DET NORSKE VERITAS

PRODUCTION QUALITY MANAGEMENT CERTIFICATE - EC

Certificate No. 32116-2008-CE-NOR
This Certificate consists of 3 pages

This is to certify that the Quality Management System of

3A HEALTH CARE SRL

Lonato, Italy

for production and final product inspection/testing of

Medical Devices for Professional and Domestic use

has been assessed with respect to
the conformity assessment procedure described in Article 11.2.b and 11.5 and Annex V (Module D1) of Council Directive 93/42/EEC on Medical Devices, as amended, and found to comply

Further details are given overleaf

Place and date:
Høvik, 09 September 2008

This Certificate is valid until:
11 July 2013

For DET NORSKE VERITAS CERTIFICATION AS
Norway

for Marianne Spæren
Certification Manager

CE

Notified Body No.:
0434

Jenny Helen Nytun
Technical Reviewer

Notice: The certificate is subject to terms and conditions overleaf. Any significant changes in design or construction may render this certificate invalid.

If any person suffers loss or damage which is proved to have been caused by any negligent act or omission of Det Norske Veritas, then Det Norske Veritas shall pay compensation to such person for his proved direct loss or damage. However, the compensation shall not exceed an amount equal to ten times the fee charged for the service in question, provided that the maximum compensation shall never exceed USD 300.000. In this provision "Det Norske Veritas" shall mean the Foundation Det Norske Veritas as well as all its subsidiaries, directors, officers, employees, agents and any other acting on behalf of Det Norske Veritas.

Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history

Revision	Description	Issue Date
	Original Certificate – Re-certification	**2008-09-09**

Products covered by this Certificate

Product Description	Product Name	Class
Nebulizers	• ATOMIZER • SPEEDAIR • MODELLO 1 • MODELLO 2 • HOSPYNEB PROFESSIONAL AT7CP/3 • HOSPYNEB PROFESSIONAL AT14P/3 • MODELLO 3 • MODELLO 4 • Fastjet • Nebjet • Nebby • AIRONE (AP10XXU) • Uniko • ARDES OVIDIO 1 TYPE 210 • ARDES OVIDIO 2 TYPE 215 • ARDES ORAZIO TYPE 200 • REALCHECK AIRCARE ECONEB-JET • REALCHECK AIRCARE PLUS-JET • FLY (AP10XXU) • DIDACTYS (AP1700U)	IIa
Suction Equipment	• ASPEED Professional • MINIASPEED Battery Plus • MINIASPEED Battery • MAXIASPEED 6.2 • MAXIASPEED 6.2P • MAXIASPEED 6.4 • MAXIASPEED 6.4P • MAXIASPEED 9.2	IIa

	• MAXIASPEED 9.2P • MAXIASPEED 9.4 • MAXIASPEED 9.4P	
Suction System Bottles	• Model ACR-2000 Code: VJ200A (2000 ml device) • Model ACR-4000 Code: VJ400A (4000 ml device)	Im

The complete list of devices is filed with the Notified Body.

Sites covered by this certificate

Via Marziale Cerutti Loc. San Tomaso, 25017 Lonato (BS), Italy

Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the local DNV Office of any intended updating of the quality system and DNV will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system DNV reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of DNV .

END OF CERTIFICATE