



452 Randolph St, Abington, MA 02351, USA



TKR

POSITIONER

Total Knee Replacement

TKR Positioner, 24" (61 cm)	800-0141
TKR Positioner, 30" (76 cm)	800-0171
Genos Split TKR Positioner, 30" (76 cm)	800-0409



Prior to using this or any other type of medical equipment with a patient, it is recommended that you read the Instructions For Use and familiarize yourself with the product.

- Please read and understand all warnings in this manual and on the device itself before using it with a patient.
- The  symbol is designed to alert users to important procedures or safety instructions regarding the use of this device.
- The techniques described in this manual are suggestions from the manufacturer. The attending physician retains the final responsibility for patient care with respect to this device.
- Check the device function before each use. Do not use this device if there are obvious signs of damage or functional issues. Consult manufacturer before reusing.
- This device should only be operated by trained personnel.
- All modifications, upgrades, or repairs must be carried out by an authorized specialist.
- Any serious incidents related to the device should be reported to the manufacturer and the national competent authority where the user is located.



**NEVER EXCEED THE WEIGHT CAPACITY OR IMPROPERLY
DISTRIBUTE THE LOAD ON THE OPERATING ROOM TABLE.**

Symbol	Description	Reference
	Manufacturer	EN ISO 15223-1
	Date of manufacture	EN ISO 15223-1
	Authorized Representative in the European Community	EN ISO 15223-1
	Authorized Representative in Switzerland	EN ISO 15223-1
	Authorized Representative in UK	EN ISO 15223-1
	Importer	EN ISO 15223-1
	Serial Number	EN ISO 15223-1
	Warning	IEC 60601-1
	Medical Device	MDR 2017/745
	Unique Device Identifier	EN ISO 15223-1
	CE Marking	MDR 2017/745
	Use-By Date	EN ISO 15223-1
	Batch Code	EN ISO 15223-1
	Manufacturer's Reference Number	EN ISO 15223-1

Indication for Use:

The TKR is used in a variety of surgical procedures including but not limited to total knee replacement, partial knee replacement and revision. This device is capable of being used with a broad patient population as deemed appropriate by medical professionals within hospitals and surgery centers.

Intended Use:

Intended use of the TKR is to allow the surgeon to position the operative leg in multiple positions (extension, flexion, tilt, and rotation) during surgical procedures.

Intended User: Surgeons, Nurses, Doctors, Physicians and/or Healthcare professionals involved in the intended procedure utilizing the device. All staff using or preparing the intended product should be properly trained and familiar with the positioning device before use.

Intended Populations: This device is intended to be used with patients that do not exceed a 500 lb. (227kg) proportional patient load (6'4" (193 cm) tall patient per 99% human body model)

Residual Risk:

This product complies with relevant performance, safety standards. However, misuse, device damage, function or mechanical hazards cannot be completely excluded. End users are responsible for ensuring device is securely attached and will operate in a safe manner.

Safety Considerations:

DO NOT USE IF THE PRODUCT SHOWS VISIBLE DAMAGE OR SIGNS OF IMPROPER FUNCTIONING.

Equipment Misuse:

Do not use the product if package is damaged. All modifications, upgrades, or repairs must be performed by a SchureMed authorized specialist.

Safe Disposal:

End users should adhere to all federal, state, regional, and/or local laws and regulations as it pertains to the safe disposal of medical devices and accessories.

System Setup 800-0141, 800-0171:

1. Drape surgical table. Use the Standard CamLoc Rail Clamp (sold separately) to use with 1-2 drapes or the Large CamLoc Rail Clamp for 3-5 drapes.
2. Attach the CamLock Rail Clamp to rail over drapes. Flip Locking Lever right or left to secure.
3. Slide Track System Crossbar into CamLoc Rail Clamp and securely tighten by turning the CamLoc Rail Clamp Star Knob clockwise.



To prevent patient or operator injury from inadvertent device movement, do not attach the CamLoc Rail Clamp over notches within side rail.

