EU QUALITY SYSTEM (MODULE D)

Certificate No.
2821-MED-0002/A

Issue date
21-Mar-2025

Issue No.
1

Re-Issue date
N/A

Expiration date
21-Mar-2030

This is to acknowledge that

Datrex Inc.

has had

EU Quality System of manufacturing sites as described in the Annex to this certificate evaluated and meets the requirements of the scope(s):

The manufacture of Lifebuoys

Deputy Head of Notified Body:

Koen Schilleman

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

This is to certify that UL International (Netherlands) B.V. did undertake the relevant conformity to type procedure surveillance activities of the quality system of the manufacturer identified below, which was found to be in compliance with the *Life Saving Appliance* requirements of the Marine Equipment Directive 2014/90/EU, as amended, for the products described in the EC Type Examination Certificates (Module B) as listed in the Annex to this certificate, subject to any conditions attached hereto.

This certificate is only valid in case it is shown including the Annex pages after this front page.

PRODUCTS RvA C 648

UL Solutions

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

Economic Operators:

Manufacturer:	Datrex Inc.
Manufacturers Address	13878 Hwy 165, Kinder, LA, 70648, USA
Authorised Representative:	Mr. Lars Lund
Authorised Representative Address:	Svanbergvägen 9, S-69141 Karlskoga. Sweden
Scope: (Item Designation)	MED/1.1 Lifebuoys

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.

Notes

1: This certificate authorises the manufacturer or his authorised representative established within the Community in conjunction with the EC TYPE EXAMINATION (MODULE B) CERTIFICATE of the equipment listed in the scope to affix the “Mark of Conformity” (‘Wheel Mark’).

2: This certificate loses its validity if the manufacturer makes any changes or modifications to the approved quality system, which have not been notified to, and agreed with the notified body named on this certificate and/or after lapse of time, withdrawal or revocation of the EU TYPE EXAMINATION (MODULE B) CERTIFICATE.

3: Example for the application of the Mark of Conformity. Wheel Mark is followed by either adjacent to or underneath the Notified Body number and the year in which the mark is affixed “xNBx/(yy)yy”. xNBx is the of the Notified Body responsible for the quality surveillance module and (yy)yy is the last two or four digits of the year in which the mark is affixed.

Place of Production:

Name:	Datrex, Inc.
Address:	PO Box 1537, 13878 Hwy, Kinder, LA, 70548. US.

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

Product details:
The following is a non-exhaustive list of PPE products, Technical Files, and/or EU type-examination certificates which were subject of the quality assurance assessment procedure noted below in 'Approval documentation' when the certificate was first established. Module B certificates issued subsequent to this certificate shall be covered by this certificate should they fall within the Scope as detailed on page 1, and are produced at those addresses listed above.

Module B Certificate No.	Technical File Ref.	Product Code(s)
2821-MED-0001/A	DatMEDTechDoc	DX0325D
2821-MED-0006/A	Version Rev. 9	DX0340D

Product and Approval documentation:
Factory Assessment report(s):
UL-CSA-DatrexInc-10-2017
UL-CSA-DatrexInc-10-2018
UL-CSA-DatrexInc-11-2019
UL-JS-DatrexInc-01-2021
UL-HP-Datrex-10.2023-signed
UL-HP-DATREX-20241016-signed

Limitations on the validity of the certificate:

None

Terms and Conditions:

1. This certificate remains the property of UL International (Netherlands) B.V., herein "UL NL", 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 NL.
4. Any system change, production/process changes, or changes in state of the art which may affect conformity shall be notified to UL NL.
5. This certificate authorizes the use of the Mark of Conformity (the 'Wheel Mark'), and should be referenced within the Manufacturer's Declaration of Conformity prior to placement on the market.
6. The certificate remains valid provided the annual audit schedule is maintained and the conclusions obtained attest to continued conformity of the product(s) and quality-control system with the Marine Equipment Directive 2014/90/EU requirements.
7. The requirements for the economic operators stipulated in Article 13 of the Directive 2014/90(EU) of the European Parliament and of the Council of 23 July 2014.

END OF CERTIFICATE

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