





Production Quality Assurance Certificate

This is to certify that: Semmco LPS Ltd

Goldsworth Park Trading Estate

9 Kestrel Way Woking **GU21 3BA** United Kingdom

Holds Certificate Number: BSI/MED/PC/754954

In respect of:

MED/3.41

Emergency Escape Breathing Devices (EEBD)

(c) Self-contained closed-circuit compressed air breathing apparatus

Product: 802.015.00, Marine 15 EEBD Orange Case, Self-contained closed-circuit, Chemical Oxygen (KO2) EEBD

On the basis that BSI carried out the relevant Conformity to type based on quality assurance of the production process procedure for the equipment identified above which was found to be in compliance with the Marine Equipment Directive (MED) 2014/90/EU as amended by Directive (EU) 2021/1206 and Regulation (EU) 2021/1158, subject to any conditions in the schedule attached hereto. The attached schedule of approval forms part of this certificate.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

First Issued: 2022-02-15 Latest Issue: 2022-02-15 Expiry Date: 2024-08-04

Page: 1 of 3



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This certificate has been issued by and remains the property of BSI Group The Netherlands B.V., John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands and should be returned immediately upon request. To check its validity telephone +31 20 3460780. An electronic certificate can be authenticated online.

Production Quality Assurance Certificate

No. BSI/MED/PC/754954

SCHEDULE OF APPROVAL

Product Designation	Model	Certificate No.	Issue Date	Notified Body
MED/3.41c – Emergency Escape Breathing Devices (EEBD) - Self-contained closed-circuit compressed air breathing apparatus	802.015.00 Marine 15 EEBD Orange Case	BSI/MED/3.41/754953	15/02/2022	2797

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
February 2022	First Issue	2797:21:3495467

Conditions of Certification

- i) This certificate remains valid unless cancelled or revoked, provided the conditions listed below are complied with and the equipment remains satisfactory in service
- ii) This certificate loses its validity if the manufacturer makes any changes or modifications to the approved quality system, which have not been notified to, and agreed with the notified body named on this certificate and/or after lapse of time, withdrawal or revocation of the Type Examination (Module B) Certificate.
- iii) The equipment detailed above is to be manufactured in accordance with Conformity to Type Based on Quality Assurance of the Production Process (Module D) Certificate of the Marine Equipment Directive.
- iv) If the specified standards are amended during the validity of this certificate, the product type are to be reapproved prior to it being supplied to vessels to which the amended standards apply.
- v) Production tests are to be conducted in accordance with the applicable requirements of the UK Marine Regulation and be recorded by the manufacturer in accordance with the approved Conformity to Type Based on Quality Assurance of the Production Process (Module D) Certificate of the Marine Equipment Directive.
- vi) This certificate authorises the manufacturer or his authorised representative established within the Community in conjunction with the EC Type Examination (Module B) Certificate of the equipment listed in the scope to affix the "Mark of Conformity".

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Page: 2 of 3

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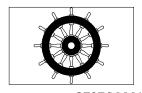
No. BSI/MED/PC/754954

SCHEDULE OF APPROVAL

Conditions of Certification (continued)

vii) Each equipment is to have the "Mark of Conformity" affixed and be issued with a "Declaration of Conformity".

Example for the Application of the "Mark of Conformity":



2797/YYYY

"Wheelmark" Format yyyy Last four digits of year mark affixed. 2797 Notified Body number undertaking surveillance module.

Page: 3 of 3

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