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Certificate No: MED 1850111-M3-05

Issue Date: 28/03/2020 Expiry Date: 31/12/2020

EC Module E Certificate Of Conformity

This is to certify that:

LLOYD'S REGISTER VERIFICATION LIMITED (LRV), designated as a "notified body" under the terms of the Merchant Shipping (Marine Equipment) Regulations 2016 (S.I. 2016 No. 1025), as amended, did undertake the relevant assessment procedures of the subject manufacturer's Quality System, which was found to be in compliance with the requirements of the 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) 2019/1397

Regulation Item MED/3.30 Portable Oxygen Analysis And Gas Detection Equipment

(No & item designation)

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 Verification or other Notified Body when they are of the above Designation(s).

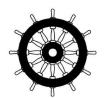
Approval is further subject to continued maintenance of the certified quality management system in accordance with the requirements of LRQA, Certificate Number LRQ 4003384 or an equivalent replacement thereof.

Authorisation is hereby given to the manufacturer to use the LRV Notified Body Registration Number 0038 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.

This certificate is issued under the authority of the MCA. 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)



0038/(yy)yy

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

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Jan de Boer

MED Project Manager CNEEA to Lloyd's Register Verification Ltd A member of the Lloyd's Register group

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EC QUALITY SYSTEM (MODULE E) CERTIFICATE No. MED 1850111-M3-05

This Certificate is an Amendment of Certificate No. MED 1850111/M3

Places of Production

Crowcon Detection Instruments Ltd, 172 Brook Drive, Milton Park, Abingdon, Oxfordshire, OX14 4SD, United Kingdom (UK)

Module B No's	Regulation Item number	Module B Date of issue	Module B Date of Expiry	Notified body
MED 1500009/M5	MED/3.30	02/09/2019	09/06/2020	0038
MED 1500019/M3	MED/3.30	02/09/2019	04/11/2020	0038
MED 1600001/M4	MED/3.30	02/09/2019	05/01/2021	0038
MED 1600002/M3	MED/3.30	02/09/2019	05/01/2021	0038

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