INSTRUCTIONS FOR USE – Manual Suction (Manual Vacuum Pump)

Intended Purpose
Manual Suction is a hand-operated medical device intended to generate negative pressure for the temporary removal of fluids and secretions from the airway in emergency and clinical situations.
The device is not intended for continuous or long-term suction.

Intended Users
Healthcare professionals and trained emergency or rescue personnel.

Patient Population
Adults and children, according to applicable clinical protocols.

Device Description
Manual Suction is a lightweight, portable, hand-powered suction device designed for single-hand operation.
The device consists of a reusable vacuum pump handle, disposable 300 ml collection container, and adult/child suction catheters.
Vacuum level is adjustable between 50% and 100% stroke.

Contraindications
Do not use if any component is damaged, missing or contaminated.
Do not use if the pump handle is cracked or broken.

Warnings
• Do not use the device upside down
• Do not autoclave any components
• Do not immerse the pump in liquid
• Use only by trained personnel

Precautions
Ensure all connections are secure before use.
If resistance increases, stop use and check for blockage.

Directions for Use

1. Select appropriate catheter size
2. Connect catheter to collection container
3. Secure container to pump handle
4. Select 50% or 100% stroke
5. Operate trigger to generate vacuum
6. Remove catheter slowly after use
7. Dispose of single-use components according to local protocols

Performance
Vacuum at 20 kPa (1 min): ≥15 kPa
Maximum vacuum: ≥39.9 kPa
Container volume: 300 ml

Technical Data
Operating temp: −20 °C to +50 °C
Storage temp: −40 °C to +60 °C
Net weight: 230 g

Cleaning
Clean reusable parts with warm water and mild detergent.
Do not autoclave.

Storage
Store dry. Protect from damage. Observe temperature limits.

Symbols
Symbols comply with ISO 15223-1. Refer to label for explanation.

UDI
UDI is provided on the device label and packaging.

Manufacturer
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Regulatory compliance
This medical device complies with Regulation (EU) 2017/745.

Document control
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