

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

**Registration No.:** HD 60127808 0001

**Report No.:** 26300174 007

**Manufacturer:** TEDISEL IBERICA, S. L.  
C/. Sant Lluc, 69-81  
08918 Badalona  
Spain

**Products:**

- Medical supply units, wall mounted and ceiling mounted
- Terminal units for use with compressed medical gases and vacuum and for anaesthetic gas scavenging systems

Replaces EC Certificate, Registration no.: HD 60084109 0001

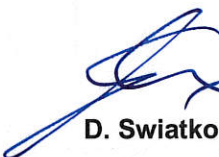
**Expiry Date:** 2023-03-11

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:** 2018-03-13

**Date:** 2018-03-13

Notified Body

  
D. Swiatko



**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.