de