



INTRODUCTION

The adult Foot Spacer provides versatility during setup and fabrication of the LightFoot and Energy foot. It may be used either temporarily to permit easier access to alignment devices or permanently to allow setup on existing standard limb systems and to accommodate patient growth. The spacer may also be used to adapt an existing exoskeletal limb to the LightFoot or Energy foot for trial purposes prior to new fabrication with an ankle block.

SELECTION

Spacer for use with adult LightFoot and Energy (M10 Foot Bolt):

Model No.	Description	Height	Product Wt	Weight Limit
SFS232	Spacer Adult Feet	2.5 cm / 1.0 in	57 g / 2.0 oz	136 kg / 300 lb

INSTALLATION AND USE

Recommended installation and use procedures must be followed for maximum safety and service life.

Warning: Never modify the spacer. It will void the warranty and can cause failure. Never redrill the mounting holes.

- When using wood or foam ankle blocks, the distal end must be reinforced with two layers of laminate.
- Failure to reinforce the ankle block can result in bolt failure.
- The foot mounting bolt must fully engage the connecting ankle block or foot adaptor threads.
- You may stack up to two spacers.

Warning: Do not stack more than two spacers in the definite setup.

Warning: Failure to follow the installation and use procedures set forth above may lead to structural failure of the components subjecting the user to a risk of serious personal injury.





QUESTIONS

Contact Customer Service at;

USA

Tel: +1 800 492 1088

Fax: +1 800 245 3765

Email: info-usa@trulife.com

Visit Trulife online at www.trulife.com.

Canada

Tel: +1 800 267 2812

Fax: +1 613 392 4139

Email: infocanada@trulife.com

Trulife has appointed Medical Device Safety Service (MDSS) of Hannover, Germany to act as our EU authorized representative. They may be contacted at:

MDSS GmbH

Schiffgraben 41

30175 Hannover

Germany

Phone (+49)-511-6262 8630

FAX (+49) -511-6262 8633

LIMITED WARRANTY

Trulife warrants that the PRODUCT will be free from defects in material and workmanship for the PRODUCT from the date of installation.

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 a PRODUCT has been modified/repared 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 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.

