



PRESTO™

Inflation Device

40
ATM

Inflating to Ultra-High Atmospheres

SINGLE solution INFLATOR

Highest Pressure Inflation Device on the Market* - **Up to 40 ATM**

Inflate to Both **Low and Ultra-High Pressures**

Inflation Device Specifications**			
Manufacturer	Device Name	Rated Inflation Pressure (ATM)	Syringe Volume (cc)
Bard	PRESTO™	40	30
Bard	CALIBER™	30	30
Boston Scientific	Encore™ 26	26	20
B Braun	AID™	30	25
Merit Medical	BasixCOMPAK™	30	20
Merit Medical	BasixTOUCH™	35	30



	PRESTO™	CALIBER™	Encore™ 26	AID™	BasixCOMPAK™	BasixTOUCH™
40 ATM Rated Pressure**	✓					
30 cc Syringe**	✓	✓				✓
17" Tubing Length†	✓					
Easy to Use††	✓	✓		✓		✓
Ergonomic††	✓	✓		✓		✓

* As of January 2015 for 30 cc inflation devices

** Information taken directly from each manufacturer's product brochure and IFU.

† 30 Bard PRESTO™ tested. Test data on file.

†† Score of 3 (defined as "acceptable") or better out of 5 in bench user evaluation. N=5, samples were evaluated twice for a total of 10 variable data points. Test data on file. Bench data may not be representative of clinical outcomes.

fast AND ADAPTABLE

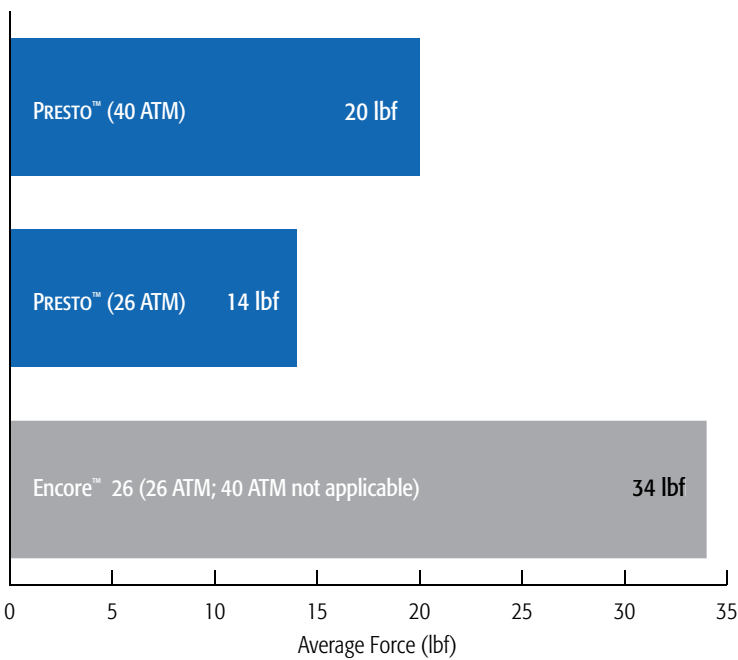
Ergonomic Design Allows for Comfortable Handling

Large Barrel Allows for **Rapid and Easy Deflation**

Designed to Inflate Large or Small Balloons with a **Single Fill of Device**



PRESTO™ Inflation Device Required Less Force at Both 26 ATM and 40 ATM than Boston Scientific Encore™ 26 at 26 ATM to Press Release Mechanism^Δ
(Less is Better)



^Δ 30 Bard Presto™ and 10 Boston Scientific Encore™ tested. Test data on file. Bench data may not be representative of clinical outcomes.

The Only Duo Designed
and Rated Up To 40 ATM -
**PRESTO™ Inflation Device &
CONQUEST® 40 PTA Dilatation Catheter**



Product Code	Description	Qty
ID4030	Presto Inflation Device	5/Box

REPRESENTATIVE'S NAME
CONTACT PHONE NO.

PHYSICIAN'S SIGNATURE

PRESTO™ Inflation Device

Indications for Use:

The PRESTO™ 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.

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 resterilization, 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 resterilization 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, state and federal 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. **6)** Do not exceed 40 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.

CONQUEST® 40 PTA Dilatation Catheter

Warning: Do not exceed the RBP recommended for this device. Balloon rupture may occur if the RBP rating is exceeded. To prevent over pressurization, use of a pressure monitoring device or indicated syringe is recommended.

Please consult product labels and instructions for use, indications, contraindications, hazards, warnings, and precautions.

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