

Instructions for Use

REF Simple Clamp 800-0228

REF Simple Clamp (EU) 800-0228-EU

REF Simple Clamp (UK) 800-0228-UK

REF Simple Clamp (DEN) 800-0228-DEN

REF Simple Clamp (JPN) 800-0228-JPN

REF Simple Clamp (SWISS) 800-0228-SWISS

INTENDED USE

Intended use is to hold surgical accessories with 1" x 3/8" (2.5 cm x .95 cm) flat mounting post vertically anywhere along table side rail or a 1" x 1/4" (2.5 cm x .64 cm) horizontally mounting blades. The intended users of this device are medical professionals within hospitals and surgery centers.



INSTRUCTIONS

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

Attaching to Side Rail

- 1. Snap Simple Clamp on surgical table side rail
- 2. Place flat bar of accessory into mounting hole
- 3. Secure clamp by turning handle clockwise to lock position

Detaching from Side Rail

- 1. Unlock clamp by turning the handle counterclockwise and remove mounting bar
- 2. Lift simple clamp off surgical table side rail

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- Length: 3" +/- 0.5" (8 cm +/- 1 cm)
- Width: 1.75" +/- 0.5" (4 cm +/- 1 cm)
- Depth: 1.375"+/- 0.5" (4 cm +/- 1 cm)
- Device Weight: 1 +/- 0.5 lbs. (.45 kg +/- .22 kg)
- Attaches to rail of a surgical table at any point on rail
- Single person installation
- Holds accessory equipment with 1" x 3/8" (2.5 cm x .95 cm) flat mounting posts

COMPONENT OVERVIEW

Simple Clamp is a surgical table clamp that attaches to all rails. Its function is to hold surgical accessories on a surgical table.

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

GENERAL INFORMATION

• Product not made with Natural Rubber Latex

- Device supports 720 lb. (327 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 $-4^{\circ}F$ to $+86^{\circ}F$ ($-20^{\circ}C$ to $30^{\circ}C$)

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 720 lb. (327 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.

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
M	Date of manufacture
EC REP	Authorized Representative in the European Community
CH REP	Authorized Representative in the Swiss
	Importer
SN	Serial Number
\triangle	Warning
MD	Medical Device
UDI	Unique Device Identifier
CE	CE Marking



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