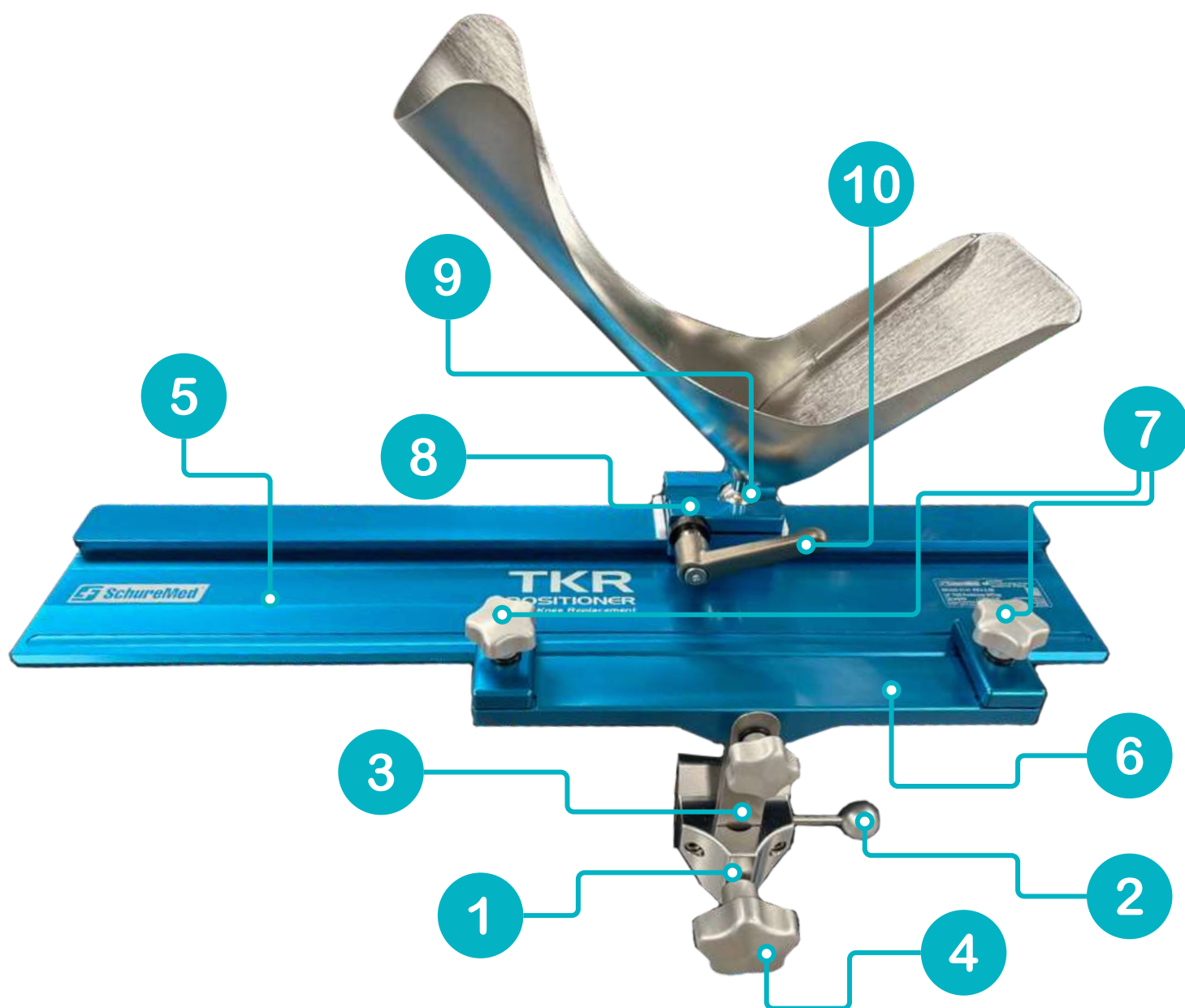
4. Slide Base Plate into Cross Support Bar making sure they are flush. Tighten two Star Knobs by turning clockwise.
5. Insert plastic slide in the opening under Adjustable Slide Block with notched side facing up. Make sure Locking Handle is rotated until it stops (don't tighten) before sliding on to the Base Plate. Locking Handle should be in vertical position pointing straight up. Depress button and pull Locking Handle to adjust as needed.
6. Slide Adjustable Slide Block onto track system while holding plastic slide up close to Adjustable Slide Block.
7. Place Adjustable Boot Ball into track system's Adjustable Slide Block.
8. Line Boot with SchureMed's TKR Boot Liner (800-0146)
9. Position patient, turn Adjustable Slide Block Locking Handle right to lock into desired position.



