EC CERTIFICATE

Full Quality Assurance System

Certificate No.: 240673-2017-CE-ARG-NA-PS Rev.1.0

Project No.: PRJC-186711-2009-MSL-ARG Valid Until: 27 May 2024

This is to certify that the quality system of:

OXIGENOTERAPIA NORTE SACIFIA

CORONEL BOGADO 1954, 1712 CASTELAR, BUENOS AIRES, ARGENTINA

For design, production and final product inspection/testing of:

MEDICAL SUPPLY UNITS

Has been assessed with respect to:

THE CONFORMITY ASSESSMENT PROCEDURE DESCRIBED IN ANNEX II EXCLUDING SECTION 4 OF COUNCIL DIRECTIVE 93/42/EEC ON MEDICAL DEVICES, AS AMENDED

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and date: Høvik, 12 March 2020



PROD 021

For: DNV GL PRESAFE AS Notified Body No.: 2460

Sholeh Gheissar

The certificate is digitally verified by blockchain technology. For more info, see www.dnvgl.com/assurance/certificates-in-the-blockchain.html



Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid. NOTIFIED BODY: DNV GL PRESAFE AS, Veritasveien 3, N-1363 Høvik, Norway - Registered Enterprise No: NO 997 067 401 MVA.

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Jurisdiction

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

Certificate history:

Revision	Description	Issue Date
0.0	Replaces certificate 188893-2015-CE-ARG-NA (NB 0434) following transfer of Notified Body functions to DNV GL Presafe AS (NB 2460).	2018-08-01
1.0	Change of EU Representative Recertification	2020-03-12

Products covered by this Certificate:

Product Description	Product Name	Class
Medical Supply Unit	Bed Head Panels Models: PCM, GCR, GRF	IIa

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address
OXIGENOTERAPIA NORTE SACIFIA	Coronel. J. F. Bogado 1954, B1712DMD, Castelar, Argentina.

EU Representative

Azary Group GmbH.

The Squaire 2, Am Flughafen, 60549. Frankfurt am Main, Germany.

+ 49 69 95932-5090



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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 Presafe of any intended updating of the quality system and Presafe 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. Presafe 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 Presafe.

End of Certificate

NOTIFIED BODY: DNV GL PRESAFE AS, Veritasveien 3, N-1363 Høvik, Norway - Registered Enterprise No: NO 997 067 401 MVA .

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