Cardiovascular Catalog - U.S.

Cordis®
A Cardinal Health company

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Cordis is a worldwide leader in the development and manufacture of interventional vascular technology with a more than 50-year history of delivering pioneering products to treat millions of patients.

Our focus is in cardiology and endovascular platforms, with high-quality products such as diagnostic and interventional catheters, balloons, self-expanding stents, guide wires, and vascular closure devices. Working with our customers, we identify solutions that provide the best possible results for both physicians and patients.

We will continue to build on our rich history as part of Cardinal Health, a company with complementary skills and expertise. By leveraging Cordis' deep experience in product innovation and Cardinal Health's business and operational expertise, we are uniquely positioned to continue meeting the evolving needs of our customers and their patients – and remain at the forefront of change in healthcare.

To learn more visit cordis.com

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CORDIS® Access Portfolio

AVANTI®+ Sheath Introducer

The Cordis AVANTI®+ Sheath Introducer is the pioneer of catheter sheath introducer technology. Featuring a patented SLIX™ Valve, the AVANTI®+ introducer provides smooth transitions, monitoring capabilities and exceptional performance for your procedural success, as well as:

- A hexacuspid design that provides a balance between catheter maneuverability and hemostasis.
- A rotating suture collar that facilitates procedural flexibility. It stays in place and allows patient movement.
- A kink-resistant cannula design integrating a soft, flexible inner layer with a stiffer outer layer for excellent bendability and support. It is also cost-effective and reliable.
- Atraumatic tip transitions for both the sheath and the vessel dilator are uniquely tapered and manicured. This results in smooth insertions and helps to minimize damage upon entry.

Obturators

- Flexible shaft helps prevent kinking of the sheath when the sheath is used in a patient for a long period of time without a product in place.
- Easy French size identification provided by a color coding system and the number on the hub*.
- Easy to store, environment friendly boxes with the smallest possible dimensions.
- Ease of insertion through the sheath and minimal vessel damage thanks to the rounded atraumatic distal tip.
- No "backing out" of the obturator by the secure snap fit between obturator and sheath.
- Low profile hub to ease placement and manipulation.

Vessel Dilators

- Secure snap to prevent "backing out" of the dilator.
- Easy insertion due to lubricious SLX™ coating.
- Minimal tissue trauma due to optimized tapering.

	Standard length	Mid-length
Cannula length (cm)	11	23
Dilator length		
4F - 5F	18	28
6F - 9F	19	28
10F - 11F	20	30
Stopcock	3-way	3-way
Sideport Extension length (cm)	22	22
Obturator length	12	26
Suture collar	Rotating color coded suture collar	

* 4F-red, 5F-grey, 6F-green, 7F-orange, 8F-blue, 9F-black, 10F-magenta, 11F-yellow.

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With Mini-Guidewire

Without Mini-Guidewire

Key Features

- Sheath assembly
- Flexible and kink-resistant cannula
- Sideport extension with 3-way stopcock
- Flexible dilator
- Units per package: 5

For information on indications, contraindications, warnings, and precautions, see page 79.

AVANTI®+ Sheath Introducer

French Size (F)	Color Code	Cannula Usable Length (cm)	Mini-Guidewire Diameter	With Mini-Guidewire	Without Mini-Guidewire
4		11	.035	504604X	504604A
5		11	.038	504605X	504605A
5.5MS		11	.038	504655X	
6		11	.038	504606X	504606A
6.5MS		11	.038	504656X	504656A
7		11	.038	504607X	504607A
7.5MS		11	.038	504657X	504657A
8		11	.038	504608X	504608A
8.5MS		11	.038	504658X	
9		11	.038	504609X	504609A
10		11	.038	504610X	504610A
11		11	.038	504611X	

AVANTI®+ Mid-Length Sheath Introducer

French Size (F)	Color Code	Cannula Usable Length (cm)	Dilator Usable Length (cm)	With Obturator	Without Obturator
4		23	30		504604T
5		23	30		504605T
6		23	30	504606D	504606T
6.5MS		23	30		504656T
7		23	30	504607D	504607T
8		23	30	504608D	504608T
10		23	30		504610T
11		23	30	504608X	504611T

MS = Monitoring sheath. Provides slightly larger ID for pressure monitoring or side port aspiration with a catheter in place; they are one-half French.
For information on indications, contraindications, warnings, and precautions, see page 79.

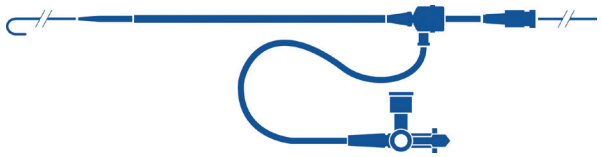
AVANTI®+ Valveless Portless Sheath Introducer With Mini-Guidewire

Key Features

- Valveless, portless
- Sheath assembly with female Luer connector hub, 45 cm mini-guidewire and tapered vessel dilator
- 11 cm usable cannula length
- 17 cm usable guidewire length
- Straight
- Units per package: 5

French Size (F)	Color Code	Cannula Usable Length (cm)	Mini-Guidewire Diameter (inch)	Product Code
7		11	.038	504607V
8		11	.038	504608V

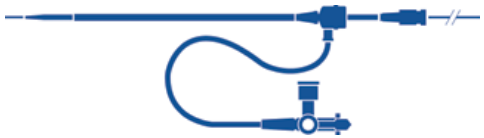
AVANTI®+ Brachial Sheath Introducer With Mini-Guidewire



French Size (F)	Color Code	Cannula Usable Length (cm)	Mini-Guidewire Diameter (inch)	Product Code
4		5.5	.035	504604P
5		5.5	.038	504605P
6		5.5	.038	504606P

For information on indications, contraindications, warnings, and precautions, see page 79.

AVANTI®+ Sheath Introducer With Mini-Guidewire



French Size (F)	Color Code	Cannula Usable Length (cm)	Mini-Guidewire Diameter (inch)	Product Code
4		7.5	.021	504604S
5		7.5	.021	504605S
6		7.5	.021	504606S

AVANTI®+ Transradial Sheath Introducer Kit

With mini-guidewire, vessel dilator and 21G needle

French Size (F)	Color Code	Cannula Usable Length (cm)	Guidewire Acceptance (inch)	Product Code
4		11	.021 70 cm	504614Z
4		23	.021 70 cm	504624Z
5		11	.021 70 cm	504615Z
5		23	.021 70 cm	504625Z
6		11	.021 70 cm	504616Z
6		23	.021 70 cm	504626Z
7		11	.021 70 cm	504617Z

With Mini-Guidewire

Without Mini-Guidewire

For information on indications, contraindications, warnings, and precautions, see page 79.

Obturators

Key Features

- For use with Sheath Introducers
- Flexible shaft helps prevent kinking of the sheath
- Easy French size identification
- Low profile hub to ease placement and manipulation
- Units per package: 10

French Size (F)	Color Code	13 cm Cannula
4		502188
5		502190
6		502191
7		502192
8		502194

Vessel Dilators

Key Features

- Facilitate the percutaneous entry of angiographic catheters
- With Luer hubs
- Units per package: 10

French Size (F)	Color Code	Guidewire Compatibility (inch)	Usable Length (cm)	Product Code
4		.021	14.5	504504S
4		.035	17	504404X
5		.021	14.5	504505S
5		.035	17	504405X
5		.038	17	504505X
6		.032	17	504406X
6		.038	17	504506X
7		.035	19	504407X
7		.038	17	504507X
8		.035	19	504408X
8		.038	17	504508X
9		.038	19	504509X

CORDIS® Diagnostic Portfolio

EMERALD® Guidewires

PTFE pre-coated guidewire
- Uniform surface finish
- Minimal insertion and withdrawal force

Product Benefits
- Easy navigation
- Versatility of tip shapes and flexibilities
- Excellent maneuverability
- Finger straightenability

Built-in safety
- Solid tensile strength minimizes the likelihood of stretching or fracturing
- Safety ribbon is welded to both ends of the wire to help it remain intact in the event of a fracture or stretching

Learn more on page 12.

7F HIGHFLOW™ Catheter

HIGHFLOW™ Catheters, available in 7F, feature polyurethane (DUCOR®) construction and a large inner diameter to maximize contrast flow.

Two-stage polyurethane construction:

- Body (Stage 1): Braided polyurethane for maximum flow and handling
- Proximal/Distal tip (Stage 2): Softer non-braided polyurethane for maximum flexibility, shape retention, and atraumatic tip

Learn more on page 19.

INFINITI® Diagnostic Catheter

The Cordis INFINITI® 4F, 5F and 6F line of diagnostic catheters is ideal for coronary angioplasty. These catheters incorporate proprietary Vestan Nylon to deliver exceptional responsiveness and flow rates, optimal torque, and shape retention.

- The TRUELUMEN Design: Provides the same inner lumen diameter from hub to tip, which eliminates contrast jetting and allows for smoother flow and excellent catheter stability.
- Large Inner Lumen: Thin-wall technology allows for larger inner lumen diameter, thus facilitating easy injections and higher flow rates.
- Radiopaque tip: Helps reduce the risk of vascular damage upon entering tortuous or fragile vessels.

Learn more on page 19.

SUPER TORQUE® Plus Diagnostic Catheter

The SUPERTORQUE® Plus line of diagnostic catheters are used in coronary angioplasty. These catheters are constructed from polyurethane to deliver exceptional responsiveness and flow rates, optimal torque, and shape retention. The SUPER TORQUE® Plus Catheter contains a soft tip which differentiates it from other SUPERTORQUE® Catheters.

- Braiding construction allows 1:1 torque control and excellent pushability without compromising kink resistance.
- Excellent visibility: Radiopaque tip improves visibility to help reduce the risk of vascular damage upon entering tortuous or fragile vessels.
- High flow rate: Thin wall technology allows for a larger inner lumen diameter, facilitating easy injections and higher flow rates.
- Custom shapes may also be available upon request.

Learn more on page 19.

EMERALD® Diagnostic Guidewires

Intended for percutaneous entry and guidance of catheters, the Cordis EMERALD® Diagnostic Guidewire complements our diagnostic catheter and catheter sheath introducer lines. Performance, endurance and safety are built into each EMERALD® Guidewire with solid tensile strength to minimize the likelihood of stretching or fracturing.

