08668
Swedish Knee Cage
Instructions and User Guide
40008-001 REV E 2022-05-23

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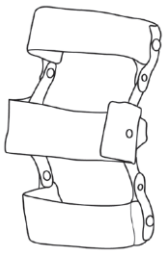

INTRODUCTION

The Swedish Knee Cage is a non-articulating knee brace. Its 3-point pressure system provides a simply yet effective way to control genu recurvatum.

INDICATIONS

Knee hyperextension, genu recurvatum.

Part Number	Description	Size	Circumference *
0866802	Lenox Hill Swedish Knee Cage	Small	25 - 30 cm • 10 -12 in
0866803	Lenox Hill Swedish Knee Cage	Medium	33 - 35 cm • 13 - 14 in
0866804	Lenox Hill Swedish Knee Cage	Large	38 - 41 cm • 15 -16 in
0866805	Lenox Hill Swedish Knee Cage	X-Large	42 - 46 cm • 17 -18 in
			***10 cm (4”) above knee center**

Accessories	Description	Size
0266802	Replacement Straps F/8668-Swedish Knee SM	Small
0266803	Replacement Straps F/8668-Swedish Knee MD	Medium
0266804	Replacement Straps F/8668-Swedish Knee LG	Large
0266805	Replacement Straps F/8668-Swedish Knee XL	X-Large

⚠ The snap rivets are nickel plated. Take appropriate action if the patient is allergic to nickel.

INSTALLATION AND USE

1. Unfasten one side of the proximal and distal straps at the snaps and slide off the uprights.
2. The patient should be seated with the knee in flexion.
3. Apply the brace from the posterior, centering the middle back pad to the knee.
4. Adjust the M/L of the brace to the knee for comfort.
5. Bring the proximal and distal straps across the front of the leg and reattach onto the uprights. The elastic portion of the straps should be against the leg.
6. Stand the patient and observe the degree of knee extension.
7. Adjust if necessary.

Note: The attachment position of the back pad controls the amount of extension lock-out i.e. attachment more anteriorly provides less lock-out (more knee extension).

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STORAGE AND USE
There are no identified restrictions on the temperature of storage and use.

DISPOSAL
There are no hazardous materials in the device. Comply with local and national laws and regulations.

LEGAL INFORMATION
MD The use of this class 1 medical device is subject to the respective national laws of the country of use and may vary accordingly. The user of this device should report any serious incident to Trulife and the competent authority of the country of use.

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LIMITED WARRANTY
Trulife warrants that the PRODUCT will be free from defects in material and workmanship from the date of installation for the warranty period stated on the PRODUCT warranty card.

This warranty will not apply if the PRODUCT has been damaged by misuse, abuse, neglect, improper care, failure to follow instructions, abnormal wear and tear, or in the event that the PRODUCT has been modified/repaired by persons unauthorized by Trulife.

If a defect in material or workmanship is found during the warranty period, Trulife will, at Trulife's option, either repair or replace the product. If it is not possible to repair or replace the product, Trulife will be limited to refunding the purchase price.

Trulife will not be liable under any legal theory for any direct, indirect, special, incidental or consequential damages arising from the use of or inability to use this product.

The application guidelines for this Trulife product are for the use of and by a certified, qualified practitioner only. Patients are not to attempt to apply or adjust the item unless expressly instructed to do so by the practitioner responsible for the prescription and/or initial fitting of the device. All patient questions should be referred to the practitioner and not to the manufacturer. The manufacturer warrants only that the enclosed product has been inspected for quality and can be effective for certain indications, but final decisions and ongoing monitoring must be made by the medical professional(s) prescribing and/or fitting the device to determine its effectiveness for an individual patient. Patient compliance is an integral part of the entire protocol and must be adhered to in order to avoid potential problems and to maximize the effectiveness of the prescribed product.

As a condition of the sale of any Trulife product, this product is restricted to a "Single Patient Use Only" by the originally fitted patient in order to protect the care provider and the patient against potentially adverse consequences of infectious disease transmission, material instability in adapting to the configuration of the original user and/or decrease in effectivity. Any express or implied warranties are voided if the product is reused or fitted to another patient. Additionally, a license of right to use under any relevant patents pertaining to the product is terminated with the cessation of use by the original patient. As with all Trulife products, this product must be prescribed and applied by a qualified practitioner to determine it meets the needs of the particular patient and accomplishes the desired results.

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