Only adjust Adjustable Slide Block Locking Handle Lever when it is on the track

Device Removal:

1. Turn Star Knob counter clockwise to loosen the Cross Support Bar.
2. Slide the TKR from the CamLoc Rail Clamp by sliding upwards.
3. Remove CamLoc Rail Clamp from accessory rail.
4. Disassemble all components and return them to the approved sterilization tray to support proper cleaning and help prevent loss of parts.



- 1 CamLoc Rail Clamp
- 2 Locking Lever
- 3 Track System Crossbar
- 4 Camlock Clamp Star Knob
- 5 Base Plate

- 6 Cross Support Bar
- 7 Star Knobs
- 8 Adjustable Slide Block
- 9 Adjustable Boot Ball
- 10 Locking Handle

System Setup 800-0409:

1. Drape surgical table. Use the Standard CamLoc Rail Clamp (sold separately) to use with 1-2 drapes or the Large CamLoc Rail Clamp for 3-5 drapes.
2. Attach the CamLock Rail Clamp to rail over drapes. Flip Locking Lever right or left to secure.
3. Slide Track System Crossbar into CamLoc Rail Clamp and securely tighten by turning the CamLoc Rail Clamp Star Knob clockwise.



To prevent patient or operator injury from inadvertent device movement, do not attach the CamLoc Rail Clamp over notches within side rail.

4. Insert plastic slide in the opening under Adjustable Slide Block with notched side facing up. Make sure Locking Handle is rotated until the opening underneath is sized to the Base Plate track before sliding on to the Base Plate. Locking Handle should be in vertical position pointing straight up. Depress button and pull Locking Handle to adjust as needed.
5. Slide Adjustable Slide Block onto either side Baseplate track system from the center while holding plastic slide up close to Adjustable Slide Block.

