EC CERTIFICATE

FULL QUALITY ASSURANCE SYSTEM CERTIFICATE

Cert no. C001254

issued to

Mirola Rescue AB

Dalagatan 5, 11123 STOCKHOLM, SWEDEN

We hereby certify that the Quality System of Mirola Rescue AB for design, production, final inspection and marketing of

Mobile oxygen devices

medical devices in class IIa has been assessed with respect to the conformity assessment procedure according to Annex II (with the exemption of section 4) of Council Directive 93/42/EEC on Medical Devices, as latest amended by Council Directive 2007/47/EC is implemented in Swedish Law by the national regulation LVFS 2003:11, and found to comply with the requirements

This certificate applies to activities performed at

Dalagatan 5, 111 23 Stockholm, Sweden

Originally issued	2020-03-26
Decision date	2020-03-26
Expiry date	2024-05-26

Issued by Notified body 0402

Signed by: Helén Dahl
I am the author of this document
2020-03-26 16:31:36 +01:00

Helén Dahl

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Ackred. no 1002
Certification of Management Systems
ISO/IEC 17021-1

Conditions

Validity

The certificate will remain valid until the expiry date, and allows the holder to use RISE notified body identification number 0402 in conjunction with the CE-mark, on products covered by this certificate, provided that the conditions stated below are fulfilled:

- that surveillance audits are performed, with approved result;
- that the company notifies RISE on all modifications on the products, and that the company does not apply the CE-mark to any new or modified products without confirmation from RISE;
- that the company notifies RISE on all significant changes in the quality system, in its activities and/or organization
- that the certificate is not used in a misleading manner, e.g. in marketing activities.
- that the company notifies RISE about vigilance actions, if any.

Basis for certificate

- The documentation presented has been examined and assessed by RISE in accordance with LVFS 2003:11, Annex II.
- An initial audit and follow-up audits of the quality system at the company's premises in Stockholm has been performed by RISE.
- RISE file 88395

Surveillance

RISE will perform surveillance inspections to ensure that the company maintains and applies the quality system that is subject of the certificate.

In accordance with the EU Commission recommendations of 2013-09-24, there will also be unannounced audits once per every three years. These audits can be performed at the manufacturers as well as at selected crucial supplier's premises.

Miscellaneous

Additional conditions appear in "RISE General Terms – Assignment" and "Rules and process assessment of medical devices as notified body LVFS 2003:11".

Certificate history

Issue	Date	Activity
1	26th March 2020	Original certificate

Register of products covered by the certificate

Product	Art.no.	Class
FIDO	**100**	**IIa**

Note: New products in **bold**