

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).

Multirall	Document Number: NPD31262; Version 7										
Manufacturer Name and Address: Liko AB and Nedre vagen 100, 975 92 Lulea, Sweden, +46 (0)920 474700											
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: Multirall 200											
<table border="1"><thead><tr><th>Reference Number</th><th>Description</th><th>Product Basic UDI-DI Number:</th></tr></thead><tbody><tr><td>1. 3130001</td><td>1. Multirall 200</td><td rowspan="3">0887761GMN000035U7</td></tr><tr><td>2. 3136001</td><td>2. HANDCONTROL MR</td></tr><tr><td>3. 3136002</td><td>3. HandControl MR 2-button</td></tr></tbody></table>	Reference Number	Description	Product Basic UDI-DI Number:	1. 3130001	1. Multirall 200	0887761GMN000035U7	2. 3136001	2. HANDCONTROL MR	3. 3136002	3. HandControl MR 2-button	
Reference Number	Description	Product Basic UDI-DI Number:									
1. 3130001	1. Multirall 200	0887761GMN000035U7									
2. 3136001	2. HANDCONTROL MR										
3. 3136002	3. HandControl MR 2-button										
Intended Purpose/Use: Multirall 200 overhead lift is intended for use in all common patient lift and transfer situations, for example, between bed/wheelchair, to/ from floor, toilet visits, gait training, and for horizontal lifts with stretchers. Intended for use in following environments: health care, intensive care, and rehabilitation.											
Device Risk Class: Class I											
MDR EU Certificate(s) No.: Not Applicable											
Conformity Assessment Description/Annexes: Annex II and III											

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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:	