

Production Quality Assurance Certificate

This is to certify that:

Draeger Safety UK Limited
Ullswater Close
Blyth Riverside Business Park
Blyth
NE24 4RG
United Kingdom

Holds Certificate Number:

BSI/MED/PC/768742

In respect of:

Please see scope page.

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, subject to any conditions in the schedule attached hereto. The attached schedule of approval forms part of this certificate.

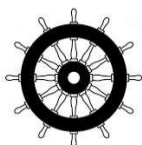
For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

First Issued: 2022-11-18

Latest Issue: 2025-11-17

Expiry Date: 2028-11-17



Production Quality Assurance Certificate

No. BSI/MED/PC/768742

Registered Scope:

MED/3.41

Emergency Escape Breathing Devices (EEBD)

(a) Self-contained open-circuit compressed air breathing apparatus with full mask or mouthed piece assembly for escape

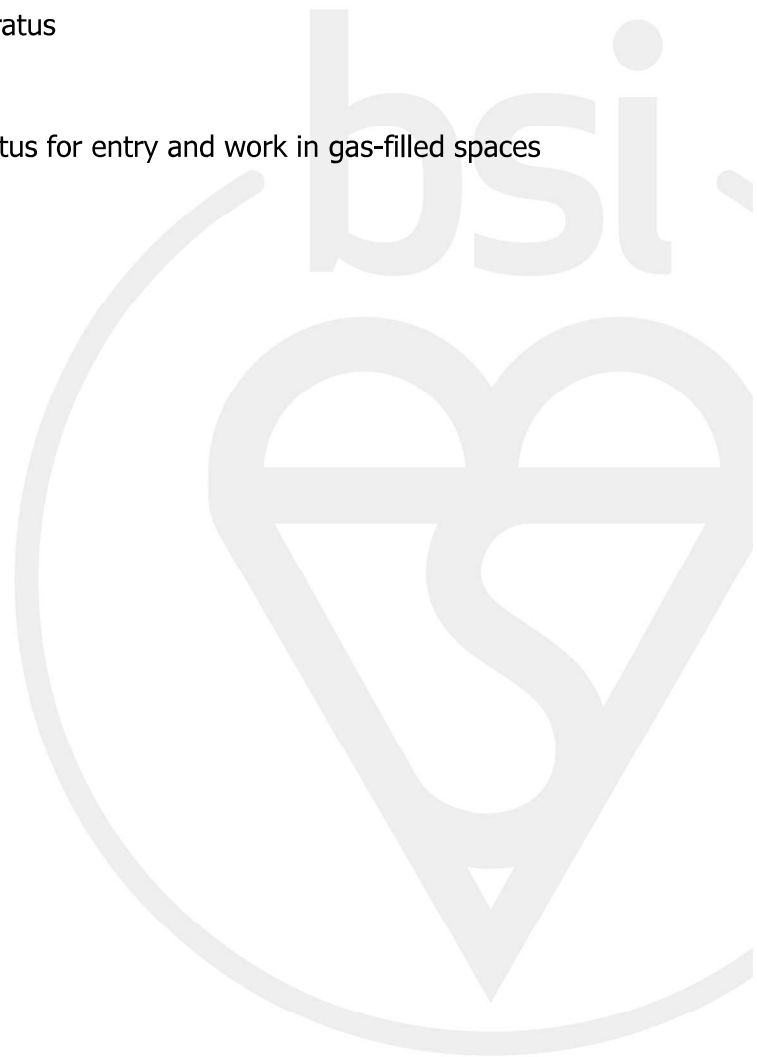
(b) Self-contained open-circuit compressed air breathing apparatus with a hood for escape

MED/3.7

Self-Contained Compressed-Air-Operated Breathing Apparatus

MED/7.1

Self-contained compressed-air-operated breathing apparatus for entry and work in gas-filled spaces



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To check its validity telephone +31 20 3460780. An electronic certificate can be authenticated [online](#).

Production Quality Assurance Certificate

No. BSI/MED/PC/768742

Schedule of Approval

Product Designation	Model	Module B Certificate No.	Issue Date	Notified Body
MED/3.41a	Saver PP	BSI/MED/3.41/768750	02/11/2022	2797
MED/3.41b	Saver CF	BSI/MED/3.41/768755	02/11/2022	2797
MED/3.7	PAS Lite, PSS 3000 & PSS 4000	BSI/MED/3.7/768746	28/10/2022	2797
MED/3.7	PSS 5000	BSI/MED/3.7/768747	02/11/2022	2797
MED/3.7	PSS 7000	BSI/MED/3.7/768748	02/11/2022	2797
MED/3.7	PSS Airboss	BSI/MED/3.7/785330	13/11/2025	2797
MED/7.1	PAS Lite, PSS 3000 & PSS 4000	BSI/MED/7.1/768756	28/10/2022	2797
MED/7.1	PSS 5000	BSI/MED/7.1/768757	02/11/2022	2797
MED/7.1	PSS 7000	BSI/MED/7.1/768758	02/11/2022	2797
MED/3.7	PSS Airboss	BSI/MED/7.1/785331	13/11/2025	2797

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Schedule of Approval

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
November 2022	First Issue	2797:22:3656173
May 2023	Re-issue to add reference of new Module B certificates	2797:23:3850702
July 2024	Re-issue to add updated Module B certificate issue dates	2797:24:30168580
October 2024	Re-issue to add updated Module B certificate issue dates	2797:24:30249530
April 2025	Re-issue to add updated Module B certificate issue dates	2797:25:30377432
November 2025	Re-issue to add updated Module B certificate issue dates	2797:25:30582650

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 re-approved 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.

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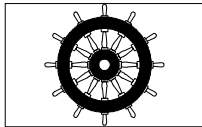
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Schedule of Approval

Conditions of Certification (continued)

- 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".
- 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.

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