



## EC TYPE EXAMINATION (MODULE B) CERTIFICATE

No.	01-015102/0111507
-----	-------------------

### THIS IS TO CERTIFY:

That Croatian Register of Shipping did undertake the relevant type approval procedures for the equipment identified below which was found to be in compliance with requirements of Marine Equipment Directive (MED) 2014/90/EU, subject to any conditions in the schedule attached hereto.

### TYPE AND DESCRIPTION OF PRODUCT

**CANOPIED REVERSIBLE LIFERAFT TYPE KHR -100**  
with brand names: **YOULONG, SEA SAFE and HERO**

### NUMBER AND ITEM DESIGNATION (in accordance with Annex of Regulation (EU) 2021/1158)

**MED / 1.15 Canopied reversible liferafts**

### MANUFACTURER:

**SHANGHAI YOULONG RUBBER PRODUCTS Co., Ltd.**

20 Xinchun Rd., Chuansha, Pudong

Shanghai 201205

People's Republic of CHINA

### REGULATIONS AND STANDARDS (in accordance with Annex of Regulation (EU) 2021/1158)

**SOLAS 74 as amended, Reg. III/4, III/26, III/34 and Reg. X/3**

**LSA Code as amended Ch. I and IV**

**IMO Res. MSC.81(70), as amended**

**IMO MSC/Circ. 809 and MSC/Circ. 811**

**1994 HSC Code 8 and 2000 HSC Code 8**

### NOTICE:

- Further details of the product and conditions for certification are given overleaf.
- This certificate will not be valid if the manufacturer makes any changes or modifications to the approved equipment, which have not been notified to, and agreed with the notified body named on this certificate.
- Should the specified regulations or standards be amended during the validity of this certificate, the product(s) is/are to be re-approved prior to it/they being placed on board vessels to which the amended regulations or standards apply.
- 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-control phase module (D, E, or F) of Annex II of the Directive is fully complied with and controlled by a written inspection agreement with a notified body.
- In case limitations of use apply, these should be indicated of in the Schedule of Approval.

Issued by Croatian Register of Shipping, notified body number 2489.

This certificate is valid until: **2025-12-20**

Place and date: **Split, 2021-12-22**

Seal

Signature

**Marinko Popović, dipl.ing.**

## THE SCHEDULE OF APPROVAL

### 1. PRODUCT DESCRIPTION

*Canopied reversible liferaft type KHR-100 with brand names YOULONG, SEA SAFE, HERO*

*Max. number of persons: 100*

*Weight: 800 kg with SOLAS A PACK and 575 kg with SOLAS B PACK*

*Required bollard pull:*

*at 2 kn / 2,6 kN*

*at 3 kn / 3,9 kN*

### 2. APPLICATION/LIMITATION OF USE

*Maximum stowage height: 25 m*

*The liferaft and container shall be marked with name of manufacturer, serial number, date of manufacture and all other markings in accordance with LSA Code item 4.2.6.3 and 4.2.7.*

*The liferaft shall be packed with a SOLAS A PACK or SOLAS B PACK.*

*Liferaft is intended to be installed with external weak link and a hydrostatic release unit.*

### 3. DESIGN DRAWINGS AND SPECIFICATIONS

*Design instruction no. YL/QI/091018-DI100, General drawing 502-101/A, Assembly drawing no. 502-106/A,*

*Construction of gas chambers & gas flow direction no. 502/102/A, Floor plan no. 502-103/A and Container dimensions no. 502-104/A.*

### 4. TYPE TEST RECORDS/LABORATORY RECOGNITION STATUS

*Inflatable liferaft material test report No. 2014048*

*Type test report no. 101-201, rev A, dated 20.10.2017.*

### 5. MATERIALS OR COMPONENTS REQUIRED TO BE TYPE APPROVED OR TYPE TESTED

*Hydrostatic release unit shall comply with the requirements of MED 2014/90 EU.*

### 6. OTHER MATERIALS AND/OR COMPONENT

*Gas cylinders shall be of an approved type.*

*Components in the gas inflation system should be approved according to ISO 15738:2002.*

### 7. PRODUCTION SURVEY REQUIREMENTS

*The manufacturer is allowed to affix the Mark of Conformity to equipment referred and to issue a Declaration of Conformity as long as either of the following is fulfilled:*

*Module D: The quality system for production and testing shall be approved by the Notified Body, or*

*Module E: The quality system for inspection and testing shall be approved by the Notified Body, or*

*Module F: Compliance of the products in this EC Type-Examination Certificate is to be verified by the Notified Body, who shall also issue a Certificate of Conformity.*

*Production testing shall be carried out according to IMO Res. MSC 81(70) Part 2.*

### 8. ONBOARD INSTALLATION AND MAINTENANCE REQUIREMENTS

*Installation shall be performed according to manufacturer's instruction.*

*Float free arrangement to be checked upon installation onboard.*

*The liferafts are to be serviced annually at approved service stations which are authorised by the manufacturer.*

### 9. MARKING AND IDENTIFICATION



*Subject to compliance with the conditions in this Schedule of Approval which forms part of certificate, and those of Articles 9, 10 and 15 of the Directive, the Manufacturer is allowed to affix the "Mark of Conformity" to the Product described herein.*

xxxx/yy

*xxxx - the number of the Notified Body undertaking surveillance module (2489 in case of CRS)*

*yy - the last two digits of year mark affixed*

### 10. OTHER

*This approval is given with the understanding that the manufacturer will accept full responsibility for informing shipbuilders, shipowners or their sub-contractors of the proper methods of fitting and general maintenance of the approved equipment and of the conditions of this approval.*

- END OF CERTIFICATE -