

REQUEST FORM FOR MED CERTIFICATION (DIRECTIVE 2014/90/EU)

1. Company Information

- Company name
- Address
- VAT / Tax ID
- Contact person
- Email
- Phone

2. Product Category (Categoria del prodotto)

- | | |
|--|---|
| <input type="checkbox"/> MED/1.1 Lifebuoys | <input type="checkbox"/> MED/1.19 Inflatable rescue boats |
| <input type="checkbox"/> MED/1.4 Lifejackets | <input type="checkbox"/> MED/1.20 Fast rescue boats a), b), c) |
| <input type="checkbox"/> MED/1.12 Inflatable liferafts | <input type="checkbox"/> MED/1.28 Means of rescue |
| <input type="checkbox"/> MED/1.13 Rigid liferafts | <input type="checkbox"/> MED/1.36 Inboard engine for rescue boat propulsion |
| <input type="checkbox"/> MED/1.14 Self-righting liferafts | <input type="checkbox"/> MED/1.37 Outboard propulsion engine |
| <input type="checkbox"/> MED/1.15 Reversible liferafts with canopy | <input type="checkbox"/> MED/1.39 Open reversible liferafts |
| <input type="checkbox"/> MED/1.17 Lifeboats a), b) | <input type="checkbox"/> MED/1.43 Rigid/inflatable rescue boats |
| <input type="checkbox"/> MED/1.18 Rigid rescue boats | |

3. Description of the product to be certified

4. Main dimensions, where applicable"

Length _____ m

Width _____ m

Freeboard _____ m

Weight _____ kg

5. Type of conformity assessment procedure requested

<input type="checkbox"/> Module B (EC-type examination)
<input type="checkbox"/> Module D (Quality assurance of production process)
<input type="checkbox"/> Module E (Product quality assurance)
<input type="checkbox"/> Module F (Product verification)
<input type="checkbox"/> Module G (Unit verification)
<input type="checkbox"/> Other:
<input type="checkbox"/> To be defined with UDICER

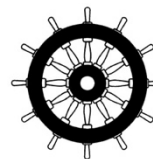
NOTE: "The CE Wheel Mark may be affixed to the device only upon completion of the procedures outlined below. Module B alone is not sufficient.

B + D

B+ E

B+ F

G



6. Additional notes or specific requests

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Please note: Following receipt and evaluation of this request, UDICER will submit a contract for certification activities, including the cost of the intervention, which will be deemed accepted by the applicant upon signing the contract within 15 days of receipt.

"UDICER reserves the right to accept the requested certification activity and will assess whether the certification procedure and the LSA device fall within the scope of Directive 2014/90/EU, or may refuse it for other reasons that will be duly justified in writing."

Date: Signature