EU Declaration of Conformity

according to the EU-Medical Device Directive 2017/745

SLK Vertriebs GmbH
Am Herdicksbach 18
D-45731 Waltrop

We

SLK Vertriebs GmbH
Deutschland/Germany

Single registration number (SRN): DE-MF-000010037

Basic UDI: 426064753

hereby declare under our sole responsibility that the products listed in Appendix 1 for the following product group:

- SLK Active Hoists

comply with the conformity assessment procedure set out in Annex IV of Regulation (EU) 2017/745 and all applicable essential safety and performance requirements of Annex I are met.
The procedures set out in Annex II and Annex III of the same Regulation were followed.
The classification of the products listed above is carried out in accordance with Annex VIII of the Regulation.

We assure that the products comply with this regulation and, where applicable, other relevant Union legislation.

This declaration is valid for products placed on the market from the date of issue.

This document is valid until: 31.12.2025

Waltrop, 13.11.2024
Place, Date

Signature: Oliver Otte, Managing Director

Annex 1

SLK Active Hoists

Risk Class: I
Basic UDI-DI: 426064753AL0100001SY
GMDN: 38050

Article-ID	Name of Article	Description
8008	SLK Multy Aktiv	Active Hoist, max. 185kg
8011	SLK Multy Conversion Kit Aktiv	Conversion Kit from Multy Universal to Multy Aktiv
8012	SLK Multy Aktiv e	Active Hoist, max. 185kg, Electrical Base Expansion
8016	SLK Eazy-up flex	Active Hoist with Base Expansion
8017	SLK Eazy-up fix	Active Hoist with fixed Base

Waltrop, 13.11.2024
Place, Date

Signature: Oliver Otte, Managing Director