

EC-TYPE EXAMINATION CERTIFICATE (MODULE B)

Certificate No: MEDB00000V1 Revision No: 3

Application of: Directive 2014/90/EU of 23 July 2014 on marine equipment (MED), issued as "Forskrift om Skipsutstyr" by the Norwegian Maritime Authority. This Certificate is issued by DNV AS under the authority of the Government of Norway.

This is to certify:

That the Marine evacuation systems

with type designation(s) LSA 9 m, 11.5 m, 14 m & 17 m - Mk1 and Mk2

Issued to

Liferaft Systems Australia Pty Ltd. Derwent Park, TAS, Australia

is found to comply with the requirements in the following Regulations/Standards: Regulation (EU) 2020/1170, item No. MED/1.27. SOLAS 74 as amended, Reg. III/4, III/15, III/26, III/34 & X/3, LSA Code and 2000 HSC Code

8.

Further details of the equipment and conditions for certification are given overleaf.

This Certificate is valid until 2026-05-17.

Issued at Høvik on 2021-05-18

DNV local station: Australia NB

Approval Engineer: Tessa Biever



Notified Body No.: 0575



for DNV AS

Digitally Signed By: Øyvind Hoff Location: DNV Høvik, Norway on behalf of

Sverre Olav Bergli Head of Notified Body

Item no. MED/1.27 Marine evacuation systems Allocated USCG Approval No. extract from Module D Certificate				
Type designation	EC Type- Examination Certificate No.	Expiry date	Notified Body No.	USCG approval number
LSA 9 m, 11.5 m, 14 m & 17 m - Mk1 and Mk2 ¹	MEDB00000V1 Rev.3	2026-05-17	0575	160.175/EC0575 /MEDB00000V1 Rev.3

A U.S. Coast Guard approval number will be assigned to the equipment when the production module has been completed and will appear on the production module certificate (module D, E or F), as allowed by the "Agreement between the United States of America and the EEA EFTA states on the mutual recognition of Certificates of Conformity for Marine Equipment" signed 17 October 2005, and amended by Decision No 1/2019 dated February 22nd, 2019. The mark of conformity may only be affixed to the above type approved equipment and a Manufacturer's Declaration of Conformity issued when the



production-surveillance module (D, E or F) of Annex B of the MED is fully complied with and controlled by a written inspection agreement with a Notified Body. The product liability rests with the manufacturer or his representative in accordance with Directive 2014/90/EU. This certificate is valid for equipment, which is conform to the approved type. The manufacturer shall inform DNV AS of any changes to the approved

This certificate is valid for equipment, which is conform to the approved type. The manufacturer shall inform DNV AS of any changes to the approved equipment. This certificate remains valid unless suspended, withdrawn, recalled or cancelled. Should the specified regulations or standards be amended during the validity of this certificate, the product is to be re-approved before being placed on

Should the specified regulations or standards be amended during the validity of this certificate, the product is to be re-approved before being placed on board a vessel to which the amended regulations or standards apply.

LEGAL DISCLAIMER: Unless otherwise stated in the applicable contract with the holder of this document, or following from mandatory law, the liability of DNV AS, its parent companies and their subsidiaries as well as their officers, directors and employees ("DNV") arising from or in connection with the services rendered for the purpose of the issuance of this document or reliance thereon, whether in contract or in tort (including negligence), shall be limited to direct losses and under any circumstance be limited to 300,000 USD.



Form code: MED 201.NOR