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: 07-280652-9-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

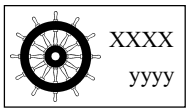
THIS CERTIFICATE IS ISSUED IN COMPLIANCE WITH CONFORMITY ASSESSMENT **MODULE D** OF THE REGULATIONS AND DIRECTIVES LISTED ABOVE.

ISSUE DATE: 24-oct-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”:



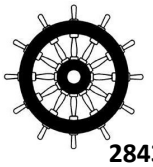
2843 Number of the Notified Body responsible for the quality surveillance module.

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:	24-oct-2025
Name of Equipment Manufacturer:	Ocenco Incorporated 10225 82nd Avenue Pleasant Prairie Wisconsin, WI 53158 USA **Tel:** +1-262 947-9000 **E-mail:** daniel.brtis@ocenco.com **Website:** www.ocenco.com
Authorized Representative:	Interspiro AB, Kemistvagen 21 18379 Taby Box 2853 18728 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) 2025/1533:** 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/3.41c.
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.
Revisions:	**Certificate Nº.** 07-280652-9-MED**, replaces Certificate Nº.** 07-280652-8-MED **Dated:** 06-feb-2025

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/3.41c	07-280652-6-EC	23-oct-2030