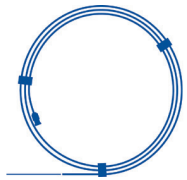
Key Features

- Excellent maneuverability, and reduced likelihood of flaking
- Finger straightenability and precise tolerances
- Proprietary PTFE Coating Process
- Available in multiple shapes and sizes

EMERALD® Fixed-Core Guidewire

Key Features

- For percutaneous entry and guidance of angiographic catheters
- Units per package: 5



PTFE-Coated Straight Tip

Diameter (inch)	Length (cm)	Flexible End (cm)	Product Code
.021	150	7	502703
.025	150	7	502549
.032	150	7	502548
.035*	150	7	502542
.035	150	10	502544
.035*	150	20	502560
.035	150	3	502542E
.038	150	7	502541

* With floppy tip

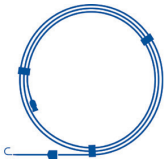
For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Fixed-Core Guidewire - PTFE-Coated Double-Ended

- 3 cm straight / 7 cm J-curve

Diameter (inch)	Length (cm)	Flexible End (cm)	Flexible End J-Curve (cm)	J-Radius (mm)	Product Code
.035	150	3	7	2	502563

EMERALD® Fixed-Core Guidewire - PTFE-Coated J-Curved



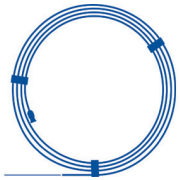
Diameter (inch)	Remarks	Length (cm)	Flexible End (cm)	J-Radius (mm)	Product Code
.025		150	7	3	502524
.025		150	7	15	502536
.030	Heavy duty, high strength	150	7	3	502522
.032		150	7	3	502526
.035		80	7	3	502701
.035		150	7	1.5	502531
.035	High strength	175	7	1.5	502534
.035		150	7	3	502521
.035		175	7	3	502585
.035		150	7	6	502589
.035		150	7	15	502535
.035		150	10	3	502587
.035		150	10	15	502576
.035	Ex Firm Tip	150	3	3	502521E
.035	Firm Tip	150	5	3	502521F
.035	Firm Tip	175	5	3	502585F
.038	Firm Tip	150	5	3	502520F
.038		150	7	3	502520
.038		175	7	3	502584
.038		150	7	6	502588
.065		150	10	6	502530

For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Fixed-Core Exchange Guidewire

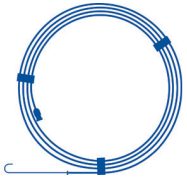
Key Features

- For use in percutaneous entry, guidance and exchange of angiographic catheters
- Units per package: 5



PTFE-Coated Straight Tip

Diameter (inch)	Length (cm)	Flexible End (cm)	Product Code
.032	260	7	502554
.035	220	7	502558
.035	260	7	502555
.038	260	7	502553



PTFE-Coated J-Curve

Diameter (inch)	Length (cm)	Flexible End (cm)	J-Radius (mm)	Product Code
.018	260	7	3	502456
.025	260	7	3	502452
.032	260	7	3	502454
.035	260	7	3	502455
.035	260	5	3	502455F
.038	260	7	3	502453

For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Fixed-Core Exchange Guidewire, *continued*

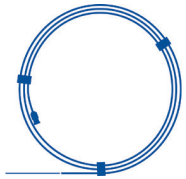
HEPARIN Coated, PTFE Coated Wires

Diameter (inch)	Length (cm)	Flexible End (cm)	J-Radius (mm)	Product Code
.038	150	7	3	502520H
.035	150	7	3	502521H
.035	150	3	3	503521H
.035	150	5	3	503521H

EMERALD® Movable-Core Guidewire

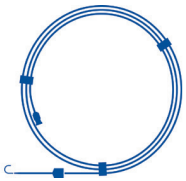
Key Features

- For use in percutaneous entry and guidance of angiographic catheters
- With 4 cm handle
- Amplatz movable core guidewires in the same configurations as below are available on request
- Units per package: 5



PTFE-Coated Straight Tip

Diameter (inch)	Length (cm)	Product Code
.035	150	502581



PTFE-Coated J-Curve

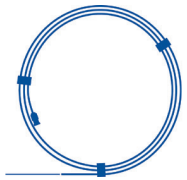
Diameter (inch)	Length (cm)	J-Radius (mm)	Product Code
.035	150	3	502571
.038	150	3	502570

For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Amplatz Guidewire

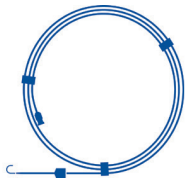
Key Features

- Exhibits little resistance through tortuous vascular curves
- For percutaneous entry and guidance of angiographic catheters
- Units per package: 5



PTFE-Coated Straight Tip

Diameter (inch)	Length (cm)	Product Code
.035	150	502581A



PTFE-Coated J-Curve

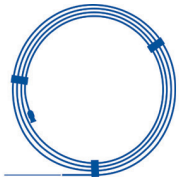
Diameter (inch)	Length (cm)	J-Radius (mm)	Product Code
.035	150	3	502571A

For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Amplatz Super Stiff Guidewires

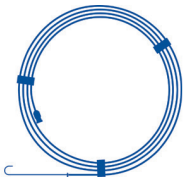
Key Features

- For percutaneous entry and guidance of angiographic catheters
- Units per package: 5



PTFE-Coated Straight Tip

Diameter (inch)	Length (cm)	Product Code
.035	260	502442E
.035	150	502726
.035	180	502728



PTFE-Coated J-Curve

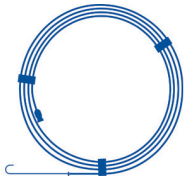
Diameter (inch)	Length (cm)	J-Radius (mm)	Product Code
.035	150	3	502731
.035	260	3	502735

For information on indications, contraindications, warnings, and precautions, see page 81.

EMERALD® Rosen Heavy Duty Guidewire

Key Features

- For percutaneous entry and guidance of angiographic catheters
- Intermediate level of body stiffness
- Highly atraumatic tip
- Maintains purchase in short vessel segments
- Units per package: 5



PTFE-Coated J-Curve

Diameter (inch)	Length (cm)	J-Radius (mm)	Product Code
.035	180	1.5	502717

EMERALD® Standard J-Tip Guidewire

Key Features

- For Biopsy Forceps procedures
- Fixed core
- Units per package: 5

PTFE-Coated J-Curve

Diameter (inch)	Length (cm)	Flexible End (cm)	J-Radius (mm)	Product Code
.035	80	7	3	502701

For information on indications, contraindications, warnings, and precautions, see page 81.

INFINITI® Diagnostic Catheter, 7F HIGHFLOW™ Diagnostic Catheter, SUPER TORQUE® Plus Diagnostic Catheter, and TEMPO AQUA® Diagnostic Catheter

Left Coronary Judkins Technique (100 cm)

- For percutaneous entry and guidance of angiographic catheters
- Units per package: 5

		4F	5F	6F	5.2F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
JL 3.5	Judkins Left 3.5	538418	534518T	534618T	533551	533618	527718
JL 4	Judkins Left 4	538420	534520T	534620T	533553	533620	527720
JL 4.5	Judkins Left 4.5	538417	534517T	534617T	533527	533627	
JL 5	Judkins Left 5	538422	534522T	534622T	533559	533622	527722
JL 6	Judkins Left 6	538424	534524T	534624T	533561	533624	527724

JL 3.5 JL 4 JL 4.5 JL 5 JL 6

Left Coronary Amplatz Technique (100 cm)

- Units per package: 5

		4F	5F	6F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
AL 1	Amplatz Left 1	538445	534545T	534645T	533645	527740
AL 2	Amplatz Left 2	538446	534546T	534646T	533646	527741
AL 3	Amplatz Left 3	538447	534547T	534647T	533647	

AL 1 AL 2 AL 3

For information on indications, contraindications, warnings, and precautions, see page 83.

Right Coronary Judkins Technique (100 cm)

		4F	5F	6F	5.2F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
JR 3.5	Judkins Right 3.5	538419	534519T	534619T	533550	533619	527719
JR 4	Judkins Right 4	538421	534521T	534621T	533552	533621	527721
JR 4 (125 cm)	Judkins Right 4, 125 cm			534615T	533565		
JR 4 Classic	Judkins Right 4 Classic, 100 cm					533687	
JR 4 RECESSED BRAID	Judkins Right 4 Recessed Braiding	534516T	534516T			533689	
JR 4 MOD	Judkins Right 4 Modified	538428	534528T	534628T	533528	533628	
JR 4 ST	Judkins Right 4 Short Tip	538427	534527T	534627T		533626	
JR 5	Judkins Right 5	538423	534523T	534623T	533558	533623	527723
JR 5 MOD	Judkins Right 5 Modified			534629T			
JR 6	Judkins Right 6, 125 cm	538425	534525T	534625T	533560	533625	

JR 3.5 | JR 4 | JR Classic | JR 4 MOD | JR 5 | JR 5 MOD | JR 6

Right Coronary Amplatz Technique Modified (100 cm)

		4F	5F	6F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
AR MOD	Amplatz Right Modified	538448	534548T	534648T	533648	527748
AR 1 MOD	Amplatz Right 1 Modified	538441	534541T	534641T	533641	
AR 2 MOD	Amplatz Right 2 Modified	538443	534543T	534643T	533643	

AR MOD | AR 1 MOD | AR 2 MOD

Right Coronary Shapes - Williams Technique

- Units per package: 3

		4F	5F	6F	6F
		INFINITI® Catheter	**INFINITI® Catheter**	**INFINITI® Catheter**	**SUPER TORQUE® Plus Catheter**
3 DRC	Right Coronary 3 Dimensional	538476	534576T	534676T	533676

3 DRC

Right Coronary Noto Technique

		4F	5F	6F
		INFINITI® Catheter	**INFINITI® Catheter**	**INFINITI® Catheter**
SRC	Right Coronary N.	538474	534574T	534674T

SRC

Multipurpose A Cournand Technique (open end, no sides holes)

