

Certificate of CE-Registration



MDSS

Medical Device Safety Service

This is to certify that, in accordance with the Medical Device Directive 93/42/EEC, Medical Device Safety Service GmbH (MDSS) agrees to perform all duties and responsibilities as the Authorized Representative for:

**The Respiratory Group
4150 Carr Lane
ST. LOUIS, MO 63119
UNITED STATES OF AMERICA**

as stipulated and demanded by the aforementioned Directive. The German Competent Authority has allocated the medical devices of the Manufacturer registration numbers as foreseen in:

Annex A dated March 17, 2008

The Manufacturer has provided MDSS with the appropriate Declaration(s) of Conformity confirming that the medical devices fulfill the applicable requirements of Directive 93/42/EEC. In compliance with German law, a safety officer has been appointed for Germany.

2008-03-17



Ludger Möller
-President-
MDSS GmbH

**Annex A dated March 17, 2008
Manufacturer: The Respiratory Group**

Notified Medical Device & UMDNS Description	UMDNS Code	Risk Class	Cat. Code	Registration Number	NB No. / NB Certificate No.	NB Cert. Valid Until YYYY-MM-DD
Regulators, High-Pressure Gas	13-323	IIb	02	DE/CA09/0170/2113	0482/3332GB410070921	2012-09-21
<i>Integral Regulators</i>						
TRG CGA 540 Regulator						
TRG CGA 870 Regulator						
Liquid Oxygen Units, Individual	16-853	IIb	02	DE/CA09/0170/041/A1	0482/3332GB410070921	2012-09-21
<i>Liquid Oxygen Systems</i>						

Sup. Staff ☆☆☆



Medical Device Safety Service ☆☆☆