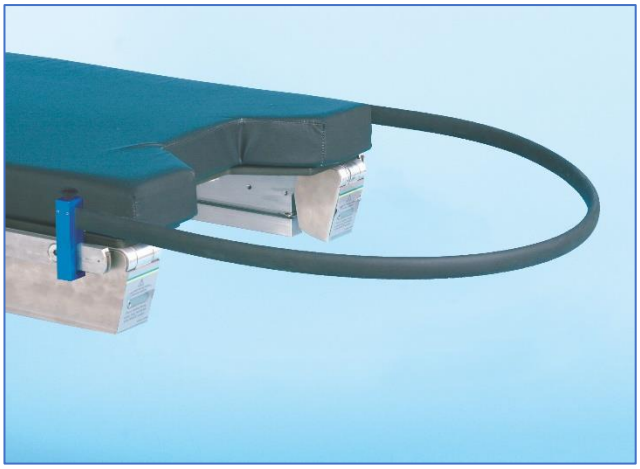
Instructions for Use



REF **Uro-Trapper System 800-0096**

Disposable Accessories
Uro-Trapper Drain Bags 800-0095
Uro-Trapper Drain Bags, Sterile 800-0098

INTENDED USE
Intended use is to hold a drain bag which collects patient's fluid during a surgical procedure. The intended users of this device are medical professionals within hospitals and surgery centers.

GENERAL SPECIFICATIONS
Device Dimensions (maximum)

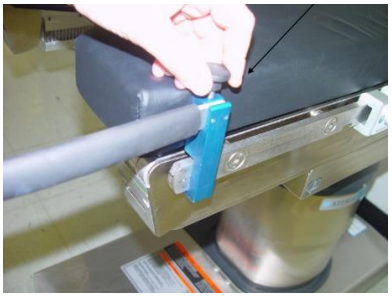
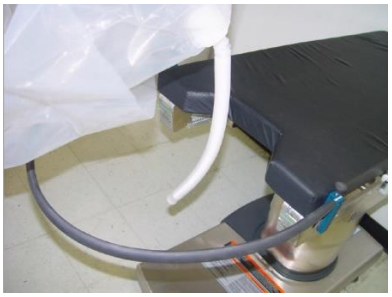
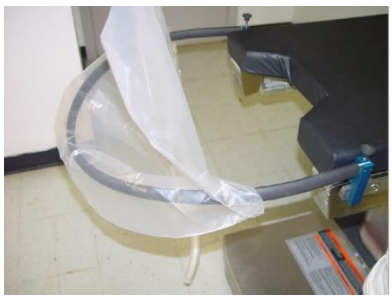
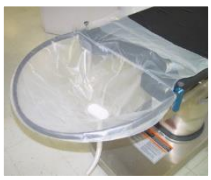
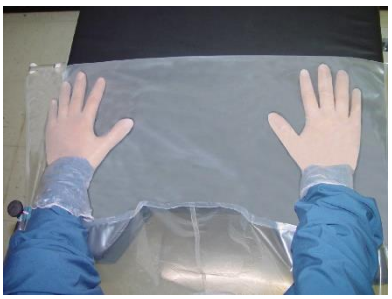
- *Length: 49" +/- 0.5" (125 cm +/- 1 cm)*
- *Width: 5" +/- 0.5 (13 cm +/- 1 cm)*
- *Device Weight: 2 +/- 0.5 lbs. (.9 +/- .22 kg)*
- *Attaches to end of surgical table side rail*
- *Single-person installation*

GENERAL INFORMATION
- *Product not made with Natural Rubber Latex*
- *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)*

COMPONENT OVERVIEW
Urology Trapper System is used to hold a drain bag that stores fluid from surgical procedures.

Other required products for use:
800-0095 Uro-Trapper Drain Bags or **800-0098** Uro-Trapper Drain Bags, Sterile

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.

	***Step 1***: *Connect mounting clamp onto side rail as shown repeat Step 1 for opposite side* ***Step 2***: *Lock down both sides using locking knobs*
	***Step 3:*** *Insert 800-0095 or 800-0098 hose inside hoop before sliding Drain Bag on*
	***Step 4:*** *Install Drain Bag onto hoop frame as shown below*
	***Step 5:*** *Slide Drain Bag down hoop until cuff reaches table pad as shown.* 
	***Step 6:*** *Tuck top of Drain Bag under patients' buttocks. Ensure drainage screen is at bottom most part of drain to allow full drainage of fluids.*

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!
Item does not hold a patient therefore item does not have any patient weight restrictions.

WARNING! *Hazard resulting 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.

UDI Basic UDI-DI: 081001460F0049EE

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
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**

CE MD

EC REP **Authorized Representative**
Emergo Europe, Prinsessegracht 20, 2514 AP The Hague, The Netherlands