		5.2F	6F	7F
		SUPER TORQUE® Plus Catheter	**SUPER TORQUE® Plus Catheter**	**7 F HIGHFLOW™ Catheter**
MPA 1 (80 cm)	Multipurpose A, 80 cm	533579	533633	
MPA 1	Multipurpose A Cournand, 100 cm		533640	527784
MPA 1	Multipurpose A, 100 cm	SR1924		
MPA 1 (125 cm)			533667	

MPA 1

Multipurpose A Curve (open end, 2 side holes)

		4F	5F	5.2F
		INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter
MPA 2 (I) SH (65 cm)	Multipurpose A-2, 65 cm 2 side holes	538440	534540T	
MPA 2 (I) SH (80 cm)	Multipurpose A-2, 80 cm 2 side holes	538449	534549T	
MPA 2 (I) SH (100 cm)	Multipurpose A-2, 100 cm 2 side holes	538442		533556
MPA 2 (I) SH (125 cm)	Multipurpose A-2, 125 cm 2 side holes	538444		

MPA 2 (I)

Multipurpose A Adult Curve (open end, 2 side holes)

		5F	6F	5.2F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
MPA 2 SH (65 cm)	Multipurpose A-2, 65 cm 2 side holes			533562		
MPA 2 SH (80 cm)	Multipurpose A-2, 80 cm 2 side holes				533629	
MPA 2 SH (100 cm)	Multipurpose A-2, 100 cm 2 side holes	534542T	534642T	533582	533642	527742
MPA 2 SH (125 cm)	Multipurpose A-2, 125 cm 2 side holes	534544T				527787

MPA 2

Multipurpose A Adult Curve (open end, no side hole)

		5.2F
		SUPER TORQUE® Plus Catheter
MPA 2	Multipurpose A-2, 100 cm	SR1924

MPA 2

Multipurpose B Gensini Technique (open end, 6 side holes, 100 cm)

		6F
		SUPER TORQUE® Plus Catheter
MPB 3	Multipurpose B Gensini	533634

MPB 3

Multipurpose B (open end, 2 side holes, 100 cm)

		5F	6F	6F
		INFINITI® Catheter	**INFINITI® Catheter**	**SUPER TORQUE® Plus Catheter**
MPB 2	Multipurpose B-2	534539T	534649T	533649

MPB 2

Sones Technique

		4F	5F	5.2F	6F
		INFINITI® Catheter	**INFINITI® Catheter**	**SUPER TORQUE® Plus Catheter**	**SUPER TORQUE® Plus Catheter**
SON 1 SH (80 cm)	Sones 1, 80 cm 2 side holes	538430	534530T		
SON 1	Sones 1, 100 cm		534562T		
SON 1	Sones 1, 100 cm 4 side holes		SR4827		533630
SON 1.5	Sones 1.5, 100 cm		534564T		
SON 2	Sones 2, 80 cm 2 side holes	538431			
SON 2	Sones 2, 100 cm			SR2360	533631
SON 3	Sones 3, 100 cm 4 side holes				533632

SON 1 SON 2 SON 3

Castillo Technique (80 cm)

		5.2F	6F
		SUPER TORQUE® Plus Catheter	**SUPER TORQUE® Plus Catheter**
CAS 1	Castillo 1, 100 cm	533584	533684
CAS 2 (80 cm)	Castillo 2, 80 cm 4 side holes		533682
CAS 2	Castillo 2, 100 cm	533585	533685
CAS 3	Castillo 3, 100 cm		533686

CAS 1 CAS 2 CAS 3

Barbeau Technique

		5F
		INFINITI® Catheter
BARBEAU 2 SH	Barbeau, 100 cm, 2 side holes	534578T

BARBEAU

El Gamal Technique (100 cm)

		6F
		SUPER TORQUE® Plus Catheter
EGB 1	El Gamal 1	533637

EGB 1

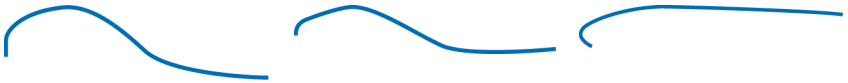
Radial Bilateral Technique and Radial/Brachial (Tilon)

		5F
		TEMPO AQUA® Catheter
RBL 4.0	Barbeau, 100 cm, 2 side holes	SRD7045
RBL 4.5	RBL 4.5, 100 cm	SRD7074
RBL 5.0	RBL 5.0, 100 cm	SRD7047
RBLA	RBL A, 100 cm (pack of 5)	SRD7075

RBL 4.0 RBL 4.5 RBL 5.0 RBMP/TILON

Coronary Bypass Techniques (100 cm)

		4F	5F	6F	6F	7F
		INFINITI® Catheter	**INFINITI® Catheter**	**INFINITI® Catheter**	**SUPER TORQUE® Plus Catheter**	**7F HIGHFLOW™ Catheter**
LCB	Left Coronary Bypass	538472	534572T	534672T	533672	527772
RCB	Right Coronary Bypass	538470	534570T	534670T	533670	527770
IM	Internal Mammary	538460	534560T	534660T	533660	527760



LCB RCB IM

Ventricular Straight Pigtail (110 cm)

		4F	5F	6F	5.2F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
PIG 4SH (125 cm)	Straight Pigtail 4 side holes				SRD5287		
PIG 5SH	Straight Pigtail 5 side holes	538451V					
PIG 6SH	Straight Pigtail 6 side holes	538450S*	534550S	534650S	533533	533650S	527750S
PIG 8SH	Straight Pigtail 8 side holes	538450E*	534550E	534650E		533650E	527750E
PIG 12SH (50cm)	Straight Pigtail 12 side holes						527750

PIG

Ventricular Angled Pigtail: Van Tassel Technique

		4F	5F	6F	5.2F	6F	7F
		INFINITI® Catheter	INFINITI® Catheter	INFINITI® Catheter	SUPER TORQUE® Plus Catheter	SUPER TORQUE® Plus Catheter	7F HIGHFLOW™ Catheter
PIG 145° MOD 5SH	Angled Pigtail 145° 5 side holes modified	538457V					
PIG 145° 6SH	Angled Pigtail 145° 6 side holes		534552S	534652S	533534A	533652S	527752S
PIG 145° MOD 6SH	Angled Pigtail 145° 6 side holes modified	538453S*	534553S	534653S			
PIG 155° 6SH	Angled Pigtail 155° 6 side holes		534554S	534654S	533533A	533654S	
PIG 155° MOD 6SH	Angled Pigtail 155° 6 side holes modified	538455S*				533655S	
PIG 155° 12SH	Angled Pigtail 155° 12 side holes						527754
PIG 155° 5 SH	Angled Pigtail 155° 5 side holes	538459V					
PIG 155° ST	Angled Pigtail 155° Short Tip		534555S*				

PIG 145°

PIG 155°

* Micro Loop

N.I.H. Paediatric Curve

		5.2F
		SUPER TORQUE® Plus Catheter
N.I.H. (65 cm)	N.I.H. 65 cm	533525
N.I.H. (80 cm)	N.I.H. 80 cm	533535
N.I.H.	N.I.H. 100 cm	533545

N.I.H

N.I.H. Adult Curve

		6F	7F
		SUPER TORQUE® Plus Catheter	**7F HIGHFLOW™ Catheter**
N.I.H. (80 cm)	N.I.H. 80 cm	533635	
N.I.H.	N.I.H. 100 cm	533636	527745

N.I.H.

Straight

		5.2F
		SUPER TORQUE® Plus Catheter
STRAIGHT (80 cm)	Straight 80cm	SR4098

Diagnostic Catheter MultiPac

Key Features

- 3 catheters JL, JR, PIG (straight or angled 145°)
- All product codes with "CP" are tailor-made packs, lead time for delivery: 6-8 weeks
- Units per package: 5 (unless otherwise noted)

4F INFINITI® Diagnostic Catheter - MultiPac

Product Code	Judkins Left 4	Judkins Right 4	Angled Pigtail 145° 5 side holes Modified
538493	538420	538421	538457V
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Angled Pigtail 155° 6 side holes Modified**
538494	538420	538421	538455S
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 8 side holes**
538498	538420	538421	538450E
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 5 side holes**
538499	538420	538421	538451V
Product Code	**Judkins Left 4**	**Right Coronary 3 Dimensional**	**Angled Pigtail 145° 6 side holes Modified**
CP0092*	538420	538476	538453S
Product Code	**Judkins Left 4**	**Right Coronary 3 Dimensional**	**Straight Pigtail 6 side holes**
CP0097	538420	538476	538450S
Product Code	**Judkins Left 5**	**Judkins Right 4**	**Angled Pigtail 145° 8 side holes**
CP0421*	538422	538421	538450E

* 10 units per package

5F INFINITI® Diagnostic Catheter - MultiPac

Product Code	Judkins Left 4	Right Coronary 3 Dimensional	Straight Pigtail 6 side holes
CP0207*	534520T	534576T	534550S
Product Code	**Judkins Left 4**	**Right Coronary 3 Dimensional**	**Angled Pigtail 145° 6 side holes**
CP0208*	534520T	534576T	534552S
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 6 side holes**
CP0388	534518T	534521T	534552S

* Units per package: 10

6F INFINITI® Diagnostic Catheter - MultiPac

Product Code	Judkins Left 4	Right Coronary 3 Dimensional	Angled Pigtail 145° 6 side holes
CP0257	534620T	534676T	534652S

5.2F SUPER TORQUE® Plus Diagnostic Catheter - MultiPac

Product Code	Judkins Left 4	Judkins Right 4	Angled Pigtail 145° 6 side holes	
533593	533553	533552	533534A	
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 6 side holes**	
533598	533553	533552	533533	
Product Code	**Judkins Left 5**	**Judkins Right 5**	**Angled Pigtail 145° 6 side holes**	
CP0313	533559	533552	533534A	
Product Code	**Judkins Left 4**	**Judkins Right 4**	**5.2F Avanti+ 11 cm .038”**	**Emerald™ .035”**
CP0401	533553	533552	504605X*	502521
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 6 side holes**	
CP0412	533551	533552	533534A	

* with mini-guidewire

6F SUPERTORQUE® Plus Diagnostic Catheter - MultiPac

Product Code	Judkins Left 4	Judkins Right 4	Angled Pigtail145° 6 side holes
533693	533620	533621	533652S
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 8 side holes**
533698	533620	533621	533650E

