SAB310, SAB314, SAB315, SAB316, SAB320
Seattle Ankle Block
Installation and User Guide
15094-001 REV J

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INTRODUCTION

The Seattle Ankle Block is designed for the exoskeletal setup of various Seattle feet. Manufactured using high strength polyurethane foam surrounding a nylon core, the Seattle Ankle Block is lightweight and durable to withstand the forces imposed by dynamic response feet.

SELECTION

Product Code	Appropriate Foot Style	Foot Size Range	Max. Patient Weight	Max. Clearance (incl. foot)	Bolt Thread
Adult					
SAB310	SLF195/198	22-30 cm	300 lbs (136 kg)	6.1 in (15.5 cm)	M10
SAB320	SNF160/163 SNF170/173 SCH110/113 SCH120/123	22-30 cm	325 lbs (102 kg)	6.1 in (15.5 cm)	M10
Child's Play Energy					
SAB314	SEF132/144	13-15 cm	77 lbs (35 kg)	3.7 in (9.4 cm)	M8
SAB315	SEF132/144	16-18 cm	99 lbs (45 kg)	4.1 in (10.4 cm)	M8
SAB316	SEF132/144	19-21 cm	144 lbs (65 kg)	4.4 in (11.2 cm)	M8

FABRICATION

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

Warning: Never modify the exposed portion of the nylon core except to grind off the rotation control rib when necessary. Other modifications will void the warranty and can cause failure. Sanding or grinding of the foam surrounding the core is permitted only on the tops and sides, and only to a depth of 1/16" (2 mm).

Rotation control can be accomplished by either of two methods. (1) If rotational alignment is sufficient, make sure the central control rib engages the keel top as the foot mounting bolt is tightened. (2) If additional adjustment is required, grind down the rib. Mold a rib on the ankle block by placing a small amount of epoxy resin in the shallow recess. You may add fillers to the resin to make a paste.

Never cut down the embedded portion of the core. If you need a shorter connection, use the Seattle Laminating Core (SLC301, 302, 303) for Child's Play Energy Feet only.

Warning: Testing an unreinforced ankle block after shaping can cause failure.

The circular recess on the bottom of the ankle block must be placed posterior to the keel centerline as close as possible.

Trulife

QUESTIONS

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Visit Trulife online at www.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
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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/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 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.