Orchestra™

Hydrophilic Nitinol Guidewires

Nitinol Shaft and Core

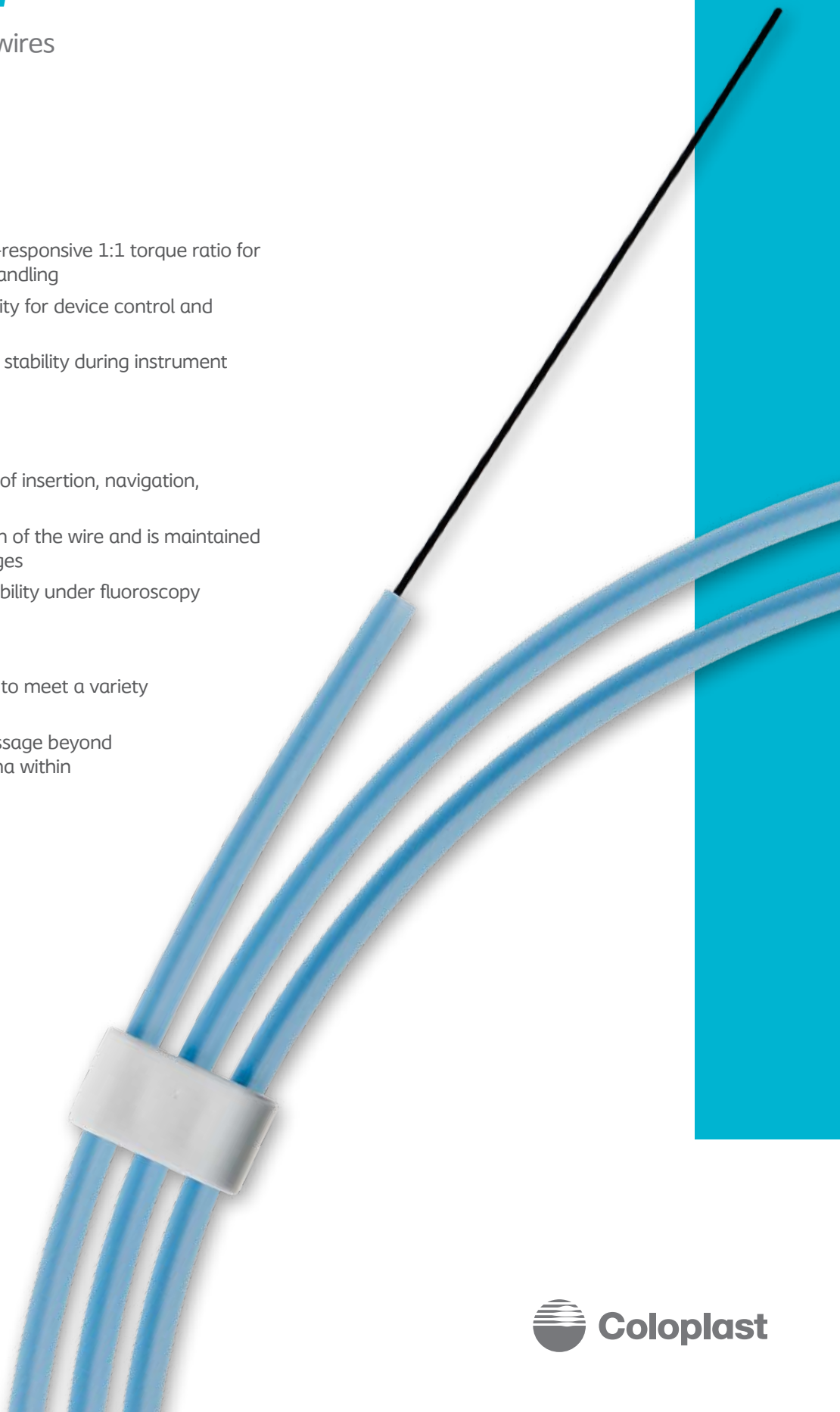
- Nitinol core design provides ultra-responsive 1:1 torque ratio for enhanced maneuverability and handling
- Ideal blend of stiffness and flexibility for device control and atraumatic navigation
- Elastic core provides support and stability during instrument advancement and exchange

Hydrophilic Outer Coating

- Reduces surface friction for ease of insertion, navigation, and withdrawal
- Coating exists on the entire length of the wire and is maintained after multiple instrument exchanges
- Outer jacket is radiopaque for visibility under fluoroscopy

Tip Talk

- Variety of available sizes and tips to meet a variety of procedural needs
- 3 cm flexible tip facilitates the passage beyond obstructions and minimizes trauma within urinary tract



Coloplast

Ordering Information

Coloplast Interventional Urology Service Support **800-258-3476**

Diameter	Length	Shaft	Tip shape	Quantity	Order Number
.035"	150 cm	Standard	Straight	Box of 5	**AEAD35**
.035"	150 cm	Standard	Angled	Box of 5	**AEAE35**
.035"	150 cm	Stiff	Straight	Box of 5	**AECD35**
.035"	150 cm	Stiff	Angled	Box of 5	**AECE35**

Indications: The hydrophilic guidewire is intended to facilitate the placement of devices through the urinary tract during endourologic procedures.

Contraindications: The hydrophilic guidewire is not intended for use other than for endourologic procedures.

Warnings: This type of instrument must only be used by trained and experienced professionals. Single Use Only. Reuse of this single use product may create a potential risk to the user.

Precautions: Excessive force against resistance may result in damage of the guidewire and/or damage of the urinary tract. Never advance the guidewire without first determining the reason for resistance. Never advance the guidewire without use of fluoroscopic guidance (or equivalent). Hydrophilic guidewires must be kept hydrated for the duration of the procedure.

Adverse Effects: Potential complications associated with device use may include perforation of the ureter, lesions of the urinary tract, trauma or shearing of the hydrophilic guidewire if the operating procedure and the warnings outlined in the instructions for use are not observed.

See the device manual for detailed information regarding the implant procedure, contraindications, warnings, precautions, and potential complications/adverse events. For further information, call Coloplast Corp at 1-800-258-3476 or consult the company website at www.coloplast.us

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

Ostomy Care
Continence Care
Wound & Skin Care
Interventional Urology

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Coloplast Corp. Minneapolis, MN 55411 / Interventional Urology Surgical Support 1-800-258-3476

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