Diagnostic Catheter PrimoPac

Key Features

- 3 catheters JL4, JR4, PIG (S or A 145°)
- 1 sheath introducer
- All product codes with "CP" are tailor-made packs, lead time for delivery: 6-8 weeks
- Units per package: 10

4F INFINITI® Diagnostic Catheter - PrimoPac

Product Code	Judkins Left 4	Judkins Right 4	Straight Pigtail 8 side holes	4F Avanti+ 11 cm .035"
538491P	538420	538421	538450E	
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 5 side holes**	**Modified 4F Avanti+ 11 cm .035"**
538493P	538420	538421	538457V	504604X*
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Angled Pigtail 155° 6 side holes**	**Modified 4F Avanti+ 11 cm .035"**
538494P	538420	538421	538455S	504604X*
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 5 side holes**	**4F Avanti+ 11 cm .035"**
538499P	538420	538421	538451V	504604X*
Product Code	**Judkins Left 4**	**Right Coronary 3 Dimensional**	**CATH 4F ST PIG**	**110 cm 4F Avanti+ 11 cm .035"**
CP0242	538420	538476	532413T	504604X*

5.2F SUPER TORQUE® Plus Diagnostic Catheter - PrimoPac

Product Code	Judkins Right 4	Judkins Left 5	Angled Pigtail 145° 6 side holes	5F Avanti+ 11 cm .038"
CP0415**	533552	533559	533534A	504605X*

* with mini-guidewire
** 5 units per package

Diagnostic Catheter CorPac

Key Features

- 3 catheters JL4, JR4, PIG (S or A 145°)
- 1 sheath introducer
- 1 guidewire: diameter (inch) as indicated
- All product codes with "CP" are tailor-made packs, lead time for delivery: 6-8 weeks
- Units per package: 10

4F INFINITI® Diagnostic Catheter - CorPac

Product Code	Judkins Left 4	Judkins Right 4	Straight Pigtail 8 side holes	4F Avanti+ 11 cm .035"	Emerald™ .035"
538-492C	538420	538421	538450E	504604X*	502521
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 5 side holes Modified**	**4F Avanti+ 11 cm .035"**	**Emerald™ .035"**
538-493C	538420	538421	538457V	504604X*	502521
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Angled Pigtail 155° 6 side holes Modified**	**4F Avanti+ 11 cm .035"**	**Emerald™ .035"**
538-494C	538420	538421	538455S	504604X*	502521
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Straight Pigtail 5 side holes**	**4F Avanti+ 11 cm .035"**	**Emerald™ .035"**
538-499C	538420	538421	538451V	504604X*	502521
Product Code	**Judkins Left 4**	**Judkins Right 4**	**4F Avanti+ 11 cm .035"**	**Emerald™ .035"**	
CP0296	538420	538421	504604X	502521	
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Angled Pigtail 145° 6SH MOD**	**4F Avanti+ 11 cm .021"**	**Emerald™ .035"**
CP0391	538420	538421	538453S	504614Z	502521
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 6 SH MOD**	**4F Avanti+ 11 cm .021"**	**Emerald™ .035"**
CP0411	538418	538421	538453S	504614Z*	502521

* with mini-guidewire

For information on indications, contraindications, warnings, and precautions, see page 83.

5F INFINITI® Diagnostic Catheter - CorPac

Product Code	Judkins Left 4	Judkins Right 4	5F Avanti+ 11 cm .038"	Emerald™ .035"	
CP0295	534520T	534521T	504605X*	502521	
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 6 side holes**	**5F Avanti+ 11 cm .021"**	**Emerald™ .035"**
CP0395	534518T	534521T	534552S	504615Z*	502521

6F INFINITI® Diagnostic Catheter - CorPac

Product Code	Judkins Left 4	Judkins Right 4	6F Avanti+ 11 cm .038"	Emerald™ .035"
CP0406	534620T	534621T	504606X*	502521

5.2F SUPER TORQUE® Plus Diagnostic Catheter - CorPac

Product Code	Judkins Left 4	Judkins Right 4	Angled Pigtail 145° 6 side holes	5F Avanti+ 11 cm .038"	Emerald™ .038"
533-593C	533553	533552	533534A	504605X*	502520

6F SUPER TORQUE® Plus Diagnostic Catheter - CorPac

Product Code	Judkins Left 4	Judkins Right 4	6F Avanti+ 11 cm .038"	Emerald™ .035"	
CP0283	533620	533621	504606X*	502521	
Product Code	**Judkins Left 5**	**Judkins Right 5**	**Angled Pigtail 145° 6 side holes**	**6F Avanti+ 11 cm .038"**	**Emerald™ .038"**
CP0306	533622	533623	533652S	504606X*	502520
Product Code	**Judkins Left 4**	**Judkins Right 4**	**Angled Pigtail 145° 6 side holes**	**6F Avanti+ 11 cm .021"**	**Emerald™ .035"**
CP0389	533620	533621	533652S	504616Z*	502521
Product Code	**Judkins Left 3.5**	**Judkins Right 4**	**Angled Pigtail 145° 6 side holes**	**6F Avanti+ 11 cm .021"**	**Emerald™ .035"**
CP0428	533618	533621	533652S	504616Z*	502585

* with mini-guidewire

Diagnostic Catheter 2-Packs

Key Features

- 2 catheters JL, JR/3DRC
- All product codes with "CP" are tailor-made packs, lead time for delivery: 6-8 weeks

2-Packs JL and JR4/3DRC

	4F	5F	6F
JL4/JR4 2-Packs order number. Units per Package: 5. Included regular codes.	CP0276 538-420 and 538-421	CP0278 534-520T and 534-521T	CP0281 534-620T and 534-621T
JL4/3DRC 2-Packs order number. Units per Package: 3. Included regular codes.		CP0279 534-520T and 534-576T	CP0282 534-620T and 534-676T
JL4/JR4 + CSI +wire CORPAC® order number. Units per Package: 10. Included regular codes.		CP0317 534-520T; 534-521T; 502-521; 504-606X and 502-652	

2-Packs JL and JR4/3DRC

	5F	6F
JL4/JR4 2-Packs order number. Units per Package: 5. Included regular codes.	CP0280, 533-553 and 533-552	CP0285, 533-620 and 533-621

JL 4 | JR 4 | JL 4 | 3DRC | JL 3.5 | JR 4

CORDIS® Interventional Portfolio

Steerable Guidewires

ATW™ Guidewire and ATW™ Marker Wire

Moderate Support / Workhorse Wire
- Broad Transition One-Piece Corewire (.0076") for excellent linear control
- Flex-Joint bond maintains strength and flexibility of the distal corewire transition
- Duraglide proximal coating and distal PTFE sleeve provide excellent lubricity
- Flexibility with support for modern Hi-Tech stents
- Available tip configuration: floppy

Learn more on page 35 and page 36.

STABILIZER® Plus Steerable Guidewire, STABILIZER® XS Steerable Guidewire and STABILIZER® Marker Steerable Guidewire

Moderate Support / Workhorse Wire
- Broad Transition One-Piece Corewire (.010") for excellent linear control
- Flex-joint bond maintains strength and flexibility of the distal corewire transition
- Duraglide proximal coating and distal PTFE sleeve provide excellent lubricity
- Balanced support and flexibility
- Available tip configuration: supersoft, soft
- An intermediate support wire for secure lesion measurement with 6 equally distanced marker bands (15 mm)

Learn more on page 37.

SHINOBI® Guidewire

- Broad Transition One-Piece Corewire (.007") for linear control
- Flex-joint bond maintains strength and flexibility of the distal corewire transition
- Extended PTFE sleeve over tip provides superior lubricity
- Flexibility with support for modern applications
- Available tip configuration: floppy

Learn more on page 38.

SHINOBI® Plus Guidewire

- Broad Transition One-Piece Corewire (.010") for excellent linear control
- Flex-joint bond maintains strength and flexibility of the distal corewire transition
- Extended PTFE sleeve over tip provides superior lubricity
- Balanced support and flexibility
- Available tip configuration: floppy

Learn more on page 38.

REFLEX® Guidewire

Light Support Wire
- An ultra-thin .0067" core wire combined with Cordis' silicone based SLX™ coating give the REFLEX® Guidewire its outstanding crossing properties
- 25 cm distal radiopaque coil
- One-piece core construction
- Distal SLX™-lubricant

- Learn more on page 39.

WIZDOM™ Guidewire

Light Support Wire
- Broad Transition One-Piece Corewire (.007") for excellent linear control
- Flex-joint bond maintains strength and flexibility of the distal corewire transition
- Duraglide proximal coating and distal PTFE sleeve provide excellent lubricity
- Available tip configurations: supersoft, soft.

Learn more on page 40.

Guiding Catheters

ADROIT® Guiding Catheters

Intended for use for intravascular introduction of interventional/diagnostic devices into the coronary or peripheral vascular systems. Learn more on page 42.

VISTA BRITE TIP® and Long VISTA BRITE TIP® Guiding Catheters

A complete system of large lumen guiding catheters that are optimized to answer clinical needs. The performance-based design of each catheter easily meets strength, control and delivery requirements in the widest range of anatomies. Learn more on page 64.

For information on indications, contraindications, warnings, and precautions, see page 78.

CORDIS® Interventional Portfolio

PTCA Balloons

EMPIRA® RX Pre-Dilatation Catheter

A low-profile balloon to help you reach and treat the lesion with excellent crossability and recrossability. Its exceptional crossability and recrossability is achieved through the highly flexible DURALYN® Flex balloon material, the proprietary pleating and folding process, and also the lubricious as well as durable hydrophilic coating. Learn more on page 65.

EMPIRA NC® RX Pre-Dilatation Catheter

Designed to deliver in challenging interventions and post-dilatation procedures. High-pressure balloon that combines exceptional crossability and recrossability with accuracy during postdilatation. Learn more on page 66.

MOZEC™ PTCA Balloon Dilatation Catheter

As the longest PTCA balloons on the U.S. market, they are available in a broad range of sizes for greater cath lab efficiency. Learn more on page 67.

