

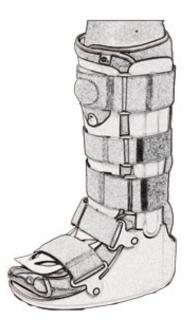
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WALKER CLAMSHELL AIR TALL

Rx Only — Single Patient Use Only



Instructions for Use



For questions, please contact your physician, the professional who fit this device or Allard USA Customer Service:

CALL: 888-678-6548 FAX: 800-289-0809 info@allardusa.com

Allard USA, Inc. 300 Forge Way, Suite 3 Rockaway, NJ 07866-2056 www.allardusa.com

CSUS

by Allard USA

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1. Pull all straps loose from plastic buckles and press each strap back onto itself.



2. Pull front shell away from liner. The top strap should be totally free from the plastic shell.



3. Open liner and step into it. Close flaps snugly starting at the toes. Insure the blue bulb is centered on the front of the leg.



7. Place the front shell centered between the uprights on front of the leg. Thread the blue bulb strap through the large hole on the shell. Insure the black valve is exposed.



8. Close the strapping on the walker, starting at the toes and up the leg. Insure you follow this strapping sequence to achieve maximum support.



9. After fastening all straps securely, inflate liner air cell by pressing the blue bulb on the top front of the boot until the walker feels comfortably secured. Do not over inflate to avoid restricting blood circulation in your foot.



4. Step into walker and place your heel as far back against the walker as possible to insure a comfortable fit.



5. Remove the "Pull Up" strips and press the uprights firmly against both sides of the leg. Insure the uprights are centered over the ankle.



6. Place the back shell on the rear of the leg against the liner surface. Insure the shell is centered between the uprights of the walker.



10. To deflate the air cell, open the small black cover below the blue bulb and press the brass button.

CAUTION

Patients without sensation need to be observed frequently for skin irritation, hot spots, etc.
This device is restricted by Federal Law to be sold only under prescription or by a licensed health-care professional.

WARNING

Overinflating air cells could cause skin irritation. If observed, discontinue the use of the device and contact your healthcare professional. Do not use this device on patients unable to communicate pain or physical discomfort.

