



This certificate is issued under the authority given in Merchant Shipping Notice No MSN 1874 (M+F) Annex 2 as amended to date.

Page 1 of 3
Certificate No: LR2431285MC
Issue Date: 18/03/2024
Expiry Date: 17/03/2029

Certificate Of Type Approval (MCA)

This is to certify that LLOYD'S REGISTER, specified as a "person" under the terms of The Merchant Shipping (Marine Equipment) Regulations 2016 (SI2016 No. 1025), did undertake the relevant type approval procedures of the equipment identified below which was found to be in compliance with the essential Life Saving Appliance requirements subject to any conditions in the Design Appraisal Document hereto.

Manufacturer	ANP Pharma Ltd
Address	Unit 9, Honywood Business Park, Honywood Road, Basildon, Essex SS14 3HW, United Kingdom
Type	SURVIVAL CRAFT EQUIPMENT
Description	Category C First Aid Kit for Survival Craft (Lifeboats & Liferrafts) and Rescue Boats
Trade Name	Category C Kit 00585-ANP
Specified Standard	MSN 1905 (M+F) Amendment 3

This certificate is not valid for equipment, the design or manufacture of which has been varied or modified from the specimen tested. The manufacturer should notify Lloyd's Register EMEA of any modification or changes to the equipment in order to obtain a valid Certificate.

The attached Design Appraisal Document forms part of this certificate.

This certificate remains valid up to the expiry date, unless cancelled or revoked, or until such date where it is superseded by the requirements of the Marine Equipment Directive, whichever is sooner, provided the conditions in the attached schedule are complied with and the equipment remains satisfactory in service.

71 Fenchurch Street, London, EC3M 4BS, United Kingdom

Lijo Thomas

Fire & Safety - Senior Surveyor to Lloyd's Register EMEA
A member of the Lloyd's Register group

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ATTACHMENT TO CERTIFICATE OF TYPE APPROVAL No. LR2431285MC

The undernoted documents have been appraised for compliance with the relevant requirements of International Conventions, and this Design Appraisal Document forms part of the Certificate.

This Design Appraisal Document forms part of the Certificate.

APPROVAL DOCUMENTATION

CAT C First Aid Kit Content List, Document No 00585-ANP V1, dated 11/03/2024.

Specification CAT C Medical Kit, Document No 00585-ANP Issue 3, dated 19/03/2024.

PQA, Product Quality Assessment Report no PRJ11100434141, witnessed by Lloyd's Register Surveyor, dated 25/02/2024.

TEST REPORTS

3m Drop test, certificate no UKI2400234/2, witnessed by Lloyd's Register Surveyor, dated 20/02/2024.

CONDITIONS OF CERTIFICATION

1. The First Aid Kit **can** be part of survival craft (Lifeboats & Liferafts) equipment on vessels flying the European and UK flag.
2. The Category C first aid kits for liferafts are to be supplied in sealed units, which should be replaced as a unit if unused at the expiry date.
3. For vessels operating within Category C limits, the machinery on board or the type of operation may give rise to risks which this kit is not adequate to treat. If a risk assessment shows that because of the nature of the work on board, or the type or pattern of operation, the range of stores required in a Category C kit may not be adequate to respond to likely medical requirements on board then the owner or master should consider carrying the additional items marked "RA" (for risk assessment) in Column 4C in Annex 1 of MSN 1905 (M+F). A copy of the risk assessment should be retained on board to provide evidence that the Category of stores and any variations from the recommended quantities of medicines are justified in accordance with Regulations made under the Merchant Shipping Act 1995.
4. The storage arrangement of the first aid kit is not part of this Design Appraisal or Certificate. All such arrangements are to be to the satisfaction of the Surveyor attending on board.
5. If the specified standards are amended during the validity of this certificate, this product type is to be re-approved prior to it being supplied to vessels to which the amended standards apply.



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6. Production items are to be manufactured in accordance with a quality control procedure and records kept as required by MSC.81(70) Part 2, Paragraph 1.2. Production tests are to be conducted to ensure compliance with SOLAS Chapter III Regulation 5. This does not preclude any further testing to additional requirements of the Marine Administration of the country where the ship is registered (i.e. the flag state) or those acting on behalf of that Administration.
7. Should a change of Place of Production from that stated below be required i.e. where the stages of manufacture/assembly/testing of this product take place, the new Place of Production is to be advised to us prior to the change taking place. This Certificate will require to be updated for Approval to be maintained.

PLACE OF PRODUCTION

ANP Pharma Ltd
Unit 9 Honywood Business Park
Honywood Road Basildon
Essex SS14 3HW
United Kingdom

Lijo Thomas
Senior Specialist
Fire & Safety, Statutory Discipline Team
UK&I Technical Support Office, Marine & Offshore
Lloyd's Register EMEA

Supplementary Type Approval Terms and Conditions

This certificate and Design Appraisal Document relates to type approval, it certifies that the prototype(s) of the product(s) referred to herein has/have been found to meet the applicable design criteria for the use specified herein, it does not mean or imply approval for any other use, nor approval of any products designed or manufactured otherwise than in strict conformity with the said prototype(s)