EU Declaration of Conformity (DoC)

NOTICE: Sections bracketed with three plus signs (+++) may not be changed or removed without approval from a Quality Director or designee within the Entity and/or function (do not delete the text in this header).

Likorall	**Document Number:** NPD31263; **Version:** 8.0.

Manufacturer Name and Address: Liko AB and Nedre vagen 100, 975 92 Lulea, Sweden

Manufacturer Single Registration Number (SRN): SE-MF-000001404

Authorised Representative Name and Address: Not Applicable, Registered place of business is within European union

Authorised Representative Single Registration Number (SRN): Not Applicable

+++ We as Manufacturer declare, under our sole responsibility, that the product(s) listed below conform to the applicable provisions of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on Medical Devices, and the following Directive(s), Regulation(s) and Common Specification(s). +++

Other relevant Directives, Regulations and Union Legislations that the device is in conformity with:

- The Directive 2011/65/EU (including amendment 2015/863) of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS)

Common Specifications Applied: Not Applicable

Product/Trade Name and Product Code or REF. number: Likorall

Reference Number	Description	Product Basic UDI-DI Number:
3121001	LIKORALL 200	0887761GMN000035U7
3122005	LIKORALL 242 ES, NATURAL	
3122006	LIKORALL 242 ES, WHITE	
3122007	LIKORALL 242 ES R2R, NATURAL	
3122008	LIKORALL 242 ES R2R, WHITE	
3122009	LIKORALL 242 S, NATURAL	
3122010	LIKORALL 242 S, WHITE	
3122011	LIKORALL 242 S R2R, NATURAL	
3122012	LIKORALL 242 S R2R, WHITE	
3122501	LIKORALL 250 ES, NATURAL	
3122502	LIKORALL 250 ES, WHITE	
3123001	LIKORALL 243 ES	
3123002	LIKORALL 243 ES, WHITE	

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3124050	LIKORALL 250 S, NATURAL, IRC	
3122005NA	LIKORALL 242 ES, NATURAL	
3122006NA	LIKORALL 242 ES, WHITE	
3122007NA	LIKORALL 242 ES R2R, NATURAL	
3122008NA	LIKORALL 242 ES R2R, WHITE	
3122009NA	LIKORALL 242 S, NATURAL	
3122010NA	LIKORALL 242 S, WHITE	
3122011NA	LIKORALL 242 S R2R, NATURAL	
3122012NA	LIKORALL 242 S R2R, WHITE	
3122501NA	LIKORALL 250 ES, NATURAL	
3122502NA	LIKORALL 250 ES, WHITE	
3122502CN	LIKORALL 250 ES, WHITE	
3123001NA	LIKORALL 243 ES	
3123002NA	LIKORALL 243 ES, WHITE	
3121660	CARRIAGE ADJUSTABLE 30-50	
3121661	CARRIAGE ADJUSTABLE 50-90	
3121662	CARRIAGE ADJUSTAB. 90-130H	
3126040	ULTRA CONTROL UNIT	
3126045	ULTRATWIST SLIM	
3126047	ULTRATWIST WIDE (N)	
3126034	HANDCONT LR 242ES 2BUTTON	
3126035	HANDCONTROL LR 242 ES-4MS	
3126036	HANDCONTR LR 242 ES 4 BUTTON	
3126038	HANDCONT LR 242 ES 6-BUTTON	
3126060	HANDCONTROL REMOTE	
3126050	HANDCONTROL LR S, ES 2-BUTTON	

Intended Purpose/Use:
Likorall overhead lift is intended for use in lifting and transferring patients, for example, from bed to a wheelchair, to or from the floor, for visits to the toilet, for gait, standing and balance training, when weighing the patient and when lifting the patient with a stretcher.

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Likorall R2R (room to room) overhead lift enables the patient to be moved between two rail systems in separate rooms, without connecting rails and without making holes over doors. Intended for use in following environments: Health care, Intensive care, Emergency ward, Rehabilitation, and Habilitation.
Device Risk Class: Class I
MDR EU Certificate(s) No.: Not Applicable
Conformity Assessment Description/Annexes: Annex II and III
Notified Body Name and Address: Not Applicable as it is class I Product **Notified Body Identification Number:** Not Applicable as it is class I Product

+++ This Declaration is made on the following basis:
• **For devices with a MDR EU Certificate issued by a Notified Body:**
  ○ **The validity of this document shall not start earlier than the validity date of the corresponding MDR EU Certificate.**
  ○ **The DoC declares conformity to all product lots released within the validity period/dates of the corresponding MDR EU Certificate.**
• **For Class I devices (*that are non-sterile, have no measurement function or are not reusable surgical instruments*) the DoC declares conformity to the product lots released after the date of signature.**
• **Compliance to standards and regulations as defined in the Technical Documentation and General Safety and Performance Requirements (GSPR).**
• **Additional information may be attached/appended to this template, such as common specifications, compliance to other union regulations/registrations, product code list or any other supporting information. +++**

Authorised Signatory:	
Name and Title:	Sofie Nybom, Sr. Manager Quality Assurance
Function:	QMR
Place of Issue:	Lulea, Sweden
Date of Issue:	04 APR 2024
Signature: