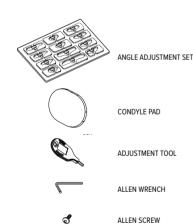


ASPEN OA KNEE+

OSTEOARTHRITIS KNEE BRACE OSTEOARTHRITIS-KNIEORTHESE ÓRTESIS DE RODILLA PARA OSTEOARTRITIS **GINOCCHIERA PER L'OSTEOARTRITE ORTHÈSE POUR ARTHROSE DU GENOU KNIEBRACE TEGEN ARTROSE** KOLENNÍ ORTÉZA PRO OSTEOARTRÓZU KNÆBØJLE TIL SLIDGIGT ΝΑΡΘΗΚΑΣ ΓΟΝΑΤΟΥ ΓΙΑ ΟΣΤΕΟΑΡΘΡΙΤΙΔΑ KNESTØTTE FOR OSTEOARTRITT ORTEZA KOLANOWA DLA PACJENTÓW Z CHOROBA ZWYRODNIENIOWĄ STAWÓW **POLVEN NIVELRIKKO-ORTOOSI**

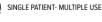
> **KNÄSTÖD FÖR ARTROS** Коленный ортез при остеоартрозе JOELHEIRA PARA OSTEOARTRITE 骨关节炎膝关节支架





MD MEDICAL DEVICE



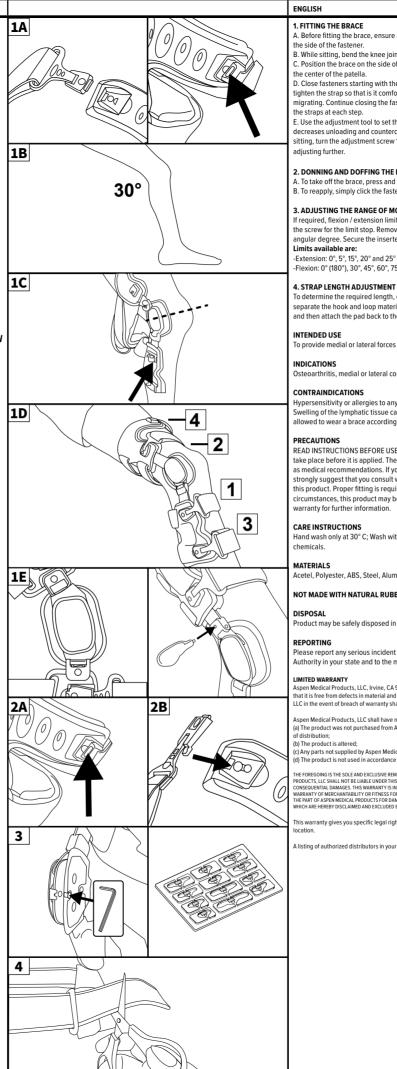






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1. FITTING THE BRACE

A. Before fitting the brace, ensure all brace fasteners are open by pressing the button on the side of the fastener

B. While sitting, bend the knee joint slightly (approximately 30°)

C. Position the brace on the side of the leg so that the center of the hinge lines up with the center of the patella.

D. Close fasteners starting with the strap directly below the knee labeled 1. Adjust and tighten the strap so that is it comfortable for the patient and prevents the brace from migrating. Continue closing the fasteners following the order sequence 2, 3, 4 adjusting the straps at each step.

E. Use the adjustment tool to set the appropriate degree of unloading. Clockwise decreases unloading and counterclockwise increases unloading. While the patient is sitting, turn the adjustment screw 1/8 to a 1/4 turn at a time and evaluate patient before adjusting further.

2. DONNING AND DOFFING THE BRACE

A. To take off the brace, press and release the fasteners until all straps are released. B. To reapply, simply click the fasteners in place.

3. ADJUSTING THE RANGE OF MOTION

If required, flexion / extension limits can be set. Remove the condule pad and remove the screw for the limit stop. Remove the limit stop from the hinge and insert the required angular degree. Secure the inserted limit stop with the screw

Limits available are:

-Extension: 0°, 5°, 15°, 20° and 25°

-Flexion: 0° (180°), 30°, 45°, 60°, 75° and 90°

To determine the required length, open the strap pad, open the end of the binding strap, separate the hook and loop material and the binding strap, cut to the required length, and then attach the pad back to the end of the binding strap.

INTENDED USE

To provide medial or lateral forces on the knee joint.

INDICATIONS

Osteoarthritis, medial or lateral compartment of the knee

CONTRAINDICATIONS

Hypersensitivity or allergies to any of the materials from which the brace is made. Swelling of the lymphatic tissue caused by circulatory disorders. Patients who are not allowed to wear a brace according to medical instructions.

PRECAUTIONS

READ INSTRUCTIONS BEFORE USE. Proper training in the use of this device should take place before it is applied. These directions are guidelines only and are not offered as medical recommendations. If you suffer from a serious medical condition, we strongly suggest that you consult with a licensed health care professional before using this product. Proper fitting is required for this product to be effective. Under some circumstances, this product may be prescribed by a physician. Please see the limited

CARE INSTRUCTIONS

Hand wash only at 30° C; Wash with mild detergent; Air dry; Do not use bleach or other chemicals.

MATERIALS

Acetel, Polyester, ABS, Steel, Aluminum, UBL, EVA, Nylon, Silicone, Polyurethane.

NOT MADE WITH NATURAL RUBBER LATEX.

DISPOSAL

Product may be safely disposed in accordance with local laws

Please report any serious incident involving the use of this device to both the Competent Authority in your state and to the manufacturer (listed on this document).

Aspen Medical Products, LLC, Irvine, CA 92618, warrants to the user who originally purchases this product that it is free from defects in material and workmanship. The sole obligation of Aspen Medical Products, LLC in the event of breach of warranty shall be to repair or replace the defective product or part(s).

Aspen Medical Products, LLC shall have no obligation under this limited warranty in the event:
(a) The product was not purchased from Aspen Medical Products, LLC or through its authorized channels of distribution;

(b) The product is altered

(c) Any parts not supplied by Aspen Medical Products. LLC are inserted into the product; or (d) The product is not used in accordance with the Aspen Medical Products, LLC Instructions for Use.

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A listing of authorized distributors in your area is available upon request.