



Thigh Garments RTG30

Universal size up to 26" (0.66m) thigh circumference

Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician.

Instructions for use

Secure the pump at the foot of the bed or in place of use. Attach the tube set to the pump connector by pushing in the silver latch on the female pump connector and then inserting the male tube connector into the pump connector until a click is heard.

Garments may be used on either leg. Unfold the garment(s) and position the inflatable center bladder directly behind the patient's calf with the cut out BEHIND the knee. The outlet tube on the garment should not be in direct contact with the patient's skin.

SNUGLY wrap the garment(s) around the patient's leg(s) and secure the hook tabs.

Attach the garment(s) to the tube set by pushing in the silver latch on the female tube connector and inserting the male garment connector firmly into the tube connector until a click is heard.

Ensure the pump is held firmly onto the bed frame while used in bed. During ambulation or transportation, ensure that the pump is secured. Rotate the outlet tube on the garment to eliminate kinks in the tubing. Tubing may become a trip hazard during ambulation if the shoulder strap is not used to secure excess tubing.

Turn the pump on by pressing the POWER button. If the LOW BATTERY light is solid red and producing a constant audible alarm, the battery will need to be charged. To charge the unit, plug the supplied power adaptor into a power outlet and then plug the round connector at the end of the cord into the top of the pump. The BATTERY CHARGE light will flash green while the unit is charging. The unit may be used while it is being charged. When fully charged, after 16-48 hours, an audible double-beep is presented once and the BATTERY CHARGE light (green) will be lit constantly.

Ensure the TYPE light indicates LEG GARMENT. If not, press the TYPE button until this light is illuminated.

Ensure the MODE light is not lit for SINGLE GARMENT. If only one garment is needed, press the MODE button to select SINGLE GARMENT and store the unused garment and tube for later use.

The pump is now ready for use. Refer to the user operating manual for complete information on the use of the system.

Indications

Intermittent Pneumatic Compression (IPC) is indicated for use for the prevention of deep vein thrombosis due to the presence of risk factors for thrombus formation during orthopedic, trauma, urologic, neurology, critical care, general medicine, obstetrics, and general surgery.

Recommendations

Garments should be removed regularly to inspect the skin.

Patients should be instructed in the proper use of the system and should report any problems to the nursing staff.

The system should be used continuously until the person is fully ambulatory. The therapy can continue with leg garments for patients with limited mobility.

If the garment cannot be applied to a limb during surgery, it may be applied to the limb once the patient reaches the recovery room. For a non-surgical patient, the system should be applied as soon as the risk of DVT is identified.

Contraindications

The Restep® system should not be used in the following conditions:

- Severe arteriosclerosis or active infection.
- Suspected or known acute DVT.
- Severe congestive heart failure or any condition where increased fluid to the heart may be detrimental.
- Existing pulmonary edema.
- Local skin or tissue conditions in which the garments would interfere.

If in doubt, refer to the patient's physician before using device.

Cautions

Garments should be removed immediately if the patient experiences any unexplained sensations, numbness or pain.

When used for DVT prophylaxis, continuous use is recommended and any prolonged interruption of therapy should be done in consultation with the patient's physician.

Garments are single-patient use, non-sterile, and latex-free. Garment bladders are PVC-free.



Pack contains one Pair Compression Sleeves
Non-Sterile (Devices HLD Processed)

For use with Restep
Pump RSP101 only.

Made in the Dominican Republic

Sustainability Solutions

Reprocessed By:
Stryker Sustainability Solutions
1810 W. Drake Dr.
Tempe, AZ (USA) 85283



DO NOT Reuse



Consult Instruction
for Use



Date of
Reprocessing

CSL Rev. C 09-2012 RM702110

Restep® is a registered trademark of Stryker Corporation.

Restep DVT System

Warranty

Reprocessed Products

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STRYKER SHALL NOT BE LIABLE FOR ANY DAMAGES TO THE EXTENT CAUSED BY ANY DEFECT IN MATERIAL, WORKMANSHIP OR DESIGN BY THE ORIGINAL MANUFACTURER OF THE PRODUCT OR ANY ACT OR OMISSION OF THE ORIGINAL MANUFACTURER OF THE PRODUCT.

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If a valid warranty claim is received within thirty (30) days of the expiration of the applicable warranty period, Stryker will, in its sole discretion: (1) replace the product at no charge with a product that is at least functionally equivalent to the original product or (2) refund the purchase price of the product. If a refund is provided by Stryker, the product for which the refund is provided must be returned to Stryker and will become Stryker's property. In any event, Stryker's liability for breach of warranty shall be limited to the replacement value of the defective or non-conforming part or component.

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