

**EC Certificate**  
**Directive 93/42/EEC Annex II, excluding Section 4**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60137935 0001

**Report No.:** 21213508 015

**Manufacturer:** Hamilton Medical AG  
Via Crusch 8  
7402 Bonaduz  
Switzerland

**Products:** Ventilators and ventilator systems  
(see attachment for additional site included)  
Replaces Approval, Registration No.: HD 60136804 0001


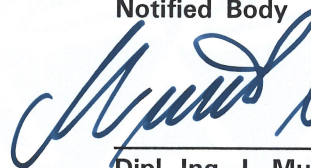
**Expiry Date:** 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2019-07-09

**Date:** 2019-04-02

Notified Body



Dipl.-Ing. I. Munkler

**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**  
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

**Attachment to  
Certificate**

**Registration No.:** HD 60137935 0001  
**Report No.:** 21213508 015


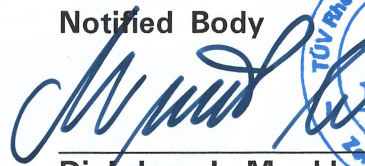
**Manufacturer:** Hamilton Medical AG  
Via Crusch 8  
7402 Bonaduz  
Switzerland

**Additional site:**

Hamilton Medical AG  
Parc Industrial Vial 10  
7013 Domat/Ems  
Switzerland

**Date:** 2019-07-09

**Notified Body**



**Dipl.-Ing. I. Munkler**