



Instructions for Use

Description

The Respendial Inflation Device is an ultra-high pressure, large volume inflation device used to inflate, monitor pressure, and deflate angioplasty balloon dilatation catheters. It is a one-piece, 30 ml disposable inflation device rated for 26 atm with a lever-lock design that controls the piston, a manometer, and a high-pressure connecting tube with a male Luer rotating adapter. Also enclosed is a high-pressure 3-way stopcock to aid in preparation and use of the device. The manometer measures pressures ranging from 0 atm up to 26 atm in 1 atm increments. The accuracy of the manometer has been determined to be within ± 1.6 atm. These products are not made with natural rubber latex.

How Supplied

The Respendial Inflation Device is supplied sterile unless the package has been opened or damaged. For single use only. Do not reuse. Do not resterilize.

Indications for Use

The Respendial Inflation Device is indicated for use with angioplasty balloon dilatation catheters to create and monitor the pressure in the angioplasty balloon dilatation catheter and to deflate the angioplasty balloon dilatation catheter.

Contraindications

None Known

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Warnings

1. Contents supplied STERILE using ethylene oxide (EO). Non-Pyrogenic. Do not use if sterile barrier is opened or damaged. Single patient use only. Do not reuse, reprocess or re-sterilize.
2. This device has been designed for single use only. Reusing this medical device bears the risk of cross-patient contamination as medical devices – particularly those with long and small lumina, joints, and/or crevices between components – are difficult or impossible to clean once body fluids or tissues with potential pyrogenic or microbial contamination have had contact with the medical device for an indeterminable period of time. The residue of biological material can promote the contamination of the device with pyrogens or microorganisms which may lead to infectious complications.
3. Do not resterilize. After reesterilization, the sterility of the device is not guaranteed because of an indeterminable degree of potential pyrogenic or microbial contamination which may lead to infectious complications. Cleaning, reprocessing and/or reesterilization of the present medical device increases the probability that the device will malfunction due to potential adverse effects on components that are influenced by thermal and/or mechanical changes.
4. After use, this device may be a potential biohazard. Handle and dispose of in accordance with acceptable medical practices and applicable local laws and regulations.
5. To reduce the potential risk of air embolism, never use air or other gaseous medium to inflate angioplasty balloon dilatation catheters. Ensure all air has been purged from the entire fluid path prior to patient use.

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6. Do not exceed 26 atm when inflating the device. Damage to the device or user injury may result.
7. Refer to the angioplasty balloon dilatation catheter instructions for use for additional warnings.

Precautions

1. Carefully inspect the device prior to use to verify that it has not been damaged during shipment. Do not use if device damage is evident.
2. Discontinue use of the device if damage, malfunction or contamination is suspected during use.
3. For experienced physician use only.
4. Refer to the angioplasty balloon dilatation catheter instructions for use for additional precautions.

Handling and Storage

Store in a cool, dry, dark place.

Equipment Required

- Contrast Medium
- Sterile Normal Saline Solution

Operation

The plunger is free to move when the lever is pushed. In this position you may pull the plunger to aspirate or push it to inject for a quick fill. Release the lever to lock the plunger. To add and hold pressure in increments, turn the handle clockwise. Press the lever and pull back on the handle to deflate.

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Preparation

1. Prepare a solution of contrast medium and normal saline in a small sterile bowl. Check angioplasty balloon dilatation catheter instructions for specific contrast mixture recommendations or volume requirements.
2. Orient the tubing downward into the contrast medium and saline solution.
3. Ensure that the stopcock is open towards the inflation device. Push the lever and pull the plunger back to aspirate the contrast and saline solution into the syringe. Release the lever to lock the plunger.
4. Hold the device upright to purge the air from the syringe, connecting tube and stopcock. With the lever in the up position, turn the handle clockwise to expel any air bubbles.
5. Inspect the syringe, tubing and stopcock to ensure that the device has been completely purged of air bubbles. Tap the syringe lightly, if necessary, to remove all the air bubbles.
6. If necessary, adjust the syringe volume to the desired amount by turning the handle clockwise to expel contrast medium and saline solution. If more solution is needed, place the stopcock into the bowl of contrast medium and saline solution and draw up additional solution as described in steps 3 through 5 above. Close the stopcock.

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Attaching the inflation device to the angioplasty balloon dilatation catheter

1. Prepare and test the angioplasty balloon dilatation catheter according to the manufacturer's instructions for use.
2. If a separate syringe was used to prepare the angioplasty balloon dilatation catheter, remove it. Create a fluid-fluid connection between the catheter hub connector and the connecting tube of the inflation device by opening the stopcock and placing a drop of contrast medium and saline solution from the syringe into the catheter hub.
3. Hand-tighten the hubs securely. Do not over tighten the connection as this may damage the catheter hub or connecting tube

Balloon inflation and deflation

1. With the lever in locked position, turn the handle clockwise to inflate the balloon to the desired pressure.
2. Alternatively, press the lever and push the handle forward to quickly fill the angioplasty balloon dilatation catheter.
3. To deflate the balloon, rotate the handle counter-clockwise to 0 atm prior to pressing the lever. Once 0 atm has been reached, press the lever and pull back on the handle to evacuate the balloon entirely.

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Warranty

Maad Zist Co. warrants that this product was manufactured according to applicable standards and specifications. Patient condition, clinical treatment, and product maintenance may affect the performance of this product. Use of this product should be in accordance with the instructions provided and as directed by the prescribing physician.



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