

EC CERTIFICATE Full Quality Assurance System

Certificate No.:

Project No.: PRJN-173524-2019-PA-CZE Valid Until 27 May 2024

10000321467-PA-NA-CZE Rev. 2.0

This is to certify that the quality system of:

Gas Control Equipment Ltd.

100 Empress Park, Penny Lane, Haydock, St Helens WA11 9DB United Kingdom

For design, production and final product inspection/testing of:

PORTABLE OXYGEN CONCENTRATORS

Has been assessed with respect to:

The conformity assessment procedure described in Annex II excluding section 4 of Council Directive 93/42/EEC on Medical Devices, as amended

and found to comply

Further details of the product(s) and conditions for certification are given overleaf.

Place and date: Høvik, 5 March 2021 For the issuing office:
Notified Body 2460



Eugenie Winger Husebye Technical Reviewer



Certificate No.: 10000321467-PA-NA-CZE Rev. 2.0

Place and date: Høvik, 5 March 2021

Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
0.0	Original Certificate	2020-06-10
1.0	Editorial changes - Correction of products names	2020-07-02
2.0	Added new sites (in bold)	2021-03-05

Products covered by this Certificate:

Product Description	Product Name	Class
Portable Oxygen Concentrators	Zen-O™ (RS-00500) Zen-O lite™ (RS-00600) Zen-O™ with Clarity (RS-00500C) Zen-O lite™ with Clarity (RS-00600C)	lla

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address	
Gas Control Equipment Ltd.	100 Empress Park, Penny Lane, Haydock, St Helens WA11 9DB, United Kingdom	
GCE, s.r.o.	Žižkova 381, 583 01 Chotěboř, Czech Republic	
Victor Equipment de Mexico S.A. de C.V.	Jesus Siqueiros 652 Colonia Alvaro Obregón 83170, Hermosillo, Sonora, Mexico	



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Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a
 defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning
 liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the Notified Body of any intended updating of the quality system and the Notified Body will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies
 the quality system. the Notified Body reserves the right, on a spot basis or based on
 suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number.

End of Certificate