Certificate No: LR21201228ME
Issue Date: 01/06/2021
Expiry Date: 31/05/2024

Notified Body authorised by
BSH 0800-S23/4822:011

EC Module E Certificate of Conformity

This is to certify that:
LLOYD'S REGISTER Deutschland GmbH (LRD), designated as a "notified body" based on the notification of the Federal Maritime and Hydrographic Agency of Germany, did undertake the relevant assessment procedures of the subject manufacturer's Quality System, which was found to be in compliance with the requirements of Marine Equipment Directive (MED) 2014/90/EU (Annex II Module E: Conformity to type based on product quality assurance) for the identified below product(s) type, subject to any conditions listed.

Manufacturer	**Crowcon Detection Instruments Ltd**
Address	172 Brook Drive, Milton Park, Abingdon, Oxfordshire, OX14 4SD, United Kingdom
Reference	Regulation (EU) 2020/1170
Regulation Item (No & item designation)	MED/3.30 Portable Oxygen Analysis And Gas Detection Equipment

Approval is subject to continued maintenance of the requirements of the above mentioned Directive and Regulation and to all products continuing to comply with the standards and conditions of EC Type Examination Certificate(s) issued by Lloyd's Register Deutschland or other Notified Body when they are of the above Designation(s).

Approval is further subject to continued maintenance of the certified quality management system in accordance with the requirements of Lloyd's Register, LRQ 4003384 or an equivalent replacement thereof.

Authorisation is hereby given to the manufacturer to use the LRD Notified Body Registration Number 0525 in accordance with the requirements of the specified Directives in relation to the described product(s).

This certificate remains valid unless cancelled or revoked, provided that product(s) manufactured under this Certificate remain satisfactory in service and the above quality management system continues to be approved.

No product shall be manufactured under this Certificate unless a valid EC Type-Examination Certificate (Module B) is held on that product's Technical File. The manufacturer shall advise the Notified Body of all proposed modifications or changes to a product for which an EC Type Examination Certificate (Module B) has been issued, and of proposed changes of manufacturing location or process, and shall retain copy of their written authorisation or Certification of such changes.

This certificate provides basis for the manufacturer or (if applicable) his authorised representative established within the European Union in conjunction with the EC Type-examination (Module B) certificate of the equipment listed in the scope to affix the "Mark of Conformity" (wheelmark).

0525/(yy)yy

(yy)yy = The year (last two or four digits) in which the mark is affixed.

Lloyd's Register Deutschland GmbH, Überseeallee
10, D-20457 Hamburg, Germany.
A member of the Lloyd's Register group.

Antje Herms-Bondzio

Senior Specialist, Head of MED for LRD
For and on behalf of Lloyd's Register
Deutschland GmbH (0525)

Certificate No: LR21201228ME
Issue Date: 01/06/2021
Expiry Date: 31/05/2024
Reference: BOS251585/LRQ4003384/3996146

Annex to

EC QUALITY SYSTEM (MODULE E) CERTIFICATE No. LR21201228ME

This Certificate is a Renewal of Certificate No. 0525-MED-DE-LR2045364ME

Places of Production				
Crowcon Detection Instruments Ltd 172 Brook Drive Milton Park Abingdon Oxfordshire, OX14 4SD United Kingdom (UK)				
Module B Certificate No.	Regulation Item No./ Description	Module B Certificate Date of Issue	Module B Certificate Date of Expiry	Notified Body
0525-MED-DE-LR2011423MB-02	MED/3.30	31/12/2020	09/06/2025	0525
0525-MED-DE-LR2011430MB-02	MED/3.30	31/12/2020	09/06/2025	0525
0525-MED-DE-MED 1600001-M4-03	MED/3.30	31/12/2020	09/06/2025	0525
0525-MED-DE-MED 1600002-M3-03	MED/3.30	31/12/2020	09/06/2025	0525