MOZEC™ NC PTCA Balloon Dilatation Catheter

A PTCA balloon dilatation catheter that combines controlled balloon growth with the longest lengths on on the U.S. market. Learn more on page 68.

Bare Metal Stents

NIRxcell™ CoCr Coronary Stent System

A bare metal stent system that includes a balloon-expandable intracoronary L-605 Cobalt Chromium (CoCr) NIRxcell stent, premounted on a rapid exchange balloon catheter (the Delivery System). Learn more on page 69.

Bifurcated Coronary Stent

TRYTON Side Branch Stent

A stent specifically designed to actively treat, protect and secure the entire bifurcation lesion, offering ease of implantation and complete main vessel stent integration. Learn more on page 70.

Drug Eluting Stents

EluNIR® Ridaforolimus Eluting Coronary Stent System

A DES that was designed to help you navigate with ease, even in highly complex anatomies. Learn more on page 71.

Cordis® Biopsy Forceps

Standard Biopsy Forceps

Choice of two forceps diameters for taking samples adapted to a wide range of clinical situations. Learn more on page 72.

BI-PAL® Biopsy Forceps

Disposable and torquable (PTFE formable/torquable shaft). Jugular or femoral access possible with two lengths of forceps. Learn more on page 73.

Note: EluNIR and NIRxcell are manufactured by Medinol and distributed by Cordis Corporation. TRYTON Side Branch Stent is manufactured by Tryton Medical and distributed by Cordis Corporation. MOZEC is manufactured by Meril and distributed by Cordis Corporation.

ATW™ Steerable Guidewire

Cordis offers a complete platform of steerable guidewires for choosing the right guidewire given a specific clinical situation.

Key Features

- Intermediate support wire
- Coating: duraglide/PTFE
- Distal tip radiopacity: 3 cm
- Tip flexibility: floppy

Units per package: 1

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	195	Straight	595014
.014	195	J-Curve	595J014
.014	300	Straight	595X014
.014	300	J-Curve	595Y014

Units per package: 5

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	195	Straight	595E014
.014	195	J-Curve	595EJ014
.014	300	Straight	595EX014
.014	300	J-Curve	595EY014

For information on indications, contraindications, warnings, and precautions, see page 78.

ATW™ Marker Wire Guidewire

An intermediate support wire for lesion measurement, with 4 radiopaque markers spaced 10 mm apart.

Key Features

- Tip flexibility: floppy

Units per package: 1

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	195	Straight	595M014
.014	195	J-Curve	595MJ014
.014	300	Straight	595MX014
.014	300	J-Curve	595MY014

Units per package: 5

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	195	Straight	595ME014
.014	195	J-Curve	595MEJ014
.014	300	Straight	595MEX014
.014	300	J-Curve	595MEY014

For information on indications, contraindications, warnings, and precautions, see page 78.

STABILIZER® Plus Steerable Guidewire

Key Features

- Super soft tip
- Radiolucent extra support steerable guidewires
- Coating: PTFE/duraglide
- Distal tip radiopacity: 3 cm
- Units per package: 5

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	507114
.014	180	J-Curve	507114J
.014	300	Straight	507114X
.014	300	J-Curve	507114Y
.014	180	Straight	507714
.014	180	J-Curve	507714J
.014	300	Straight	507714X
.014	300	J-Curve	507714Y
.014	180	Straight	507914
.014	180	J-Curve	507914J
.014	300	Straight	507914X
.014	300	J-Curve	507914Y

STABILIZER® XS Steerable Guidewire

Key Features

- Coating: PTFE/duraglide
- Distal tip radiopacity: 3 cm
- Units per package: 5

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	527914
.014	180	J-Curve	527914J
.014	300	Straight	527914X
.014	300	J-Curve	527914Y

STABILIZER® Marker Steerable Guidewire

Key Features

- Super soft tip
- Number of markers: 6
- Marker spacing: 15mm apart
- Marker width: 1.5mm
- Most distal marker: 4.5cm from tip
- Distal tip radiopacity: 3 cm
- Units per package: 5

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	518224
.014	180	J-Curve	518224J
.014	300	Straight	518224X
.014	300	J-Curve	518224Y

For information on indications, contraindications, warnings, and precautions, see page 78.

SHINOBI® Steerable Guidewire

Key Features

- Inner corewire diameter: .007″
- Shapeable
- Radiopaque: 3 cm
- Coating: PTFE
- Units per package: 1

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	547114
.014	300	Straight	547114X

SHINOBI® Plus Steerable Guidewire

Key Features

- Inner corewire diameter: .01″
- Shapeable
- Radiopaque: 3 cm
- Coating: PTFE
- Units per package: 1

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	547214
.014	300	Straight	547214X

For information on indications, contraindications, warnings, and precautions, see page 78.

REFLEX® Steerable Guidewire

An ultra-thin .0067" core wire combined with Cordis' silicone based SLX™ coating give the REFLEX® Guidewire its outstanding crossing properties.

Key Features

- Coating: PTFE & SLX™
- 25 cm distal radiopaque coil
- One-piece core construction
- Distal SLX™ lubricant
- Units per package: 5

Super Soft Tip

- Color code: Orange

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	175	J-Curve	502014CJ

Soft Tip

- Color code: Blue

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	175	Straight	502014B

Standard Tip

- Color code: Green

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	175	J-Curve	502014AJ

For information on indications, contraindications, warnings, and precautions, see page 78.

WIZDOM® Steerable Guidewire

Key Features

- Moderate support wire
- Coating: PTFE/duraglide
- Distal tip radiopacity: 3 cm
- Units per package: 5

Super Soft Tip

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	512143
.014	180	J-curve	512143J
.014	300	Straight	512143X
.014	300	J-curve	512143Y

Soft Tip

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	512142
.014	180	J-curve	512142J
.014	300	Straight	512142X

WIZDOM® Steerable Guidewire ST - Short Transition

Key Features

- Coating: PTFE/duraglide

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	180	Straight	512114
.014	180	J-curve	512114J
.014	300	Straight	512114X

For information on indications, contraindications, warnings, and precautions, see page 78.

Extension Wire - CINCH® QR

Diameter (inch)	Length (cm)	Tip Shape	Product Code
.014	145	Straight	502144

Catheter Extensions

Key Features

- Flexible, translucent polyurethane
- Braided
- Designed for 1000 psi
- Luer lock hubs
- Male-female
- Units per package: 5

Length	Inner Diameter (mm)	Product Code
45	1.8	502100D
65	1.8	502101D
100	1.8	502102D

ADROIT® Guiding Catheter

Key Features

- Improves device compatibility and provides excellent visualization
- Easier to perform kissing balloon technique
- Innovative hybrid braided wire technology enables larger lumen with optimal back-up support
- PTFE lining provides lubricious lumen for smoother delivery
- TRUELUMEN® Technology ensures consistent ID from hub to tip, for added confidence
- 5F to 6F platform, transradial shapes, and extra back-up shapes
- Available with a .058" inner diameter (ID) for 5F
- Available with a .072" inner diameter (ID) for 6F
- Long Brite Tip shapes

Judkins Left

Shape	Shape Description	5F	6F
JL 3	Judkins Left 3	55800000	67200000
JL 3.5	Judkins Left 3.5	55800200	67200200
JL 3.5 ST	Judkins Left 3.5 - Short Tip		67201200
JL 3.5 SH	Judkins Left 3.5 - 2 Side Holes		67200300
JL 4 (90 cm)	Judkins Left 4		67200490
JL 4	Judkins Left 4	55800400	67200400
JL 4 ST	Judkins Left 4 - Short Tip		67201400
JL 4 SH	Judkins Left 4 - 2 Side Holes		67200500
JL 4 (125 cm)	Judkins Left 4		67204025
JL 4 LBT	Judkins Left 4 - Long Brite Tip		6720040L
JL 4.5	Judkins Left 4.5	55800600	67200600
JL 4.5 ST	Judkins Left 4.5 - Short Tip		67201600
JL 5	Judkins Left 5	55800800	67200800
JL 6	Judkins Left 6		67201000

JL 3 JL 3.5 JL 4 JL 4.5 JL 5 JL 6

SH = Side Holes. ST = Short Tip. For information on indications, contraindications, warnings, and precautions, see page 78.

Judkins Curved Left

Shape	Shape Description	5F	6F
JCL 3.5	Judkins Curved Left 3.5	55802600	67202600
JCL 3.5 SH	Judkins Curved Left 3.5 - 2 side holes		67202700
JCL 4	Judkins Curved Left 4		67202800

JCL 3.5 JCL 3.5 SH JCL 4

Judkins Right

Shape	Shape Description	5F	6F
JR 3.5	Judkins Right 3.5	55808000	67208000
JR 3.5 SH	Judkins Right 3.5 - 2 Side Holes		67208100
JR 4 (90cm)	Judkins Right 4		67208290
JR 4	Judkins Right 4	55808200	67208200
JR 4 ST	Judkins Right 4 - Short Tip		67209000
JR 4 SH	Judkins Right 4 - 2 Side Holes		67208300
JR 4 (125cm)	Judkins Right 4		67208225
JR 4 LBT	Judkins Right 4 - Long Brite Tip		6720820L
JR 5	Judkins Right 5	55808400	67208400

JR 3.5 JR 4 JR 5

SH = Side Holes. ST = Short Tip.

Judkins Curved Right

Shape	Shape Description	5F	6F
JCR 4	Judkins Curved Right 4		67209800

JCR 4

Extra Back-up

Shape	Shape Description	5F	6F
XB 2	Extra Back-up 2		67246400
XB 2.5	Extra Back-up 2.5		67246500
XB 2.75	Extra Back-up 2.75		67273000
XB 3	Extra Back-up 3	55805200	67205200
XB 3 SH	Extra Back-up 3 - 2 side holes		67205300
XB 3.25	Extra Back-up 3.25	55840000	67240000
XB 3.5	Extra Back-up 3.5	55805400	67205400
XB 3.5 (90 cm)	Extra Back-up 3.5 (90 cm)		67205490
XB 3.5 (125 cm)	Extra Back-up 3.5		67205425
XB 3.5 SH	Extra Back-up 3.5 - 2 side holes		67205500
XB 3.5 LBT	Extra Back-up 3.5 - Long Brite Tip		6720540L
XB 4	Extra Back-up 4	55805600	67205600
XB 4 SH	Extra Back-up 4 - 2 side holes		67205700
XB 4.5	Extra Back-up 4.5	55805800	67205800
XB 4.5 SH	Extra Back-up 4.5 - 2 side holes		67205900

XB 2

XB 2.5

XB 2.75

XB 3

XB 3.25

XB 3.5

XB 4

XB 4.5

SH = Side Holes.