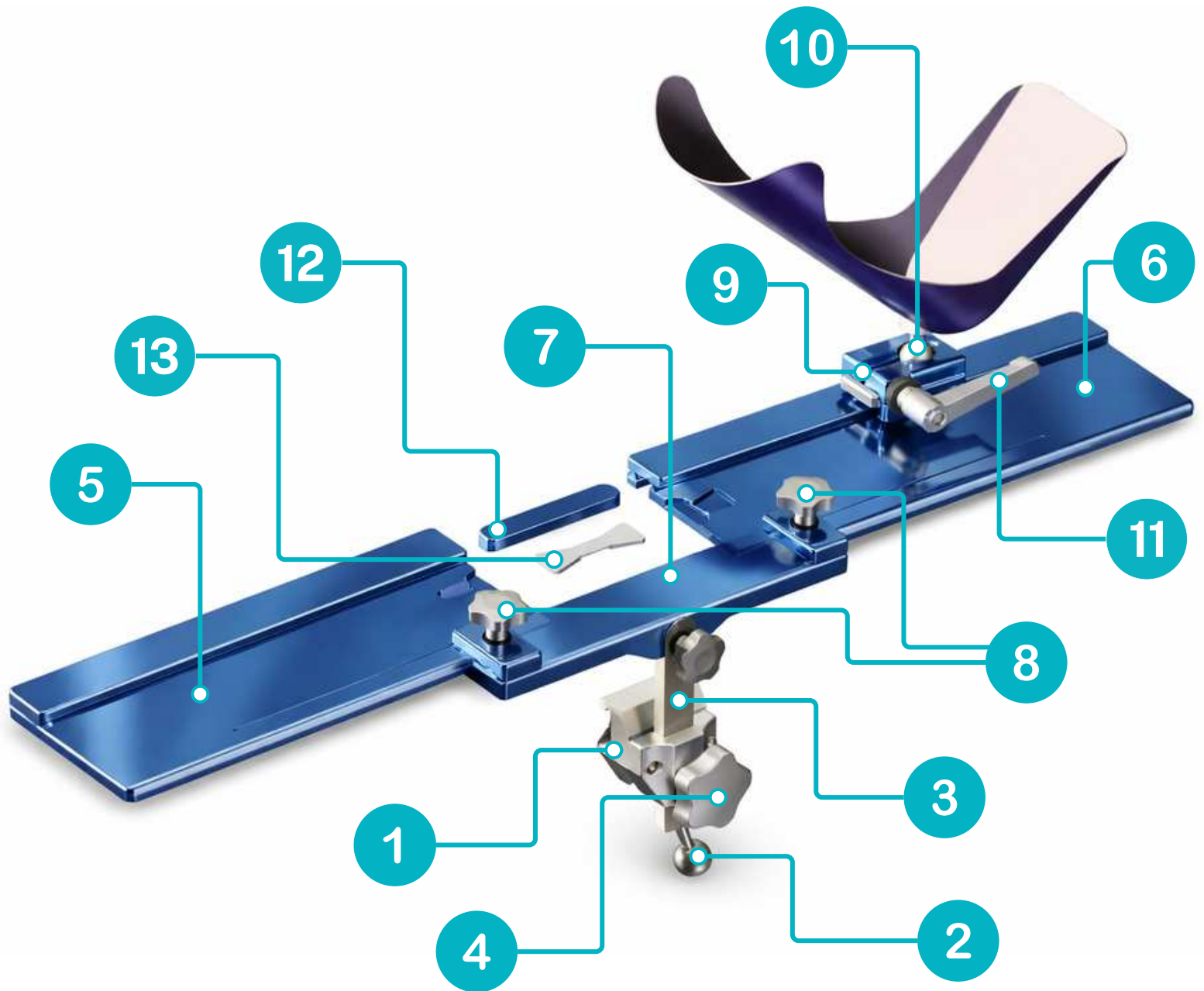


Adjustable Slide Block cannot be placed onto Baseplate track system after the Baseplate is assembled. It must be placed on the track from the center.

6. Place the Key into the opening under one side of the Base Plate and slide both Baseplates together until they are flush in the center. Then slide the assembled Base Plate into Cross Support Bar making sure they are flush. Tighten two Star Knobs by turning clockwise.
7. Insert the Joint Lock into the cutout in the center of the Baseplate with the smooth side facing up. The Joint Lock should be seated flush to the Base Plate surface.
8. Place Adjustable Boot Ball into track system's Adjustable Slide Block.
8. Line Boot with SchureMed's TKR Boot Liner (800-0146).
9. Verify all connections and knobs are securely tightened, and confirm the TKR is fully secured and will not move prior to patient placement in the device.
10. Position patient, turn Adjustable Slide Block Locking Handle right to lock into desired position.

