

## **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).

|                                                                                                                                                                                                                                                                                                                                              |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------|
| <b>HygieneSlings &amp; HygieneVest</b>                                                                                                                                                                                                                                                                                                       | Document Number: NPD39193; Version: 3.0                                                                                                                                                                                                                                                                                                                                                                                                                          |                                     |
| <b>Manufacturer Name and Address:</b> Liko AB and Nedre vagen 100, 975 92 Lulea, Sweden, +46 (0)920 474700<br><b>Manufacturer Single Registration Number (SRN):</b> SE-MF-000001404                                                                                                                                                          |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>Authorised Representative Name and Address:</b> Not Applicable, Registered place of business is within European union<br><b>Authorised Representative Single Registration Number (SRN):</b> Not Applicable                                                                                                                                |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>+++ 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). +++</b> |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>Other relevant Directives, Regulations and Union Legislations that the device is in conformity with:</b><br>Not Applicable                                                                                                                                                                                                                |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>Common Specifications Applied:</b> Not Applicable                                                                                                                                                                                                                                                                                         |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>Product/Trade Name and Product Code or REF. number:</b> HygieneSlings + & HygieneVest (Mod.41, Mod.45 & Mod.55)                                                                                                                                                                                                                           |                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                     |
| <b>Reference Number</b>                                                                                                                                                                                                                                                                                                                      | <b>Description</b>                                                                                                                                                                                                                                                                                                                                                                                                                                               | <b>Product Basic UDI-DI Number:</b> |
| 1. 3541133<br>2. 3541134<br>3. 3541135<br>4. 3541136<br>5. 3541137<br>6. 3545134<br>7. 3545135<br>8. 3545136<br>9. 3545137<br>10.3555112<br>11.3555113<br>12.3555114<br>13.3555115<br>14.3555116<br>15.3555117                                                                                                                               | 1. HYGIENE SLING 41 PES XS<br>2. HYGIENE SLING, MOD 41, S<br>3. HYGIENE SLING, MOD 41, M<br>4. HYGIENE SLING, MOD 41, L<br>5. HYGIENE SLING, MOD 41, XL<br>6. HYGIENE SLING 45 PES S<br>7. HYGIENE SLING 45 PES M<br>8. HYGIENE SLING 45 PES L<br>9. HYGIENE SLING 45 PES XL<br>10.HYGIENEVEST HB 55 PES<br>XXS<br>11.HYGIENEVEST HB 55 PES XS<br>12.HYGIENEVEST HB 55 PES S<br>13.HYGIENEVEST HB 55 PES M<br>14.HYGIENEVEST HB 55 PES L<br>15.HYGIENEVEST HB 55 | 0887761GMN000038UD                  |

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### **Intended Purpose/Use:**

Liko Hygiene Sling (Mod 41) is designed in order to assist with dressing and undressing when lifting to and from the toilet

Liko Hygiene Sling (Mod 45) has been designed to facilitate dressing and undressing in connection with lifts to and from the lavatory.

Liko Hygiene Vest (Mod. 55) is designed for safe lifting and transfer to and from the toilet.

It is 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:**

|                        |               |
|------------------------|---------------|
| <b>Name and Title:</b> | Sofie Nybom   |
| <b>Function:</b>       | QMR           |
| <b>Place of Issue:</b> | Luleå, Sweden |
| <b>Date of Issue:</b>  | 27-NOV-2023   |

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|                   |                                                                                    |
|-------------------|------------------------------------------------------------------------------------|
| <b>Signature:</b> |  |
|-------------------|------------------------------------------------------------------------------------|