Extra Back-up Contralateral

Shape	Shape Description	5F	6F
XBC 2.5	Extra Back-up Contralateral 2.5		67273200
XBC 2.75	Extra Back-up Contralateral 2.75		67273400
XBC 3	Extra Back-up Contralateral 3		67207000
XBC 3 SH	Extra Back-up Contralateral 3 - 2 side holes		67207100
XBC 3.25	Extra Back-up Contralateral 3.25		67273600
XBC 3.5	Extra Back-up Contralateral 3.5	55807200	67207200
XBC 3.5 SH	Extra Back-up Contralateral 3.5 - 2 side holes		67207300
XBC 4	Extra Back-up Contralateral 4	55807400	67207400
XBC 4 SH	Extra Back-up 4 Contralateral - 2 side holes		67207500
XBC 4.5	Extra Back-up 4.5 Contralateral		67207600
XBC 4.5 SH	Extra Back-up 4.5 Contralateral - 2 side holes		67207700

XBC 2.5 XBC 2.75 XBC 3 XBC 3.25 XBC 3.5 XBC 4 XBC 4.5

Extra Back-up Lad

Shape	Shape Description	5F	6F
XBLAD 3	Extra Back-up LAD 3		67206600
XBLAD 3 SH	Extra Back-up LAD 3 -2 side holes		67206700
XBLAD 3.5	Extra Back-up LAD 3.5	55806000	67206000
XBLAD 3.5 SH	Extra Back-up LAD 3.5 - 2 side holes		67206100
XBLAD 4	Extra Back-up LAD 4	55806200	67206200
XBLAD 4 SH	Extra Back-up LAD 4 - 2 side holes		67206300
XBLAD 4.5	Extra Back-up LAD 4.5		67206400

XBLAD 3 XBLAD 3.5 XBLAD 4 XBLAD 4.5

SH = Side Holes. ST = Short Tip.

Extra Back-up RCA

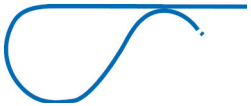
Shape	Shape Description	5F	6F
XBRCA	Extra Back-up Right CA	55812600	67212600
XBRCA SH	Extra Back-up Right CA - 2 side holes		67212700

XBRCA

Extra Back-up Right

Shape	Shape Description	5F	6F
XBR 1	Extra Back-up Right 1		67212200
XBR 1 SH	Extra Back-up Right 1 - 2 side holes		67212300
XBR 2	Extra Back-up Right 2		67212400

XBR 1



XBR 2

Amplatz Left

Shape	Shape Description	5F	6F
AL .75	Amplatz Left 0.75	55803400	67203400
AL .75 SH	Amplatz Left 0.75 - 2 side holes		67203500
AL 1	Amplatz Left 1	55803600	67203600
AL 1 ST	Amplatz Left 1 - Short tip		67204400
AL 1 SH	Amplatz Left 1 - 2 side holes		67203700
AL 1.5	Amplatz Left 1.5	55803800	67203800
AL 2	Amplatz Left 2	55804000	67204000
AL 2 SH	Amplatz Left 2 - 2 side holes		67204100
AL 2 LBT	Amplatz Left 2 - Long Brite Tip		6720400L
AL 3	Amplatz Left 3	55804200	67204200

AL .75

AL 1

AL 1.5

AL 2

AL 3

SH = Side Holes. ST = Short Tip.

Amplatz Right

Shape	Shape Description	5F	6F
AR 1	Amplatz Right 1	55811000	67211000
AR 1 SH	Amplatz Right 1 - 2 side holes		67211100
AR 1 LBT	Amplatz Right 1 - Long Brite Tip		6721100L
AR 2	Amplatz Right 2	55811200	67211200
AR 2 SH	Amplatz Right 2 - 2 side holes		67211300
AR 2 LBT	Amplatz Right 2 - Long Brite Tip		6721120L


AR 1 AR 2

3-Dimensional Right Coronary - Williams Technique

Shape	Shape Description	5F	6F
3DRC	Right Coronary 3 Dimensional	55813000	67213000
3DRC SH	Right Coronary 3 Dimensional - 2 side holes		67213100


3 DRC

Noto Technique

Shape	Shape Description	5F	6F
NR 4	Noto Technique Right 4		67212000


NR 4

Hockey Stick

Shape	Shape Description	5F	6F
H -STICK	Hockey Stick	55827800	67227800
H -STICK SH	Hockey Stick - 2 side holes		67227900
H -STICK (90 cm)	Hockey Stick - Short Shaft		67227890
H -STICK LBT	Hockey Stick - Long Brite Tip		6722780L


H-STICK

SH = Side Holes. ST = Short Tip.

Multipurpose A 1

Shape	Shape Description	5F	6F
MPA 1	Multipurpose A 1	55827000	67227000
MPA 1 SH	Multipurpose A 1 - 2 side holes		67227100
MPA 1 (125cm)	Multipurpose A 1		67227025
MPA 1 LBT	Multipurpose A 1 - Long Brite Tip		6722700L

MPA 1

Multipurpose B 1

Shape	Shape Description	5F	6F
MPB 1	Multipurpose B 1		67227200

MPB 1

Shapes For Coronary Bypass: Left

Shape	Shape Description	5F	6F
LCB	Left Coronary Bypass	55818000	67218000
LCB (90cm)	Left Coronary Bypass - Short Shaft		67218090
LCB SH	Left Coronary Bypass - 2 side holes		67218100

LCB

Shapes For Coronary Bypass: Right

Shape	Shape Description	5F	6F
RCB	Right Coronary Bypass		67218200
RCB (90cm)	Right Coronary Bypass - Short shaft		67218290
RCB SH	Right Coronary Bypass - 2 side holes		67218300

RCB

SH = Side Holes.

Dedicated Radial Shapes

Shape	Shape Description	5F	6F
JFL	Left Transradial	55816200	67216200
JFL -ST	Left Transradial - short tip		67216400
JFR	Right Transradial		67216800
JFR -ST	Right Transradial - short tip		67217000
RB	Radiobrachial		67217200
BARBEAU	Barbeau	55817400	67217400
RBL 4	Radial Bilateral 4.0		67271700
RBL 4.5	Radial Bilateral 4.5		67272900
RBL 5.0	Radial Bilateral 5.0		67271800
RRAD5	Right Radial 5	55832000	
RRAD6	Right Radial 6		67230000
RRAD6 XB	Right Radial 6 Extra Back-up		67230100

JFL

JFR

RB

BARBEAU

RBL 4

RBL 4.5

RBL 5.0

RRAD5

RRAD6

RRAD6 XB

ST = Short Tip.

Internal Mammary

Shape	Shape Description	5F	6F
IM (90 cm)	Internal Mammary, 90 cm		67219090
IM	Internal Mammary	55819000	67219000
IM SH	Internal Mammary, 2 side holes		67219100

IM

Renal Curve

Shape	Shape Description	5F	6F
RDC (55cm)	Renal Double Curve		67221255

RDC

SH = Side Holes.

ADROIT® Guiding Catheter - EcoPac Five Pacs

Key Features

- Units per package: 5

Judkins Left

Shape	Shape Description	6F
JL 4	Judkins Left 4 ECOPAC	6720040E

Extra Back-up

Shape	Shape Description	6F
XB 3.5	Extra Back-up 3.5 ECOPAC	6720540E

Judkins Right

Shape	Shape Description	6F
JR 4	Judkins Right 4 ECOPAC	6720820E

VISTA BRITE TIP® and Long VISTA BRITE TIP® Guiding Catheters

A complete system of large lumen guiding catheters that are optimized to answer clinical needs. The performance-based design of each catheter easily meets strength, control and delivery requirements in the widest range of anatomies.

Key Features

- The multisegment technology and hybrid braiding provides large inner diameters and excellent deliverability
- TRUELUMEN® Technology facilitates consistent ID from hub to tip, for added confidence
- PTFE lining provides lubricious lumen for smoother delivery
- Complete choice of shapes
- 5F to 9F platform
- Transradial shapes
- Extra back-up shapes
- Long Brite Tip shapes
- All catheters 100 cm except when noted
- Units per package: 1

For information on indications, contraindications, warnings, and precautions, see page 89.

Judkins Left

Shape	Shape Description	5F	6F	7F	8F
JL 3	Judkins Left 3		67000000	77800000	588812
JL 3 ST	Judkins Left 3 - short tip		SM7338		
JL 3 SH	Judkins Left 3 - 2 side holes			77800100	
JL 3.5	Judkins Left 3.5	55600200	67000200	77800200	588851
JL 3.5 LBT	Judkins Left 3.5 - Long BRITE TIP®	5560020L			
JL 3.5 ST	Judkins Left 3.5 - short tip		67001200	77801200	
JL 3.5 SH	Judkins Left 3.5 - 2 side holes		67000300	77800300	588832
JL 4 (90 cm)	Judkins Left 4		67000490	77800490	588823T
JL 4	Judkins Left 4	55600400	67000400	77800400	588823
JL 4 LBT	Judkins Left 4 - Long BRITE TIP®	5560040L	6700040L		
JL 4 ST	Judkins Left 4 - short tip		67001400	77801400	
JL 4 SH	Judkins Left 4 - 2 side holes		67000500	77800500	588834
JL 4 ST SH	Judkins Left 4 - short tip - 2 side holes			77801500	
JL 4 (125 cm)	Judkins Left 4 - 125 cm length		SM7435		
JL 4.5	Judkins Left 4.5		67000600	77800600	588852
JL 4.5 LBT	Judkins Left 4.5 - Long BRITE TIP®	SM7329			
JL 4.5 ST	Judkins Left 4.5 - short tip		67001600		
JL 4.5 SH	Judkins Left 4.5 - 2 side holes			77800700	588810
JL 5	Judkins Left 5	55600800	67000800	77800800	588853
JL 5 LBT	Judkins Left 5 - Long BRITE TIP®	5560080L			
JL 5 SH	Judkins Left 5 - 2 side holes			77800900	588800
JL 5 (125 cm)	Judkins Left - 125 cm length		SM7361		
JL 6	Judkins Left 6		67001000	77801000	588854

