$\mathsf{DNV} \cdot \mathsf{GL}$

Certificate No: MEDD00001HJ

QS - CERTIFICATE OF ASSESSMENT - EC (MODULE D)

Application of: Directive 2014/90/EU of 23 July 2014 on marine equipment (MED). This Certificate is issued by DNV GL SE based on the notification of the Federal Maritime and Hydrographic Agency of Germany.

This is to certify:

That the Quality System for the products

with type designation(s) as specified in the Appendix to this Certificate

Issued to

Northrop Grumman LITEF GmbH Freiburg, Germany

is found to comply with the applicable requirements.

The quality system has been assessed with respect to the procedure of conformity assessment described in Annex II, Module D in the directive 2014/90/EU and regulation (EU) 2018/773.

This Certificate is valid until 2023-12-13.

Issued at Hamburg on 2018-12-14

DNV GL local station:

Augsburg

Approval Engineer:

Jörg Rebel

for **DNV GL SE**

Notified Body No.: **0098**

Sven Dudszus Head of Notified Body

The manufacturer is allowed to affix the U.S. Coast Guard approval number(s) as stated in the appendix attached hereto and as allowed by the "Agreement between the European Community and the United States of America on Mutual Recognition of Certificates of Conformity for Marine Equipment", signed February 27th, 2004.



0098/yyyy

0098:

Notified Body number undertaking quality surveillance

The year in which the mark is affixed yyyy:

The product liability rests with the manufacturer or his representative in accordance with Directive 2014/90/FU. This certificate authorizes the manufacturer in conjunction with the valid EC Type Examination (Module B) Certificate(s) of the equipment listed before to affix the Mark of Conformity (wheelmark) to the product described herein.

This certificate loses its validity if the manufacturer makes any changes to the approved quality system, which have not been notified to, and agreed with the notified body named on this certificate. This certificate remains valid unless suspended, withdrawn, recalled or cancelled. The Manufacturer has to apply for periodical audits to verify the maintenance and application of the quality system every 12 months.



Form code: MED 211.DEU Revision: 2016-12 Page 1 of 2