Production Quality Assurance Certificate

This is to certify that:

Scott Health & Safety Ltd
Pimbo Road
West Pimbo
Skelmersdale
WN8 9RA
United Kingdom

Holds Certificate Number: BSI/MED/PC/754995

In respect of:

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 space

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):

Denelise L'Ecluse, Managing Director Assurance - Continental Europe

First Issued: 2023-05-03 Latest Issue: 2023-05-03 Expiry Date: 2026-05-02

Production Quality Assurance Certificate

No. BSI/MED/PC/754995

Schedule of Approval

Product Designation	Model	Module B Certificate No.	Issue Date	Approved Body
MED/3.7	Propak SIGMA, Propak I, Propak I DUO, Propak F, Propak F DUO, Propak FX, Propak FX DUO	BSI/MED/3.7/754997	22/03/2023	2797
MED/3.7	Sigma 2 Type 2	BSI/MED/3.7/754998	29/03/2023	2797
MED/7.1	Propak SIGMA, Propak I, Propak I DUO, Propak F, Propak F DUO, Propak FX, Propak FX DUO	BSI/MED/7.1/755000	30/03/2023	2797
MED/7.1	Sigma 2 Type 2	BSI/MED/7.1/754999	30/03/2023	2797

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
May 2023	First Issue	2797:23:3495927

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Schedule of Approval

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.

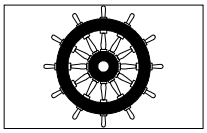
iv) If the specified standards are amended during the validity of this certificate, the product type are to be re-approved prior to it being supplied to vessels to which the amended standards apply.

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.

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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