JL 3 JL 3.5 JL 4 JL 4.5 JL 5 JL 6

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Judkins Curved Left

Shape	Shape Description	5F	6F	7F	8F
JCL 3.5	Judkins Curved Left 3.5		67002600		
JCL 3.5 SH	Judkins Curved Left 3.5 - Side Holes		67002700		
JCL 4	Judkins Curved Left 4		67002800		5888108

JCL 3.5

Judkins Right

Shape	Shape Description	5F	6F	7F	8F
JR 3.5	Judkins Right 3.5	55608000	67008000	77808000	588855
JR 3.5 (125 cm)	Judkins Right 3.5, 125 cm		SM7371		
JR 3.5 LBT	Judkins Right 3.5 - Long Brite Tip	5560800L			
JR 3.5 SH	Judkins Right 3.5, 2 side holes		67008100	77808100	588828
JR 4 (90 cm)	Judkins Right 4, 90 cm		67008290	77808290	588830T
JR 4 (98 cm)	Judkins Right 4 - 98cm				
JR 4	Judkins Right 4	55608200	67008200	77808200	588830
JR 4 LBT	Judkins Right 4 Long BRITE TIP®	5560280L	6700820L		
JR 4 LBT (110 cm)	Judkins Right 4 Long BRITE TIP®, 110 cm	SM7426			
JR 4 ST	Judkins Right 4 short tip		67009000	77809000	
JR 4 SH	Judkins Right 4, 2 side holes		67008300	77808300	588831
JR 4 SH (90 cm)	Judkins Right 4 - 2 side holes - 90cm			77808390	588831T
JR 4 (125 cm)	Judkins Right 4, 125 cm	SM7739	SM7436		
JR 4.5 ST	Judkins Right 4.5 - Short Tip			77809200	
JR 5	Judkins Right 5		67008400	77808400	588-856
JR 5 LBT	Judkins Right 5 Long BRITE TIP®	SM7330			
JR 5 SH	Judkins Right 5, 2 side holes			77808500	

JR 3.5 JR 4 JR 5

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Judkins Curved Right

Shape	Shape Description	5F	6F	7F	8F
JCR 4	Judkins Curved Right 4		67009800		

JCR 4

Extra Back-up Left

Shape	Shape Description	5F	6F	7F	8F
XB 2.5	Extra Back-up 2.5		SM7465		
XB 3	Extra Back-up 3	SM7328	67005200	77805200	588829
XB 3 (90 cm)	Extra Back-up 3, 90 cm				SM7720
XB 3 (125 cm)	Extra Back-up 3, 125 cm			SM7395	
XB 3 LBT	Extra Back-up 3 Long BRITE TIP®				
XB 3 SH	Extra Back-up 3, 2 side holes		67005300	77805300	588875
XB 3.5 (90 cm)	Extra Back-up 3.5, 90 cm		67005490	SM7710	SM7721
XB 3.5	Extra Back-up 3.5	55605400	67005400	77805400	588882
XB 3.5 LBT	Extra Back-up 3.5 Long BRITE TIP®	5560540L	6700540L		
XB 3.5 SH	Extra Back-up 3.5, 2 side holes		67005500	77805500	588885
XB 3.5 (98 cm)	Extra Back-up 3.5, 98 cm				
XB 4	Extra Back-up 4	55905600	67005600	77805600	588894
XB 4 (90 cm)	Extra Back-up 4, 90 cm				SM7712
XB 4 LBT	Extra Back-up 4 Long BRITE TIP®	5560560L	SM7148		
XB 4 SH	Extra Back-up 4, 2 side holes		67005700	77805700	588896
XB 4 SH (85cm)					SM7621
XB 4.5	Extra Back-up 4.5		67005800	77805800	588898
XB 4.5 SH	Extra Back-up 4.5, 2 side holes		67005900	77805900	

XB 2.5 XB 3 XB 3.5 XB 4 XB 4.5

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Extra Back-up Contralateral

Shape	Shape Description	5F	6F
XBC 3	Extra Back-up Circumflexa 3		67007000
XBC 3 SH	Extra Back-Up 3.5 - 2 side holes		67007100
XBC 3.5	Extra Back-up Circumflexa 3.5		67007200
XBC 3.5 LBT	Extra Back-Up 3.5 - Long Brite Tip	SM7466	
XBC 3.5 SH	Extra Back-up Circumflexa 3.5, 2 side holes		67007300
XBC 4 SH	Extra Back-Up 4		67007400
XBC 4 SH	Extra Back-up Circumflexa 4, 2 side holes		67007500
XBC 4.5 SH	Extra Back-Up 4.5 - 2 side holes		67007700

XBC 3 XBC 3.5 XBC 4 XBC 4.5

Extra Back-up LAD

Shape	Shape Description	5F	6F	7F	8F
XB LAD 3	Extra Back-Up LAD 3		67006600		
XB LAD 3 SH	Extra Back-Up LAD 3 - 2 side holes		67006700		
XB LAD 3.5	Extra Back-Up LAD 3.5	55606000	67006000	77806000	5888100
XB LAD 3.5 LBT	Extra Back-Up LAD 3.5 - Long Brite Tip	5560600L			
XB LAD 3.5 SH	Extra Back-Up LAD 3.5 - 2 side holes		67006100	77806100	5888103
XB LAD 4	Extra Back-Up LAD 4		67006200	77806200	5888101
XB LAD 4 SH	Extra Back-Up LAD 4 - 2 side holes		67006300	77806300	5888104
XB LAD 4.5	Extra Back-Up LAD 4.5		67006400	77806400	

XB LAD 3 XB LAD 3.5 XB LAD 4 XB LAD 4.5

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Extra Back-up RCA

Shape	Shape Description	5F	6F	7F	8F
XBR CA 90cm	Extra Back-Up Right CA 90cm				SM7746
XBR CA	Extra Back-Up Right CA	SM7744	67012600	G778126	
XBR CA SH	Extra Back-up Right CA - 2 side holes		67012700		

XBR CA

Extra Back-up Right

Shape	Shape Description	5F	6F	7F	8F
XBR 1	Extra Back-up Right 1		67012200		
XBR 1 SH	Extra Back-Up Right 1 - 2 side holes		67012300		
XBR 2	Extra Back-up Right 2		67012400		

XBR 1 XBR 2

SH = Side Holes.

Amplatz Left

Shape	Shape Description	5F	6F	7F	8F
AL .75	Amplatz Left 0.75	SM7722	67003400	77803400	588890
AL .75 (90 cm)	Amplatz Left 0.75, 90 cm		SM7719		
AL .75 SH	Amplatz Left 0.75 - 2 side holes		67003500	G782AL75	SM7649
AL .75 ST	Amplatz Left 0.75 - short tip		SM7664		
AL 1	Amplatz Left 1		67003600	77803600	588843
AL 1 (90 cm)	Amplatz Left 1 - 90 cm catheter length			77803690	588843T
AL 1 LBT	Amplatz Left 1 - Long BRITE TIP®	SM7331			
AL 1 ST	Amplatz Left 1 - short tip	SM7503	67004400	77804400	588870
AL 1 SH	Amplatz Left 1 - 2 side holes		67003700	77803700	588847
AL 1 SH (85cm)	Amplatz Left 1 - 2 side holes - 85cm				SM7622
AL 1 ST SH	Amplatz Left 1 - short tip - 2 side holes		SM7443	77804500	588873
AL 1 (90 Degree Angle)	Amplatz Left 1 - 90 Degree Angle - 100cm			SM7603	
AL 1 (110 cm)	Amplatz Left 1 - 110cm catheter length		SM7705		
AL 1.5	Amplatz Left 1.5		67003800	77803800	
AL 1.5 SH	Amplatz Left 1.5 - 2 side holes			77803900	588878
AL 2 (80 cm)	Amplatz Left 2 - 80 cm catheter length				SM7342
AL 2 (90 cm)	Amplatz Left 2, 90 cm catheter length			77804090	588844T
AL 2	Amplatz Left 2	55604000	67004000	77804000	588844
AL 2 LBT	Amplatz Left 2 - Long BRITE TIP®	5560400L	6700400L		
AL 2 ST	Amplatz Left 2 - short tip			77804800	
AL 2 SH	Amplatz Left 2 - 2 side holes		67004100	77804100	588848
AL 3	Amplatz Left 3		67004200	77804200	588849
AL 3 SH	Amplatz Left 3 - 2 side holes			77804300	588891

AL .75 AL 1 AL 1.5 AL 2 AL 3

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Amplatz Right

Shape	Shape Description	5F	6F	7F	8F
AR 1	Amplatz Right 1		67011000	77811000	588845
AR 1 LBT	Amplatz Right 1 Long BRITE TIP®	SM7333	6701100L		
AR 1 SH	Amplatz Right 1 - 2 side holes			77811100	588836
AR 2	Amplatz Right 2	55611200	67011200	77811200	588846
AR 2 LBT	Amplatz Right 2 Long BRITE TIP®	5561120L	6701120L		
AR 2 SH	Amplatz Right 2 - 2 side holes		67011300	77811300	588837
AR 3	Amplatz Right 3	SM7750			
AR-MOD	Amplatz Right Mod, 100 cm		SM7398		

AR 1 AR 2 AR MOD

3-Dimensional Right Coronary - Williams Technique

Shape	Shape Description	5F	6F	7F	8F
3DRC	Right Coronary 3 Dimensional		67013000		
3DRC LBT	Right Coronary 3 Dimensional - Long Brite Tip	SM7599			
3DRC SH	Right Coronary 3 Dimensional - 2 side holes		67013100	G7823DRC	

