ENGLISH

DESCRIPTION OF SYMBOLS

MD

Medical Device

INDICATION:

For foot and ankle conditions that may benefit from motion restriction, such as:

- Grade 2 and 3 ankle sprains.
- · Stable fractures.
- · During postoperative or rehab phase.

FEATURES

- High quality rigid outer skeleton for protection.
- Metal upright design allow for bars to be contoured to the patient for a custom fit.
- Dense foam lining within the sole to provide comfort and support.
- Shock absorbing sole to improve comfort and protects the user from slips, providing additional security in each step.
- · Rocker bottom to simulate natural gait.
- Uprights give additional stability by restricting of motion of the leg.
- Large air bladder creates a larger surface for contact pressure to help control swelling and ensure a snug fit.

TO PUT ON THE PNEUMATIC WALKER WITH METAL UPRIGHTS

- Open the boot liner without removing it from the boot
- Place the boot flat on the floor and put your foot inside the boot. Make sure your heel is touching the back of the boot.
- Secure the boot lining to your leg by overlapping the lining on top of your foot and at the front of your lower leg.
- Secure the remaining Velcro straps, beginning with the straps closest to your toes.

TO CONTROL THE AIR PRESSURE INSIDE THE PNEUMATIC WALKER WITH METAL UPRIGHTS.

- To put air inside the boot, turn the valve clockwise until secure, then squeeze the grey bulb until you feel adequate pressure.
- To remove air from the boot, simply turn the valve counterclockwise.

TO TAKE OFF THE PNEUMATIC WALKER WITH METAL UPRIGHTS

- Open the boot liner by detaching all of the Velcro straps.
- Without removing the liner from the base of the boot, take your foot out of the Walker.

WASHING INSTRUCTIONS

Wash the liner with mild soap and water and lay out to dry. DO NOT machine dry.

Size	Men's Shoe Size	Women's Shoe Size
X-Small	<4	<5.5
Small	4.5-7	6-8
Medium	7.5-10.5	8.5-11.5
Large	10.5-12.5	11.5-13.5
X-Large	>12.5	>13.5

GENERAL SAFETY INFORMATION

WARNING: In cases of functional change or functional loss, the patient should stop using the device and contact a healthcare professional.

WARNING: If the device shows signs of damage or wear hindering its normal functions, the patient should stop using the device and contact a healthcare professional.

REPORT A SERIOUS INCIDENT

Important notice to users and/or patients established in Europe:

The user and/or patient must report any serious incident that has occurred in relation to the device to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

DISPOSAL

The device and packaging must be disposed of in accordance with respective local or national environmental regulations.

LIABILITY TEXT

Össur does not assume liability for the following:

- Device not maintained as instructed by the instructions for use.
- Device assembled with components from other manufacturers.
- Device used outside of recommended use condition, application, or environment.