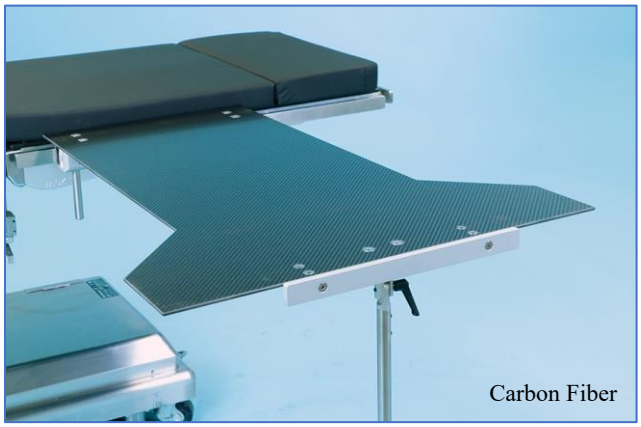
Instructions for Use

REF **End Rest Procedure Table w/Leg, Phenolic 800-0026-ER-P**

REF **End Rest Procedure Table w/Leg, Carbon Fiber 800-0026-ER-CF**

Replacement Pad
End Rest Procedure Table Deluxe Pad
508-0101


Carbon Fiber

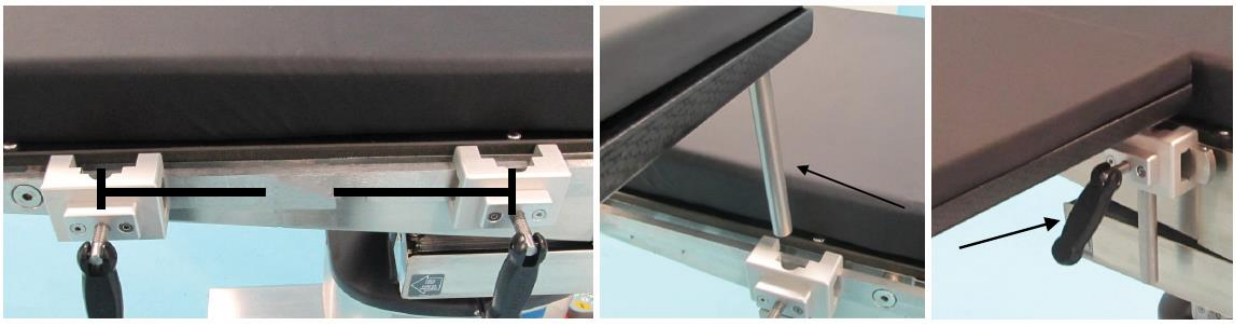
INTENDED USE
Its function is to provide a stable platform for all arm and hand surgeries. The intended users of this device are medical professionals within hospitals and surgery centers.

GENERAL SPECIFICATIONS
Device Dimensions (maximum)

- *Length: 34" +/- 0.5" (86 cm +/- 1 cm)*
- *Width: 16"+/- 0.5" (41 cm +/- 1 cm)*
- *Depth: 42" +/- 0.5" (107 cm +/- 1 cm)*
- *Phenolic Device Weight: 17 +/- 0.5 lbs. (7.7 +/- .22 kg)*
- *Carbon Fiber Device Weight: 13 +/- 0.5 lbs. (5.8 +/- .22 kg)*
- *Attaches to surgical table anywhere on the rail using two rail clamps*
- *Single-person installation*
- *Clean with hospital grade disinfectant*

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 Procedure Table to Surgical Table



1. *Attach clamps (sold separately) to side rail and place them approximately 1' (31 cm) apart (as shown)*
2. *Insert mounting rods into clamps*
3. *Adjust procedure table so that it is level with surgical table pad. Secure table to accessory rail using clamp locking knobs*
4. *Includes 1" (3 cm) pressure management pad*
5. *Clamps sold separately*

Adjusting Table Leg

(1) End Rail • **(2)** Adjustable Swivel Leg Locking Handle • **(3)** Adjustable Height Locking Handle

1. *Swivel leg in 360° to obtain unlimited C-arm axis (see above)*
2. *Loosen Adjustable Leg Handle (3) and extend leg until it sits firmly against floor then lock handle tight by turning it to the right*
3. *End rail accommodates traction attachments*

COMPONENT OVERVIEW

End Rest Procedure Table w/Leg, Phenolic or Carbon Fiber, are surgical table extensions that attach to all rails.

Replacement Pad

End Rest Procedure Table Deluxe Pad 508-0101

Two Clamps Required: Choose Universal Clamp for US or Deluxe Rail Clamp for Outside US

US – Universal Rail Clamp

- *US: 0.374" x 1.122" (9.5 mm x 28.5 mm) PN# 800-0085*

Outside US – Deluxe Rail Clamp

- *Denyer: 0.236” x 1.496” (6 mm x 38 mm) PN# 800-0248-DEN*
- *Europe: 0.394” x 0.984” (10 mm x 25 mm) PN# 800-0248-EU*
- *Eschmann (UK): 0.236” x 1.260” (6 mm x 32 mm) PN# 800-0248-UK*
- *Japan: 0.354” x 1.260” (9 mm x 32 mm) PN# 800-0248-JPN*
- *Swiss: 0.394” x 1.181” (10 mm x 30 mm) PN# 800-0248-SWISS*

GENERAL INFORMATION

- *Product not made with Natural Rubber Latex*
- *Device supports 500 lb. (227 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 500 lbs. (227 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 procedure.

WARNING!

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

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: 800-0026-ER-P – 081005737F0022KP

UDI Basic UDI-DI: 800-0026-ER-CF – 081005737F0023KR

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