



EC CERTIFICATE

EC TYPE-EXAMINATION (MODULE B) CERTIFICATE No. 2821-MED-0003

This is to certify that: UL International (Netherlands) B.V. did undertake the relevant type approval procedures for the equipment identified below which was found to be in compliance with the relevant requirements of Marine Equipment Directive (MED) 2014/90/EU as amended, subject to any conditions in the schedule attached hereto.

Manufacturer: **Kent Water Sports LLC**
Address: **433 Park Avenue South, New London, Ohio 44851. USA**
Authorised Representative: **SEACOTEC GmbH & Co. KG**
Address: **Millerntorplatz 1, 20359 Hamburg, Germany**
Directive Reference: **MED 2014/90/EU, as amended by Regulation (EU) 2020/1170**
MED Item: **MED/1.6b Immersion suits and anti-exposure suits to be worn without a lifejacket; immersion suit with inherent insulation**
Product Type: **Immersion suit**
Product Description: **Model 1541; Adult Universal, Adult Intermediate, Adult Oversize**
Specified Standard: **Life Saving Appliances Code, 2nd Edition, 2017**

The attached (*schedule of approval*) forms part of this certificate.

This certificate remains valid unless cancelled or revoked, provided the conditions in the attached schedule are complied with and the equipment remains satisfactory in service.

Date of issue: **14 December 2020** Issued by: **UL International (Netherlands) B.V.**
Notified Body 2821

Expiry date: **14 December 2025**

Signed:

This Certificate consists of 2 pages

Name:

Horst Thelen
Head of Notified Body

Notes

1: 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.

2: 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.

3: 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 B of the Directive is fully complied with and controlled by a written inspection agreement with a notified body.

4: In case limitations of use apply, these are indicated in the Annex.



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Schedule of Approval (Annex) No. 2821-MED-0003

Date of issue: 14 December 2020

Issued by: **UL International
(Netherlands) B.V.
Notified Body 2821**

Expiry date: 14 December 2025

Product and Approval documentation:

Technical File 1541 MED Technical Documentation 20201120 version 3
 Type Report MQ6228-D1-European Directive-Original; MQ6228-D1-European Directive-Original-Addendum1; MQ6228-D1-European Directive-Original-Addendum2
 Test Reports ULInternalDataSheet-1; ULInternalDataSheet-2; ULInternalDataSheet-3; ULInternalDataSheet-4; ULInternalDataSheet-5; ULInternalDataSheet-6; ULInternalDataSheet-7; ULInternalDataSheet-8; ULInternalDataSheet-9; ULInternalDataSheet-10; ULInternalDataSheet-11; ULInternalDataSheet-12; ULInternalDataSheet-13; ULInternalDataSheet-14; ULInternalDataSheet-15; ULInternalDataSheet-16; ULInternalDataSheet-17; Test Report – Project 4751132.1101240; RTP Test House Report – Project 4751132.1101240 – Kent

MED/1.6 Immersion suits / anti-exposure suits to be worn without a lifejacket

Regulation SOLAS 74, as amended:
Reg. III/4; Reg. X/3

Regulation of SOLAS 74, as amended, and the relevant resolutions and circulars of the IMO:
Reg. III/7; Reg. III/22; Reg. III/32; Reg. III/34; IMO Res. MSC.36(63)-(1994 HSC Code) 8
IMO Res. MSC.48(66)-(LSA Code) I, II; IMO Res. MSC.97(73)-(2000 HSC Code) 8; IMO MSC/Circ.1046

Testing standard:
IMO Res. MSC.81(70), as amended

Limitations on the acceptance or use of the product(s):

MODEL	SIZE	WEIGHT	HEIGHT
1541	ADULT INTERMEDIATE	110-250 lbs. (50-113 kg)	62-70 in (157-178 cm)
1541	ADULT UNIVERSAL	110-305 lbs. (50-139 kg)	59-75 in (150-191 cm)
1541	ADULT OVERSIZE	250-350 lbs. (114-159 kg)	75-82 in (191-208 cm)

Terms and Conditions:

1. This certificate remains the property of UL International (Netherlands) B.V., herein "UL Netherlands", and will be withdrawn if any conditions attached to its issue are not complied with.
2. This certificate is issued subject to the Global Services Agreement (GSA) and MED Service Terms.
3. Production is limited to the site(s) as listed, or detailed within the Technical documentation held by UL Netherlands.
4. Any product change, production/process changes, or changes in state of the art which may affect conformity shall be notified to UL Netherlands.
5. This certificate does not authorize the use of the Mark of Conformity (the 'Wheel mark'), which may only be affixed to the above type approved equipment and a Manufacturer's Declaration of Conformity issued when Module D, E or F of the Directive is fully complied with and controlled by a written agreement with a notified body.

END OF CERTIFICATE