



Instructions for Use

- REF** SpringLoc Clamp 800-0338
- REF** SpringLoc Clamp (EU) 800-0338-EU
- REF** SpringLoc Clamp (JPN) 800-0338-JPN
- REF** SpringLoc Clamp (SWISS) 800-0338-SWISS

INTENDED USE

Intended use is to hold operating room accessories on surgical table. Clamp is primarily used together with our Great White Stirrups. The intended users of this device are medical professionals within hospitals and surgery centers.



GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- *Length: 3" +/- 0.5" (8 cm +/- 1 cm)*
- *Width: 2.5" +/- 0.5" (6 cm +/- 1 cm)*
- *Depth: 2.25" +/- 0.5" (6 cm +/- 1 cm)*
- *Handle Collapsed: 5.5" +/- 0.5" (14 cm +/- 1 cm)*
- *Handle Extended: 5.5" +/- 0.5" (14 cm +/- 1 cm)*
- *Device Weight: 0.5 +/- 0.5 lbs. (.22 kg +/- .22 kg)*
- *Attaches to surgical table anywhere on rail*
- *Single-person installation*

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.

SpringLoc Clamp is spring-loaded and 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
2. Push clamp down on rail until springs click
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 clicks, then grab handle and lift clamp from side rail

COMPONENT OVERVIEW

SpringLoc 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# 800-0338

Europe: 0.394" x 0.984" (10 mm x 25 mm) PN# 800-0338-EU

Japan: 0.354" x 1.260" (9 mm x 32 mm) PN# 800-0338-JPN

Swiss: 0.394" x 1.181" (10 mm x 30 mm) PN# 800-0338-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.
- 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)

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.

**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 (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.

UDI Basic UDI-DI: 081001460F0041DW

eIFU Language Versions

To download and print the Instructions for Use, please go to <http://www.schuremed.com/schuremed-eifu>.

Symbol Glossary

Symbol	Title	Symbol Description
	Manufacturer	Indicates the medical device manufacturer.
	Authorized Representative in the European Community	Indicates the authorized representative in the european community.
	Use-by Date	Indicates the date after which the medical device is not to be used.
	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Catalogue or Item Number	Indicates the manufacturer's catalogue or item number so that the medical device can be identified.
	Serial Number	Indicates the manufacturer's serial number so that a specific medical device can be identified.
	Caution	To indicate that caution is necessary when operating the device or control close to where the symbol is placed, or to indicate that the current situation needs operator awareness or operator action in order to avoid undesirable consequences.
	Medical Device	Indicates the item is a medical device.
	Unique Device Identifier	Indicates a barcode as containing unique device identifier information.
	CE Marking	European Conformity.
	Single Patient Use	Indicates the item is a single patient use medical device.



Manufacturer

SchureMed (081001460)

452 Randolph Street, Abington, MA 02351 USA

Toll Free (888) 724-8763 | **Ph** (781) 982-7000 | **Fax** (781) 982-7001 | **orders@schuremed.com**



Authorized Representative

Emergo Europe, Prinsessegracht 20, 2514 AP The Hague, The Netherlands