Instructions for Use

REF	**Stirrup Clamp**	**800-0396**
REF	**Stirrup Clamp (EU)**	**800-0396-EU**
REF	**Stirrup Clamp (UK)**	**800-0396-UK**
REF	**Stirrup Clamp (JPN)**	**800-0396-JPN**
REF	**Stirrup Clamp (SWISS)**	**800-0396-SWISS**
REF	**Stirrup Clamp (DEN)**	**800-0396-DEN**

INTENDED USE

Intended use is to hold operating room accessories on surgical table. The clamp is primarily used together with our Great White Stirrups. 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 use on a nurse, physician or appropriate volunteer prior to using clinically.

Stirrup Clamp attaches anywhere along surgical table's side rail. Accepts 1" x 1/4" (2.5 cm x .64 cm) flat bar mounting post. Holds all accessories in vertical position.

Attaching to Side Rail

1. *Place clamp on rail so flat bottom is in down position and the SchureMed logo sticker is facing up*
2. *Push clamp down on rail until it attaches to the rail.*
3. *Secure clamp by turning handle clockwise to locked position*

Detaching from Side Rail

1. *Unlock clamp by turning handle counterclockwise and remove accessory*
2. *Slide clamp away from standoffs, then push top of clamp down with thumb until it releases from rail, then grab handle and lift clamp from side rail*

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- *Length:2.75" +/- 0.5" (7 cm +/- 1 cm)*
- *Height: 1.94" +/- 0.5" (4.9 cm +/- 1 cm)*
- *Depth: 2.98"+/- 0.5" (6.8 cm +/- 1 cm)*
- *Device Weight: 0.5 +/- 0.5 lbs. (.22 kg +/- .22 kg)*
- *Attaches to surgical table anywhere on rail*
- *Single-person installation*

COMPONENT OVERVIEW

Stirrup Clamp is a surgical table clamp that attaches to surgical table side rails. Clamp is primarily used in conjunction with our Great White Stirrups.

See side rail dimensions below:
***US**: 0.374" x 1.122" (9.5 mm x 28.5 mm) PN# 800-0396*
***Denver**: 0.236" x 1.496" (6 mm x 38 mm) PN# 800-0396-DEN*
***Europe**: 0.394" x 0.984" (10 mm x 25 mm) PN# 800-0396-EU*
***Eschmann (UK)**: 0.236" x 1.260" (6 mm x 32 mm) PN# 800-0396-UK*
***Japan**: 0.354" x 1.260" (9 mm x 32 mm) PN# 800-0396-JPN*
***Swiss**: 0.394" x 1.181" (10 mm x 30 mm) PN# 800-0396-SWISS*

GENERAL INFORMATION

- *Product not made with Natural Rubber Latex*
- *Device supports 1,000 lb. (454 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.*
- *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*

Product Use Warnings

WARNING!

Maximum load should not exceed appropriate proportion of a patient weighing 1,000 lbs. (454 kg). Use care with low-maximum load capacity surgical tables that accessory rails are not overloaded.

Cleaning Recommendation

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

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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