



Instructions for Use

REF SchureFoot, Lexan 800-0011

Replacement Pads

SchureFoot Disposable Pads 508-0111

INTENDED USE

The SchureFoot, Lexan is specifically designed to leverage the knee, allowing access to the medial compartment of the operative leg during arthroscopic procedures. The intended users of this device are medical professionals within hospitals and surgery centers.

INSTRUCTIONS

Become familiar with the features of patient positioning device before use with a patient. Always practice using it on a nurse, physician or appropriate volunteer prior to using clinically.



Set Up and Use of SchureFoot

Attach socket to surgical table side rail near end of torso section, on side of operative knee

Slide disposable pad #508-0111 (sold separately) onto SchureFoot tube

Insert SchureFoot mounting post into socket

Once patient is on surgical table, slide SchureFoot along rail to achieve desired position for leg

Turn socket handle clockwise securing SchureFoot in desired position

Prep and drape in usual method

After completing procedure, discard the SchureFoot disposable pad. Spray SchureFoot with hospital-approved disinfectant and wipe clean

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- *Length: 12" +/- 0.5" (31 cm +/- 1 cm)*
- *Height: 16" +/- 0.5" (41 cm +/- 1 cm)*
- *Diameter: 2" +/- 0.5" (5 cm +/- 1 cm) (with pad 2 3/4" (7 cm))*
- *Device Weight: 2 +/- 0.5 lbs. (.9 +/- .22 kg)*
- *Operates on leg section of surgical table*
- *Requires a socket to attach to surgical table*

COMPONENT OVERVIEW

SchureFoot, Lexan is used in a variety of surgical procedures including , but not limited to arthroscopic surgery.

Replacement Pads

508-0111 Schurefoot Disp Pads

Other required products for use: 800-0134 Schure Socket XL

US: 0.374" x 1.122" (9.5 mm x 28.5 mm) PN# 800-0134

Denyer: 0.236" x 1.496" (6 mm x 38 mm) PN# 800-0134-DEN

Europe: 0.394" x 0.984" (10 mm x 25 mm) PN# 800-0134-EU

Eschmann (UK): 0.236" x 1.260" (6 mm x 32 mm) PN# 800-0134-UK

Japan: 0.354" x 1.260" (9 mm x 32 mm) PN# 800-0134-JPN

Swiss: 0.394" x 1.181" (10 mm x 30 mm) PN# 800-0134-SWISS

GENERAL INFORMATION

- *Product not made with Natural Rubber Latex*
- *Product warranty covers product from manufacturing defects for period of 2 years*
- *If damaged in shipping, call Customer Service at (888) 724-8763 or (781) 982-7000 to obtain Return Material Authorization number. For product warranty issues, contact Customer Service.*
- *Product is maintenance-free, check product condition before next use*
- *Life of device is 5 years under normal use*
- *Store device between -4°F to +86°F (-20°C to 30°C)*

DISPOSAL

- **General** - *Prevent infection by cleaning and disinfecting product before disposal*
- **Packaging** - *Dispose packaging material via household waste according to national requirements*
- *SchureMed accepts back used or retired products - or dispose of product in accordance with national requirements*



WARNING!

Hazard resulting from incorrect use. Strictly follow instructions for your Operating Table system.



WARNING! *Do not reuse device if there are obvious signs of damage or functional issues. Consult manufacturer before reusing.*

CLEANING RECOMMENDATION

Follow current Association of periOperative Registered Nurses Journal Guidelines for proper cleaning and disinfection procedures.



WARNING!

Adhere to standards for blood-borne pathogens from the Occupational Safety and Health Administration. Use recommended protective clothing, gloves, masks and eye protection to clean accessory.

CAUTION

Strictly read/follow manufacturer's directions for cleaning fluids. **DO NOT** use cleaners containing phenolics.

1. Remove major contaminants from accessory with disposable materials. Follow appropriate bio-hazard waste disposal procedures.
2. Apply cleaning fluid liberally to entire accessory and wipe with clean, lint-free cloth until all moisture and cleaning fluid is removed from accessory
3. Let accessory dry












USER NOTICE

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

eIFU Language Versions

To download and print the Instructions for Use, please go to <https://www.schuremed.com>, select the product, and click on the drop-down menu for currently available translations for the IFU.

Symbol Glossary

Symbol	Title
	Manufacturer
	Date of manufacture
	Authorized Representative in the European Community
	Authorized Representative in the Swiss
	Importer
	Catalogue Number
	Serial Number
	Warning
	Medical Device
	Unique Device Identifier
	CE Marking



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