Quality Management System Approval

Number: 2191/4/CD/2026/RST/EN/250051/V01

In accordance with Directive (EU) 2016/797 of 11 May 2016 (as amended)

Assessment according to the Technical Document of ERA 000MRA1044 version 2.0 of December 2022

Object of Assessment	Quality Management System for the production of the Interoperability Constituent on board lifts
Applicant/ Manufacturer	U-Lift AB
Manufacturing Location	Värmansmålavägen 11, SE-372 77, Backaryd, Sweden
Assessment Requirements within the framework of Directive (EU) 2016/797 (as amended)	TSI PRM, (EU) 1300/2014 amended by: Commission Implementing Regulation (EU) 2019/772 of 2019-05-27, Commission Implementing Regulation (EU) 2022/721 of 2022-05-11 Commission Implementing Regulation (EU) 2023/62 of 2023-01-05 Commission Implementing Regulation (EU) 2023/1694 of 2023-08-10 in combination with those Harmonised Standards, Voluntary Standards (or parts thereof), other European or national rules authorized by TSI's and Alternative Solutions as identified is the [Audit Report] (Section 4)
Scope of /Exemptions from Assessment	See [Audit report] (chapter 4)
Module applied	CD of the relevant decision adopted pursuant to the Directive
Assessment / Audit result	The Quality Management System of the aforementioned Manufacturer at the indicated location, has been audited and was shown to comply with the Assessment Requirements, subject to any Conditions and Limits of use as listed below. The Assessment Results are provided in detail within the accompanying [Audit Report] (section 6). The Essential Requirements have been assessed as being met through compliance with the requirements of the relevant TSI only.
The following conditions and limits of use apply	No Conditions or Limits of use
Annex	No
Documentation accompanying this QMS approval	Audit report : DR/26/250051/14, Rev 1.0, 25/01/2026 This documentation is an integral part of this QMS Approval.
Validity	Start: 25/01/2026 End : 24/01/2028 The validity of this QMS Approval is subject to the continued compliance with the Type Examination Certificate as listed in the NoBo Conformity Assessment Report and the continued maintenance of the Quality Management System in accordance with the requirements of the above Directive. This QMS Approval is valid as long as compliance of the Quality Management System with certification requirements is maintained. If certification requirements are affected, then the NoBo must be informed. Within the validity duration of this QMS Approval the applicant can perform production/installation and final product/installation inspection of the object of the assessment. This validity duration may be extended on the basis of future auditing.
Date of Issue / Done on	25/01/2026

Notified Body
DEKRA Rail b.v
ID No. 2191
Concordiastraat 67
3551EM Utrecht
Netherlands

Signature : _____
Armand Verweij, Manager Certification

PRODUCTS
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