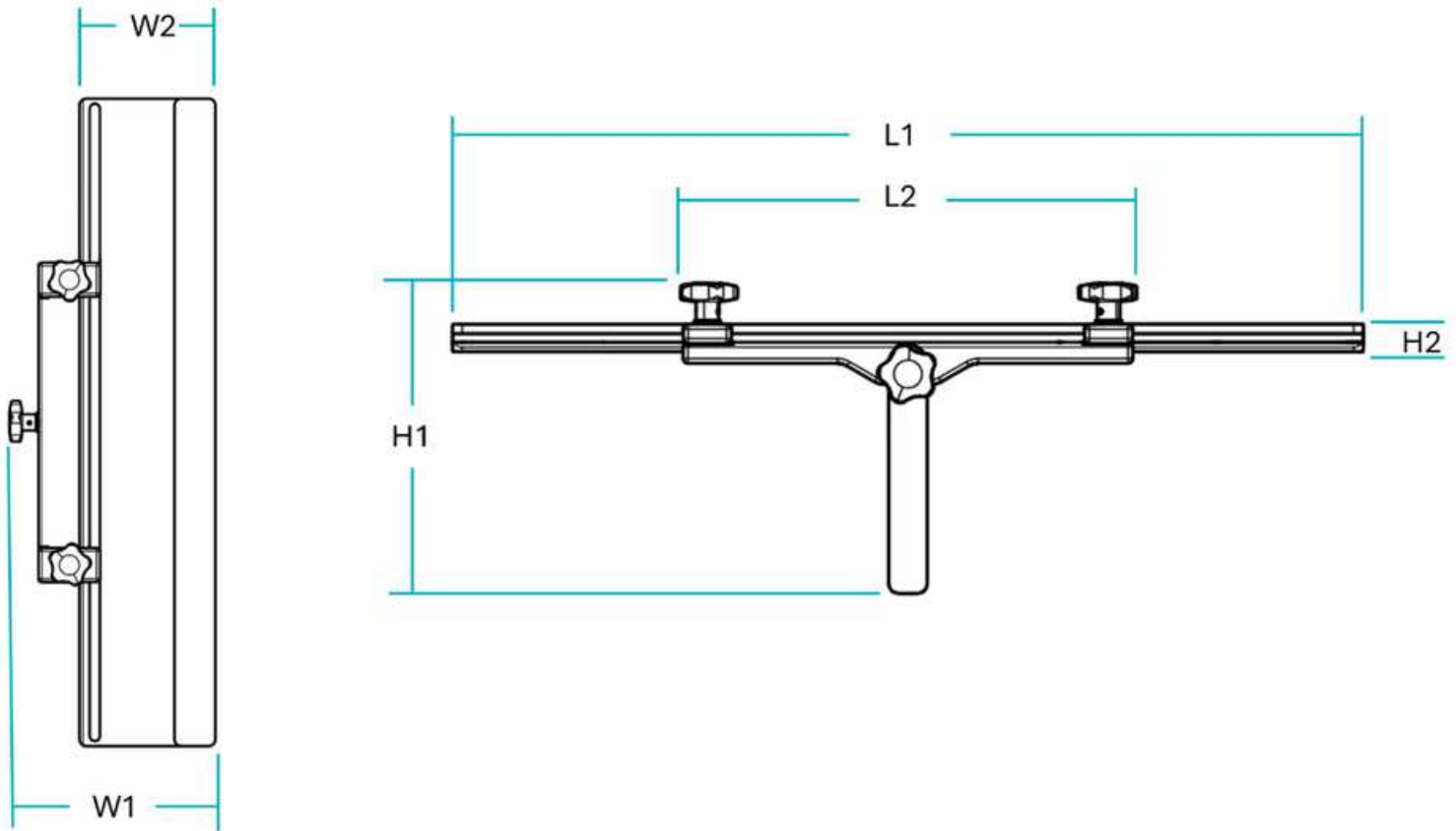
Device Removal:

1. Turn Star Knob counter clockwise to loosen the Cross Support Bar.
2. Slide the TKR from the CamLoc Rail Clamp by sliding upwards.
3. Remove CamLoc Rail Clamp from accessory rail.
4. Disassemble all components and return them to the approved sterilization tray to support proper cleaning and help prevent loss of parts.

