



Product Service

# EC-CERTIFICATE

## Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 05 05 55491 001

**Manufacturer:** **Linde Medicale S.r.l.**  
Via Pio Semeghini, 38  
00155 Roma  
ITALY

**Facility(ies):** Linde Medicale S.r.l.  
Via Pio Semeghini, 38, 00155 Roma, ITALY  
  
Linde Medicale S.r.l.  
Via G. Rossa, 3, 20010 Arluno (MI), ITALY

**Product Category(ies):** **Medical gases and vacuum supply systems, anaesthetic gas scavenging systems**

The Certification Body of TÜV Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

**Report No.:** ITA148269

**Valid until:** 2009-08-12



**Date,** 2005-05-10

TÜV Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

Page 1 of 1