EU MDR DECLARATION OF CONFORMITY

REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017

We, Guangzhou Landswick Medical Technologies Limited.
Add: NO. 1302/13F, Block C2. No.182 science avenue, Guangzhou Science City, Guangzhou Hi-tech Industrial Development Zone, Guangzhou, China. Post Code: 510663
Phone: +86 20 28065346 www.landswick.com
SRN number: CN-MF-000022972
Basic UDI-DI: 69742962100369

hereby declares being the manufacturer of the medical devices Vacuum Mattress and Vacuum Splint according to the enclosed article number list, and that the devices are in conformity with Medical Device Regulation (EU) 2017/745 (MDR) and fulfill all relevant requirements in MDR.
The devices are intended for patient transfer and are classified in Class I according to Annex VIII in MDR

Name: Vacuum Mattress
Type or model: VMA-0B01

Name: Vacuum KED Extrication Device
Type or model: VMA-1B01

Name: Vacuum Splint
Type or model: VSA-3A00, VSA-4A00, VSA-5A00, VSA-001, VSA-002, VSA-003 ,VSA-4B00.

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

We hereby explicitly appoint Share Info Consultant Service LLC Repräsentanzbüro located at Heerdter Lohweg 83, 40549 Düsseldorf to act as our European Authorized Representative as defined in the aforementioned Directive.

Date of issue: 05th September 2023
Place of issue: Guangzhou, PR China
Signature:
Typed name: **Dong Hui**
Position/Title: Legal Representative
Stamp/Seal: