



Quality Assessment Certificate

Notified Body 2843

This is to certify that ABS ITALY S.R.L. did undertake the relevant type approval procedures for the type of equipment identified below which was found to be in compliance with the requirements of Marine Equipment Directive (MED) 2014/90/EU, subject to any conditions in the schedule attached hereto.

CERTIFICATE NUMBER: 07-280652-8-MED

MANUFACTURER: Ocenco Incorporated

MANUFACTURER PLANT LOCATION: Pleasant Prairie, Wisconsin, USA

AUTHORISED REPRESENTATIVE: Interspiro AB

EC TYPE EXAMINATION CERTIFICATE,

NUMBER: see page 3

DATED: see page 3

THIS CERTIFICATE IS ISSUED IN COMPLIANCE WITH CONFORMITY ASSESSMENT **MODULE D** OF THE REGULATIONS AND DIRECTIVES LISTED ABOVE.

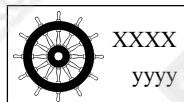
ISSUE DATE: 06-feb-2025

EXPIRATION DATE: 05-feb-2030

ELECTRONICALLY SIGNED BY: E. Brina

This certificate authorizes the manufacturer or his authorised representatives, in conjunction with the EC Type Examination (**Module B**) Certificates listed, to affix the "Mark of Conformity" in accordance with articles 9 & 10 of the Directive.

Example for the application of the "Mark of Conformity":



2843 Number of the Notified Body responsible for the quality surveillance module.

XXXX The year in which the mark is affixed.

The manufacturer shall issue a Declaration of Conformity for each product with reference to the EC Type Examination Certificate and the Quality Assessment Certificate.

Entry Date:

06-feb-2025

**Name of Equipment
Manufacturer:**

Ocenco Incorporated
10225 82nd Avenue
Pleasant Prairie
Wisconsin, WI
53158 USA

Tel: +1-262 947-9000**E-mail:** mikekay@ocenco.com**Website:** www.ocenco.com**Authorized Representative:**

Interspiro AB,
Kemistvagen 12 18379 Taby
Box 2853 18728 Taby
Sweden
Tel. +46 8 636 5100

Equipment/Component:

Emergency Escape Breathing Device (EEBD)

Model:

M-20.2

Scope:

**European Union Marine Equipment Directive 2014/90/EU,
Commission implementing Regulation (EU) 2024/1975:**
Item MED/3.41 - Emergency Escape Breathing Device (EEBD)

Comments:

Quality system approved in accordance with the requirements
of the European Union Marine Equipment Directive
2014/90/EU for Item MED/3.41.

Limitations:

This certificate authorizes the manufacturer or his authorized
representative, in conjunction with valid EC Type Examination
(Module B) Certificates detailed on page 3 to affix the "Mark of
Conformity" in accordance with Articles 9 and 10 of the
Directive.

This certificate loses its validity if the manufacturer makes any
unapproved changes to the approved quality system.

The manufacturer shall issue a Declaration of Conformity for
each product with reference to the EC Type Examination
Certificate and this Quality Assessment Certificate in
accordance with Article 16 of the Directive.

Markings & Declaration of Conformity:

2843/YYYY

Example for the application of the "Mark of Conformity":**2843** Number of the Notified Body responsible for the quality surveillance module.**YYYY** The year in which the mark is affixed.

The following products are covered by this Quality Assessment Certificate:

MED item No.	MED Certificate No.	Expiry date
MED/3.41	07-280652-5-EC	26-oct-2025