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 Zertifizierungsstelle

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.