



Marine & Offshore

Certificate number: SMS.MED2.D/643/E.3

Notified Body N° 2690 www.veristar.com

MED 2014/90/EU QUALITY SYSTEM MODULE D CERTIFICATE

This certificate is issued under the French Maritime Authority, in compliance with the Directive 2014/90/EU of the European Parliament and of the Council of 23 July 2014 as transposed in the French Regulations and Commission Implementing Regulation (EU) 2020/1170 of 16 July 2020, to:

RIKEN KEIKI CO., LTD. Tokyo - JAPAN

Summary of the range of the recognition which is detailed in the subsequent page(s):

FIXED OXYGEN ANALYSIS AND GAS DETECTION EQUIPMENT PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIPMENT

This certificate is issued to attest that Bureau Veritas Marine & Offshore, notified body number 2690, did undertake, at the above company's request, an assessment of the quality system for production quality assurance related to the equipment of the type described in EC type-examination (Module B) certificate(s) listed in the subsequent page(s). Bureau Veritas Marine & Offshore, notified body number 2690, has considered that the quality system operated was satisfying the applicable requirements of the Marine Equipment Directive 2014/90/EU as amended.

This certificate will expire on: 26 Aug 2022

For Bureau Veritas Marine & Offshore, At BV KOBE, on 21 Jan 2021, Shinichi Takemoto



This certificate remains valid until the date stated above, unless cancelled or revoked, provided the conditions indicated in the subsequent page(s) are complied with. This certificate is issued within the scope of the General Conditions of Bureau Veritas Marine & Offshore available on the internet site www.veristar.com. Any Person not a party to the contract pursuant to which this document is delivered may not assert a claim against Bureau Veritas Marine & Offshore for any liability arising out of errors or omissions which may be contained in said document, or for errors of judgement, fault or negligence committed by personnel of the Society or of its Agents in establishment or issuance of this document, and in connection with any activities for which it may provide.

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ANNEX

1. SCOPE

Products

App. number	Sign date	Validity date	Item designation	Trade name	Comment
20988/D1 MED	01 Dec 2020	06 Jul 2021	PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIP- MENT - MED/3.30	GX-2009	
23502/C1 MED	01 Dec 2020	06 Jul 2021	PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIP- MENT - MED/3.30	Portable Multi Gas Monit- or Model: GX-8000	
38435/B1 MED	01 Dec 2020	12 Aug 2023	FIXED OXYGEN ANA- LYSIS AND GAS DE- TECTION EQUIP- MENT - MED/3.54	Smart transmitter / gas detector, Model: SD- 1(TypeGP) and SD-1D	
39093/B1 MED	01 Dec 2020	06 Jul 2021	PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIP- MENT - MED/3.30	Portable gas detector, RX-8000	

App. number	Date of issue	Validity date	Item designation	Trade name	Comment
MEDB00004SX	29 Jan 2020	28 Jan 2025	PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIP- MENT - MED/3.30	RX-8500 & RX-8700	
MEDB00004SY REVISION NO.1	13 Nov 2020	26 Nov 2024	FIXED OXYGEN ANA- LYSIS AND GAS DE- TECTION EQUIP- MENT - MED/3.54	SD-10X/1DOX & SD- 1RI/1DRI & SD-1RI-T	
MEDB000074D	07 Dec 2020	06 Dec 2025	PORTABLE OXYGEN ANALYSIS AND GAS DETECTION EQUIP- MENT - MED/3.30	GX-3R & GX-3R Pro	

2. LIMITATIONS

The EC type-examination certificates listed in the scope are to be valid.

Bureau Veritas Marine & Offshore is to be informed immediately of any modification to the quality system in order to agree on appropriate actions.

RIKEN KEIKI CO., LTD. has to apply for the periodical audits as agreed with Bureau Veritas Marine & Offshore.

RIKEN KEIKI CO., LTD. must draw up a written EC declaration of conformity to type in compliance with the article 16 of the MED 2014/90/EU.

RIKEN KEIKI CO., LTD. must affix the mark of conformity in compliance with the articles 9 and 10 of the MED 2014/90/EU taking into consideration the 2 following examples:



2690/XXXX Where XXXX is the year in which the mark is affixed or



2690/ZZ

Where ZZ are the two last digits of the year in which the mark is affixed.

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3. PERIMETER OF CERTIFICATION

Quality system of following site(s) has been assessed:

RIKEN KEIKI CO., LTD. - Tokyo - JAPAN RIKEN KEIKI CO., LTD. Development Center - Kasukabe - JAPAN

4. REMARKS

The certificate supersedes certificate No.SMS.MED2.D/643/E.2.

*** END OF CERTIFICATE ***