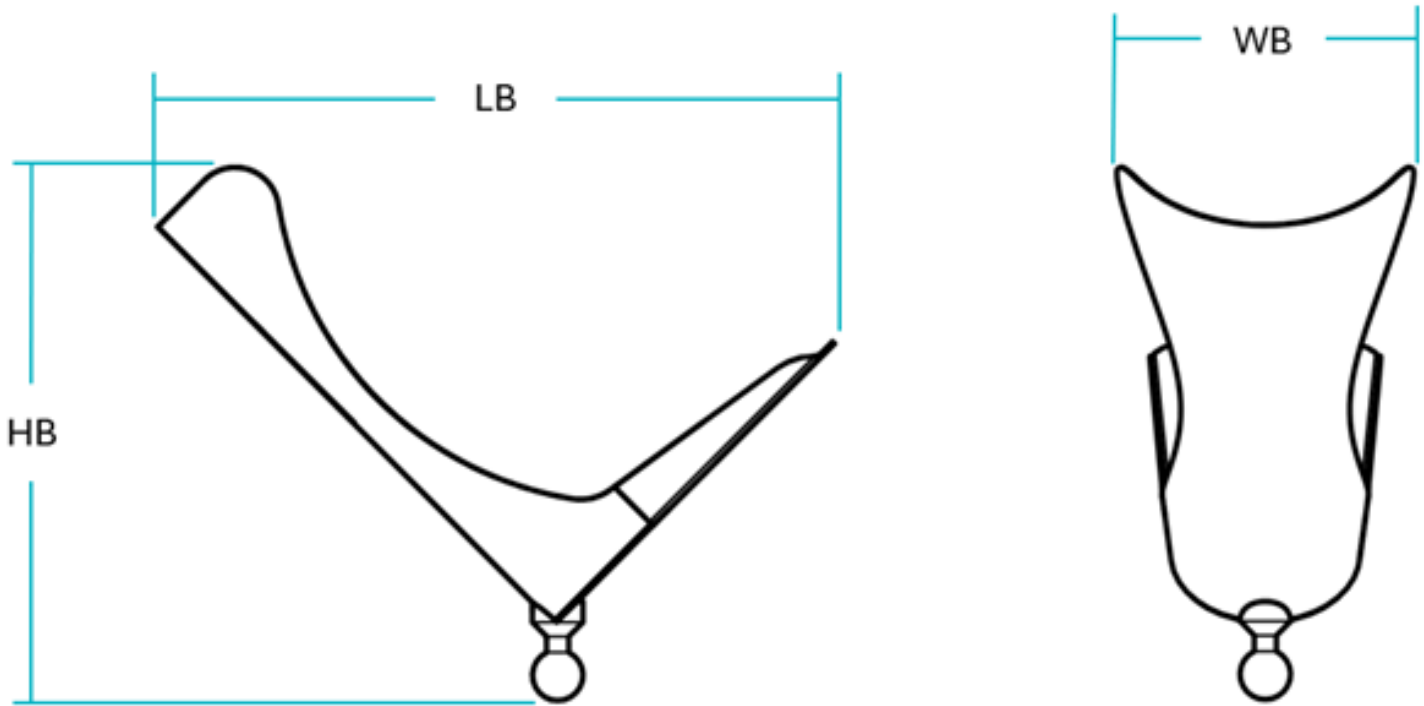


- 1 CamLoc Rail Clamp
- 2 Locking Lever
- 3 Track System Crossbar
- 4 Camlock Clamp Star Knob
- 5 Base Plate Left
- 6 Base Plate Right

- 7 Cross Support Bar
- 8 Star Knobs
- 9 Adjustable Slide Block
- 10 Adjustable Boot Ball
- 11 Locking Handle
- 12 Key
- 13 Joint Lock



Device Dimensions	TKR 24" (800-0141)	TKR 30" (800-0171) Genos 30" (800-0409)
Total Length (L1)	23.7" +/- 0.5" (61cm +/- 1cm)	29.7" +/- 0.5" (76cm +/- 1cm)
Mount Length (L2)	11.7" +/- 0.5" (29.7cm +/- 1cm)	11.7" +/- 0.5" (29.7cm +/- 1cm)
Total Width (W1)	7.5" +/- 0.5" (19cm +/- 1cm)	7.5" +/- 0.5" (19cm +/- 1cm)
Base Width (W2)	5.0" +/- 0.5" (13cm +/- 1cm)	5.0" +/- 0.5" (13cm +/- 1cm)
Total Height (H1)	8.1" +/- 0.5" (20cm +/- 1cm)	8.1" +/- 0.5" (20cm +/- 1cm)
Base Height (H2)	0.7" +/- 0.5" (1.8cm +/- 1cm)	0.7" +/- 0.5" (1.8cm +/- 1cm)
Net Weight	9lbs +/- 0.5lbs (4kg +/- .22kg)	10lbs +/- 0.5lbs (4.5kg +/- .22kg)
Gross Weight	12lbs +/- 0.5lbs (5.6kg +/- .22kg)	14.5lbs +/- 0.5lbs (6.5kg +/- .22kg)



