



Instructions for Use

REF Foot Extension, 20 x 10
800-0036

REF Foot Extension, 20 x 10 (EU)
800-0036-EU

Replacement Pad
Foot Extension Pad 508-0090



INTENDED USE

Foot Extension mounts horizontally as table extension or vertically as foot support when using reverse Trendelenburg. The intended users of this device are medical professionals within hospitals and surgery centers.

GENERAL INFORMATION

- *Product not made with Natural Rubber Latex*
- *Device supports 500 lb. (227 kg) proportional patient load (6'4" (193 cm) tall patient per 99% human body model)*
- *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.*
- *CE marked medical device according to MDR (EU) 2017/745*
- *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)*

INSTRUCTIONS

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

Attaching Foot Extension to Surgical Table

1. *Attach Simple Clamps P/N 800-0228 (sold separately) onto side rails*
2. *Insert mounting posts into Simple Clamps on both sides of table*
3. *Turn handle clockwise to tighten clamps*

Detaching Foot Extension from Surgical Table

1. *Turn simple clamp handle counterclockwise to loosen foot extension*
2. *Lift foot extension from operating room table and store properly*

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- *Length: 10" +/- 0.5" (25 cm +/- 1 cm)*
- *Width: 20" +/- 0.5" (51 cm +/- 1 cm)*
- *Depth: 3.75" +/- 0.5" (10 cm +/- 1 cm) (with pad)*
- *Device Weight: 5.25 +/- 0.5 lbs. (2.4 kg +/- .22 kg)*
- *Attaches to rail of surgical table at foot section*
- *Single-person installation*

COMPONENT OVERVIEW

Foot Extension is operating room table extension or used as foot support. Simple Clamp (sold separately) required for proper use of Foot Extension.

See side rail dimensions below:

US: 0.374" x 1.122" (9.5 mm x 28.5 mm) PN# 800-0228

Denyer: 0.236" x 1.496" (6 mm x 38 mm) PN# 800-0228-DEN

Europe: 0.394" x 0.984" (10 mm x 25 mm) PN# 800-0228-EU

Eschmann (UK): 0.236" x 1.260" (6 mm x 32 mm) PN# 800-0228-UK

Japan: 0.354" x 1.260" (9 mm x 32 mm) PN# 800-0228-JPN

Swiss: 0.394" x 1.181" (10 mm x 30 mm) PN# 800-0228-SWISS

Replacement Pad: Foot Extension Pad 508-0090

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



PRODUCT USE WARNINGS

WARNING!

Maximum load should not exceed appropriate proportion of a patient weighing 500 lbs. (227 kg). Use care with low-maximum load capacity surgical tables that accessory rails are not overloaded.



WARNING!

Hazards result from incorrect use. Strictly follow Instructions for Use with 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 www.schuremed.com

Symbol Glossary

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



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