

High Pressure. Uniform Ureteral Dilation.

Dual Lumens

- Balloon lumen is fitted with a stopcock for connection to an inflation device
- Distal lumen is designed to pass a guidewire up to 0.035"
- Distal tip can be used for infusion of contrast medium

Balloon

- 17 ATM maximum burst pressure
- 12 18 Fr options; 4 cm length
- Features two radiopaque markers for fluoroscopic visualization and more accurate positioning

Kit Configurations

- BD40 kits include screw syringe with a manometer
- BD41 kits include screw syringe with luer-lock
- In-Ka[®] balloon catheters are single-use and packaged sterile

For ordering and additional information

please contact your sales representative or Coloplast Interventional Urology Surgical Support at **800-258-3476**



Ordering Information

ltem	Catheter Diameter	Catheter Length	Balloon Diameter	Balloon Length	Inflation Device	Sales UOM	EA/Sales UOM
BD4044	5 Fr	75 cm	12 Fr	4 cm	Screw Syringe with Manometer	ΚT	1
BD4045	5 Fr	75 cm	15 Fr	4 cm	Screw Syringe with Manometer	ΚT	1
BD4046	5 Fr	75 cm	18 Fr	4 cm	Screw Syringe with Manometer	ΚT	1
BD4144	5 Fr	75 cm	12 Fr	4 cm	Screw Syringe with Luer Lock	EA	1
BD4145	5 Fr	75 cm	15 Fr	4 cm	Screw Syringe with Luer Lock	EA	1
BD4146	5 Fr	75 cm	18 Fr	4 cm	Screw Syringe with Luer Lock	EA	1

Coloplast Interventional Urology Surgical Support 800-258-3476

Indications:

In-Ka® ureteral balloon dilatation catheters are intended for:

Dilation of ureteral meatus and/or ureteral canal during endoscopic procedures

Contraindications:

Any contraindication to endoscopy (untreated urinary tract infection)

• Do not use when, in the judgement of the physician, such a procedure would be contrary to the best interest of the patient

Warnings and Precautions:

Treatment of ureteral stenosis

- Never inflate the balloon with air or gas
- Do not exceed the maximum rated burst pressure (17ATM)
- Do not inflate balloon catheter while the balloon is directly beside and in contact with a stone
- Do not resterilize this product
- The risks and benefits of using In-Ka® Ureteral Balloon Catheters should be considered in patients

Adverse Events:

- As with any dilation procedure, the use of a dilatation balloon may be associated with several risks including, but not limited to:
- Hematuria
- Damage to the urinary tract (tissue trauma, ureteral perforation), particularly if the instructions for use have not been complied with, and especially if the catheter has been positioned without fluoroscopic control

The information provided is not comprehensive with regard to product risks. For a comprehensive listing of indications, contraindications, warnings and precautions refer to the product's Instructions for Use. Alternatively, you may contact a Coloplast representative at 1-800-258-3476 and/or visit the company Website at www.coloplast.us.

Complications from the use of this device should be brought to the attention of your Coloplast Representative and your physician.

For Rx Only

Caution: Federal (USA) law restricts this device to sale by or on the order of a physician. Minneapolis, MN 04/23/2020

Ostomy Care / Continence Care / Wound & Skin Care / Interventional Urology