Boot Dimensions	STAINLESS STEEL TKR BOOT (700-1107)
Boot Length (LB)	13.3" +/- 0.5" (34cm +/- 1cm)
Boot Width (WB)	5.9" +/- 0.5" (15cm +/- 1cm)
Boot Height (HB)	10.5" +/- 0.5" (27cm +/- 1cm)
Boot Weight	1.5lbs +/- 0.5lbs (0.68kg +/- .22kg)

Storage Specifications	Description
Storage Temperature	-20° F to 140° F (-29° C to +60° C)
Storage Relative Humidity Range	15% to 85%
Operating Temperature	This device is intended to be used in a controlled Operating Room environment.
Operating Relative Humidity Range	

Cleaning and Disinfection:



- **Clean the device after each use as directed.**
- **Do not submerge the device in liquid.**
- **Use caution in areas where liquid can get into the mechanism.**
- **Do not clean the device with bleach or products that contain bleach.**

Hand and Washer Machine Cleaning Instructions

- Clean TKR Positioner thoroughly before using (before sterilization) and after each use
- Recommended cleaning products: (All solutions should be used in accordance with manufacturer's instructions)
 - Enzymatic neutral pH detergent solution
 - (Enzymatic cleaners are compatible with metal devices/instruments and are preferred)
 - Non-enzyme, hydrogen peroxide based solution
- Disassemble TKR Positioner into separate parts for cleaning:
 - Boot Ball
 - Mounting Plate with Clamp
 - Base Plate with Track
 - Slide Block with Handle
 - Plastic Slide

Hand Cleaning

- Scrub each piece with a soft, plastic scrub brush and a recommended cleaning solution (see above)
- Rinse all parts thoroughly to remove any chemical residue
- Wipe each part completely dry with a clean, lint-free cloth

Washer Cleaning

- Put through washer/decontaminator according to manufacturer's instruction (Select cycle that does not include lubrication)
- Prepare for sterilization (device must be cleaned before sterilization)
- See sterilization instructions below

Steam Sterilization

- Place the TKR into the sterilization tray and wrap using envelope method or accepted wrap method at your facility.
- Follow your hospital or facility's policy on steam sterilization procedures

#800-0141-TRAY 24" TKR Sterilization Tray



#800-0171-TRAY 30" TKR Sterilization Tray



Read the cleaning product's directions and follow the instructions on the label. Use caution in areas where fluid migration may occur.

Wipe device with a clean, dry cloth. Be sure that the product is dry prior to reinstalling and storage to avoid damage.

CAUTION: Damage may result if product is cleaned with caustic chemicals or harsh abrasives

ATTENTION: If any SchureMed product is damaged or does not function normally, discontinue use and contact SchureMed or an authorized SchureMed Certified Distributor.

Compatible Product Table:

Accessory Description	Product Number
CamLoc Rail Clamp	800-0271, 800-0271-DEN, 800-0271-EU, 800-0271-JPN, 800-0271-UK, 800-0271-SWISS
CamLoc Rail Clamp (Large)	800-0271-L
CamLoc Rail Clamp (Low Profile)	800-0271-LP
TKR Boot Liner	800-0146
Sterilization Tray 24"	800-0141-TRAY
Sterilization Tray 30"	800-0171-TRAY
Genos Sterilization Tray	800-0409-TRAY

Compliance with Medical Device Regulations



These products are non-invasive, Class I Medical Devices and are CE-marked according to AnnexVIII, Rule 1, of the Medical Device Regulations (REGULATION (EU) 2017/745).

EC Authorized Representative



Emergo Europe

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Manufacturing Information



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