EC CERTIFICATE

EC TYPE-EXAMINATION (MODULE B) CERTIFICATE
No. 2821-MED-0001

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:	**Datrex Inc.**
Address:	**13878 Hwy 165, Kinder, LA 70648 USA**
Authorised Representative:	**Mr. Lars Lund**
Address:	**Svanbergvägen 9, S-69141 Karlskoga. Sweden**
Directive Reference:	**MED 2014/90/EU, as amended by Regulation (EU) 2020/1170**
MED Item:	**MED1.1 Lifebuoys**
Product Type:	**Lifebuoy**
Product Description**:**	**DX0325D**
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: 28 February 2020
Date of re-issue: 14 December 2020
Expiry date: 28 February 2025

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

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.

UL International (Netherlands) B.V.
Westervoortsedijk 60, 6827AT Arnhem
The Netherlands

EC CERTIFICATE

Schedule of Approval (Annex)
No. 2821-MED-0001

Date of issue:	28 February 2020	Issued by:	**UL International (Netherlands) B.V.**
Date of re-issue:	14 December 2020		**Notified Body 2821**
Expiry date:	28 February 2025		

Product and Approval documentation:

Technical File	DatMedTechDoc, Revision 4, Issued July 13, 2017; Revised Sept. 10 2020.
Type Report	MQ1058-D1-European Directive-Original, as amended
Test Reports	RTP Test House Report – Project 4787937009; RTP Test House Report – Project 4786501819.

MED/1.1 Lifebuoys
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/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

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

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

Maximum stowage height above waterline: 30 Metres.

DX0325D is not intended for use with Quick Release with Light and Smoke Signal and has not been tested as such.

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.

NOTE: This Certificate cancels and replaces certificate 2821-MED-0001 dated 28 February 2020.

END OF CERTIFICATE

UL International (Netherlands) B.V.
Westervoortsedijk 60, 6827AT Arnhem
The Netherlands