Restep®

DVT System

Calf Garments RCG10

Universal size up to 20” (0.51m) 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 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. 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 system use.

Indications

Intermittent Pneumatic Compression (IPC) is indicated for use of 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 calf 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. 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.

stryker®

Sustainability Solutions

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

Pack contains one Pair Compression Sleeves
Non-Sterile

DO NOT Reuse | Consult Instruction for Use | Date of Manufacture

For use with Restep Pump RSP101 only.

Made in the Dominican Republic

RCS Rev. D 09-2012 RM705000

Restep DVT System

Warranty

Reprocessed Products

Stryker warrants all reprocessed products, subject to the exceptions provided herein, to be free from defects in reprocessing and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for one use in accordance with the instructions for use of such product.

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.

Products for which Stryker is the Original Manufacturer

Stryker warrants all products for which it is the original manufacturer, subject to the exceptions provided herein, to be free from defects in design, materials and workmanship and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for a period of one year from the date of purchase.

General Warranty Terms Applicable to All Products

TO THE FULLEST EXTENT PERMITTED BY LAW, THE EXPRESS WARRANTY SET FORTH HEREIN IS THE ONLY WARRANTY APPLICABLE TO THE PRODUCTS AND IS EXPRESSLY IN LIEU OF ANY OTHER WARRANTY BY STRYKER, EXPRESSED OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, ANY IMPLIED WARRANTY OR MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. IN NO EVENT WILL STRYKER'S LIABILITY ARISING IN CONNECTION WITH THE SALE OF THE PRODUCT (WHETHER UNDER THE THEORIES OF BREACH OF CONTRACT, TORT, MISREPRESENTATION, FRAUD, WARRANTY, NEGLIGENCE, STRICT LIABILITY OR ANY OTHER THEORY OF LAW) EXCEED THE PURCHASE PRICE, CURRENT MARKET VALUE OR RESIDUAL VALUE OF THE PRODUCTS, WHICHEVER IS LESS. STRYKER SHALL NOT BE LIABLE FOR INDIRECT, SPECIAL, INCIDENTAL, PUNITIVE OR CONSEQUENTIAL DAMAGES RESULTING FROM ANY BREACH OF WARRANTY OR UNDER ANY OTHER LEGAL THEORY.

This warranty shall apply only to the original end-user purchaser of products directly from Stryker or a Stryker authorized distributor. This warranty may not be transferred or assigned without the express written consent of Stryker.

This warranty does not apply to: (1) products that have been misused, neglected, modified, altered, adjusted, tampered with, improperly installed or refurbished; (2) products that have been repaired by any person other than Stryker personnel without the prior written consent of Stryker; (3) products that have been subjected to unusual stress or have not been maintained in accordance with the instructions in the user manual or as demonstrated by a Stryker representative; (4) products on which any original serial numbers or other identification marks have been removed or destroyed; or (5) products that have been repaired with any unauthorized or non-Stryker components.

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.

If Stryker determines in its reasonable discretion that the claimed defect or non-conformance in the product is excluded from warranty coverage as described hereunder, it will notify the customer of such determination and will provide an estimate of the cost of repair of the product. In such an event, any repair would be performed at Stryker's standard rates.

Products and product components repaired or replaced under this warranty continue to be warranted as described herein during the initial applicable warranty period or, if the initial warranty period has expired by the time the product is repaired or replaced, for thirty (30) days after delivery of the repaired or replaced product. When a product or component is replaced, the item provided in replacement will be the customer's property and the replaced item will be Stryker's property. 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.

Restep®

DVT System

Bariatric Calf Garment RBG60

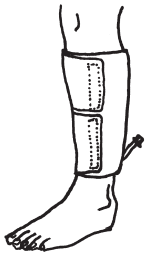
Size 20” to 26” (0.51 to 0.66m) calf circumference

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 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. 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 calf 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.

stryker®

Sustainability Solutions

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

Pack contains one Pair Compression Sleeves
Non-Sterile

DO NOT Reuse | Consult Instruction for Use | Date of Manufacture

For use with Restep Pump RSP101 only.

Made in the Dominican Republic

RCS Rev. D 09-2012 RM705001

Restep DVT System

Warranty

Reprocessed Products

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Restep®

DVT System

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.

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Sustainability Solutions

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

Pack contains one Pair Compression Sleeves
Non-Sterile

DO NOT Reuse | Consult Instruction for Use | Date of Manufacture

For use with Restep Pump RSP101 only.

Made in the Dominican Republic

RCS Rev. D 09-2012 RM705003

Restep DVT System

Warranty

Reprocessed Products

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Restep®

DVT System

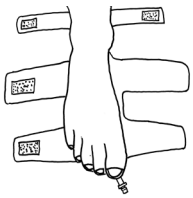
Foot Garment RFG210

Universal size

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

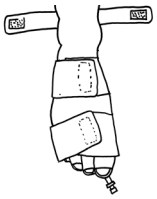
Instructions for use

Secure the pump at the foot of the bed or in place of use. Attach the tube set to the pump 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 foot. Unfold the garment(s) and position the inflatable center bladder directly below the arch of the patient's foot.

SNUGLY wrap the garment(s) around the patient's foot and secure the hook tabs. Wrap the ankle straps around the back of the ankle and on top the foot.



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. Foot garments should be removed prior to patient ambulation or transportation.

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 FOOT GARMENT. If not, press the TYPE button until the 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.

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 report any problems to the nursing staff.

The system should be used continuously until the person is fully ambulatory. Foot garments should be removed before ambulating.

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.

stryker®

Sustainability Solutions

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

Pack contains one Single Compression Sleeve
Non-Sterile

DO NOT Reuse | Consult Instruction for Use | Date of Manufacture

For use with Restep Pump RSP101 only.

Made in the Dominican Republic

RCS Rev. C 04-2012 RM705002

Restep DVT System

Warranty

Reprocessed Products

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This warranty shall apply only to the original end-user purchaser of products directly from Stryker or a Stryker authorized distributor. This warranty may not be transferred or assigned without the express written consent of Stryker.

This warranty does not apply to: (1) products that have been misused, neglected, modified, altered, adjusted, tampered with, improperly installed or refurbished; (2) products that have been repaired by any person other than Stryker personnel without the prior written consent of Stryker; (3) products that have been subjected to unusual stress or have not been maintained in accordance with the instructions in the user manual or as demonstrated by a Stryker representative; (4) products on which any original serial numbers or other identification marks have been removed or destroyed; or (5) products that have been repaired with any unauthorized or non-Stryker components.

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.

If Stryker determines in its reasonable discretion that the claimed defect or non-conformance in the product is excluded from warranty coverage as described hereunder, it will notify the customer of such determination and will provide an estimate of the cost of repair of the product. In such an event, any repair would be performed at Stryker's standard rates.

Products and product components repaired or replaced under this warranty continue to be warranted as described herein during the initial applicable warranty period or, if the initial warranty period has expired by the time the product is repaired or replaced, for thirty (30) days after delivery of the repaired or replaced product. When a product or component is replaced, the item provided in replacement will be the customer's property and the replaced item will be Stryker's property. 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.