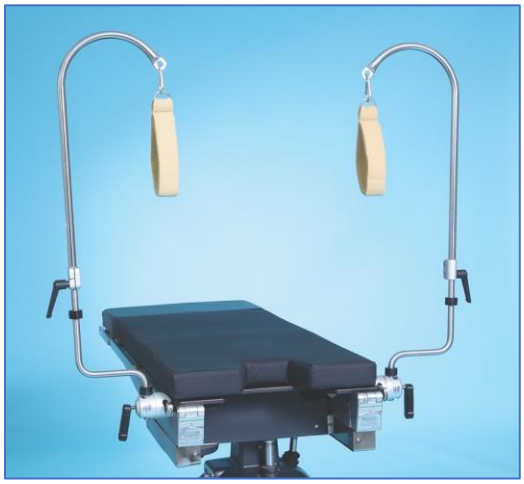
Instructions for Use



REF **Candy Cane Stirrups 800-0012**

Replacement Part
Candy Cane Stirrups Repl. Strap 790-0006

INTENDED USE

Intended use is to hold patient's legs during short or long gynecologic, urological or laparoscopic 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 use on a nurse, physician or appropriate volunteer prior to using clinically.

Attaching Stirrups to Surgical Table

1. *Attach candy cane stirrup mounting post into Schure Socket XL P/N 800-0134 (sold separately), turning handle of Schure Socket XL clockwise to tighten.*
2. *Guide patient's leg into ankle stirrup*

Adjusting Stirrup

1. *Stainless Steel Candy Cane Stirrups adjust vertically from 28" to 44" (71 cm to 112 cm) with an 8" (20 cm) leg clearance. Loosen handle to adjust to desired position.*
2. *Lock stirrup height by twisting ergonomic handle*

Detaching Stirrup from Table

1. *Turn handle of Schure Socket XL counter clockwise to loosen and remove stirrup from socket*

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- *Length: 44" +/- .5" (112 cm +/- 1 cm)*
- *Extended: 28" +/- .5" (71 cm +/- 1 cm) Collapsed*
- *Width: 17" +/- .5" (43 cm +/- 1 cm)*
- *Depth: 10.5" +/- .5" (27 cm +/- 1 cm)*
- *Device Weight Per Stirrup: 5 +/- 0.5 lbs. (2.2 +/- .22 kg)*
- *Connects to side rails of a surgical table*
- *Single-person installation*
- *Stirrup has 5/8" (1.6 cm) mounting post which attaches with Schure Socket XL P/N 800-0134 (sold separately)*
- *Ankle Strap: 18"L x 2"W (46 cm +/- 5 cm)*

- *Replacement Ankle Strap P/N 790-0006 (sold separately as an each)*

COMPONENT OVERVIEW
Candy Cane Stirrup is a surgical table accessory aiding in positioning the legs for gynecological, urological, or laparoscopic procedures.

Replacement Part
790-0006 Candy Cane Stirrup Strap

Other required products for use: 800-0134 Schure Socket XL, 2 each (sold separately)

***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*
- *Device supports 350 lb. (159 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 350 lbs. (159 kg). Use care with low-maximum load capacity surgical tables that accessory rails are not overloaded.

WARNING!
Hazard resulting from incorrect use. Strictly follow Instructions for Use for your Operating Table system.

WARNING!
Surgical table load capacities may be less. Never overload surgical table. Device is intended for mounting on side rail of surgical table only.

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 procedure.

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.

UDI Basic UDI-DI: 081001460F0031DT

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.
EC REP	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.
LOT	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified.
REF	Catalogue or Item Number	Indicates the manufacturer's catalogue or item number so that the medical device can be identified.
SN	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.
MD	Medical Device	Indicates the item is a medical device.
UDI	Unique Device Identifier	Indicates a barcode as containing unique device identifier information.
CE	CE Marking	European Conformity.
	Single Patient Use	Indicates the item is a single patient use medical device.

Manufacturer
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CE MD

EC REP **Authorized Representative**
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