3 DRC

Noto Technique

Shape	Shape Description	5F	6F	7F	8F
NR 4	Noto Technique Right 4		67012000		

NR 4

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Hockey Stick

Shape	Shape Description	5F	6F	7F	8F
H-STICK (90cm)	Hockey Stick - 90cm catheter length		67027890	77827890	588841T
H -STICK	Hockey Stick		67027800	77827800	588841
H-STICK LBT	Hockey Stick - Long Brite Tip	5562780L	6702780L		
H -STICK SH	Hockey Stick, 2 side holes		67027900	77827900	588850
H -STICK SH (90cm)	Hockey Stick, 2 side holes - 90cm catheter length			77827990	588850T

H-STICK

Multipurpose A 1

Shape	Shape Description	5F	6F	7F	8F	9F
MPA (120)	Multipurpose A - 120 cm catheter length		SM7470			
MPA 1 LBT	Multipurpose A 1 - Long BRITE TIP®	5562700L	6702700L			
MPA 1 LBT (110cm)	Multipurpose A 1 - Long BRITE TIP® - 110 cm					
MPA 1	Multipurpose A 1		67027000	77827000	588842	
MPA 1 (115cm)	Multipurpose A 1 - 115 cm catheter length		SM7462			
MPA 1 SH	Multipurpose A 1 - 2 side holes		67027100	77827100	588892	
MPA 1 (125cm)	Multipurpose A 1 - 125 cm catheter length		SM7477	SM7206		
MPA 1 (98cm)	Multipurpose A-1 - 98 cm catheter length					598942
MPA 1 (55cm)	Multipurpose A-1 - 55 cm catheter length			77827055	588840P	
MPA 2 (125cm)	Multipurpose A 2 - 125 cm catheter length		SM7394			

MPA 1 MPA 2

Multipurpose B 1

Shape	Shape Description	5F	6F	7F	8F
MPB 1	Multipurpose B 1		67027200		
MPB 1 SH	Multipurpose B 1 - 2 side holes			77827300	

MPB 1

SH = Side Holes. ST = Short Tip. LBT = Long BRITE TIP.

Dedicated Radial Shapes

Shape	Shape Description	5F	6F
JFL	Left Transradial	55616200	67016200
JFL -ST	Left Transradial - short tip		67016400
JFL -LBT	Left Transradial - Long BRITE TIP®	5561620L	67016600
JFR	Right Transradial	55616800	67016800
JFR -ST	Right Transradial - short tip		67017000
JFR - LT	Right Transradial - Long BRITE TIP®	5561680L	
RB	Radiobrachial	55617200	67017200
RB LBT	Radiobrachial - Long BRITE TIP®	5561720L	
BARBEAU	Barbeau	55617400	67017400
BARBEAU LBT	Barbeau - Long BRITE TIP®	556174-0L	SM7542
RBL 4.0	Radial Bilateral 4.0		SM7717
RBL 4.5	Radial Bilateral 4.5		SM7729
RBL 5.0	Radial Bilateral 5.0		SM7718
BBL	Brachial Bilateral		SM7636

JFL

JFR

RB

BARBEAU

RBL 4.0

RBL 4.5

RBL 5.0

RRAD 5

RRAD 6

ST = Short Tip. LBT = Long BRITE TIP.

Shapes For Coronary Bypass: Left

Shape	Shape Description	5F	6F	7F	8F
LCB	Left Coronary Bypass		67018000	77818000	588815
LCB (90cm)	Left Coronary Bypass - 90 cm catheter length		67018090	77818090	588815T
LCB LBT	Left Coronary Bypass - Long Brite Tip	SM7327			
LCB SH	Left Coronary Bypass - 2 side holes		67018100	77818100	


LCB

Shapes For Coronary Bypass: Right

Shape	Shape Description	5F	6F	7F	8F
RCB	Right Coronary Bypass		67018200	77818200	588816
RCB (90 cm)	Right Coronary Bypass - 90 cm catheter length		67018290	G780RCBH	588816T
RCB SH	Right Coronary Bypass - 2 side holes		67018300	77818300	


RCB

Coronary Bypass: Left Internal Mammary

Shape	Shape Description	5F	6F	7F	8F
LIMA (90 cm)	3D LIMA Left Internal Mammary Artery, 90 cm		SM7501		
LIMA (125cm)	3D LIMA Left Internal Mammary Artery, 125 cm	SM7740	SM7741		


LIMA

Internal Mammary

Shape	Shape Description	5F	6F	7F	8F
IM (90 cm)	Internal Mammary - 90 cm shaft length		67019090	77819090	588801
IM	Internal Mammary		67019000	77819000	588817
IM LBT	Internal Mammary - Long BRITE TIP	5561900L			
IM SH	Internal Mammary - 2 side holes		67019100	77819100	588820
IM SH (90cm)	Internal Mammary - 2 side holes - 90cm catheter			77819190	
IM VB-1	Internal Mammary - 100cm		67019200		


IM

SH = Side Holes. LBT = Long BRITE TIP.

Champ - Bypass Backup Support

Shape	Shape Description	5F	6F	7F	8F
CHAMP	Champ 1.0		SM7467		

Sones

Shape	Shape Description	5F	6F	7F	8F
SON 1	Sones 1				588880

Castillo

Shape	Shape Description	5F	6F	7F	8F
CAS 1	Castillo 1		G667683		588883
CAS 2	Castillo 2				588884

VISTA BRITE TIP® and Long VISTA BRITE TIP® Guiding Catheters - EcoPac Five Pacs

Key Features

- Units per package: 5

Judkins Left

Shape	Shape Description	6F
JL 4	Judkins Left 4	6700040E
JL 4 SH	Judkins Left 4 - 2 side holes	SM7427

Extra Back-up

Shape	Shape Description	6F
XB 3.5	Extra Back-up 3.5	6700540E

Judkins Right

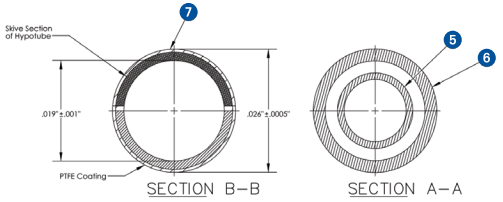
Shape	Shape Description	6F
JR 4	Judkins Right 4	6700820E

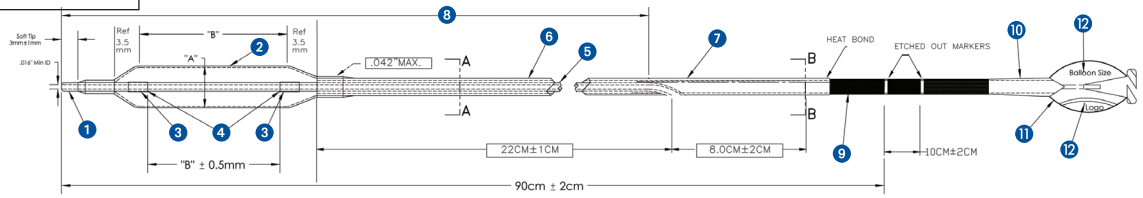
EMPIRA® RX Pre-Dilatation Catheter

Key Features

- Help you reach and treat the lesion with excellent crossability and recrossability.
- Low-profile balloon for crossing even the most difficult lesions.
- Exceptional crossability and recrossability is achieved through the highly flexible DURALYN® Flex balloon material, the proprietary pleating and folding process, and also the lubricious as well as durable hydrophilic coating.
- Versatile by bringing crossability and controlled growth to a wide range of lesions.
- Guidewire diameter: .014"
- Nominal pressure: 8 atm
- Rated burst pressure: 14 atm, 1/4 size at 16 atm
- Usable length: 139 cm
- Units per package: 1

1. Formed Soft Tip – Yellow
2. Balloon
3. Radiopaque Marker Bands
4. Adhesive Loctite 4041
5. Inner Member Tubing - Black
6. Outer Member Tubing - Natural .029"/.035" + /- .001"
7. Intermediate Shaft - Natural .029"/.035" + /- .001"
8. Hydrophilic Coating
9. Skived Proximal Shaft
10. Strain Relief - White
11. Molded Luer Hub
12. Ink - White





Balloon Diameter	Balloon Length					
	6	10	12	15	20	30
1.50	85R06150S	85R10150S	85R12150S	85R15150S	85R20150S	
2.00	85R06200S	85R10200S	85R12200S	85R15200S	85R20200S	85R30200S
2.25		85R10225S	85R12225S	85R15225S	85R20225S	
2.50	85R06250S	85R10250S	85R12250S	85R15250S	85R20250S	85R30250S
2.75		85R10275S	85R12275S	85R15275S	85R20275S	
3.00		85R10300S	85R12300S	85R15300S	85R20300S	85R30300S
3.25				85R15325S		
3.50		85R10350S	85R12350S	85R15350S	85R20350S	85R30350S
4.00		85R10400S	85R12400S	85R15400S	85R20400S	

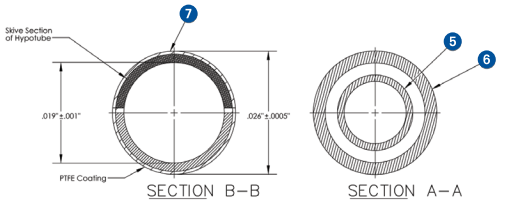
For information on indications, contraindications, warnings, and precautions, see page 76.

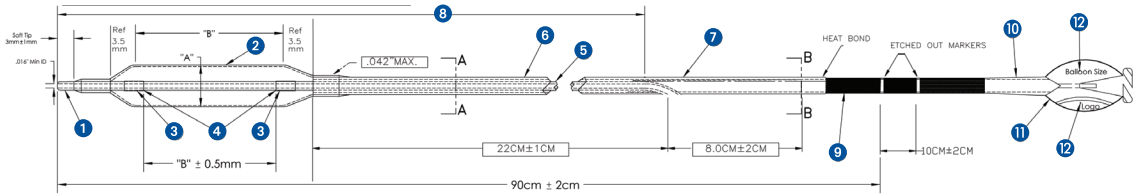
EMPIRA NC® RX Pre-Dilatation Catheter

Key Features

- Designed to deliver in challenging interventions and post-dilatation procedures.
- High-pressure balloon that combines exceptional crossability and recrossability with accuracy during postdilatation.
- Exceptional crossability and recrossability is achieved through the highly flexible DURALYN® Flex balloon material, the proprietary pleating and folding process, and also the lubricious as well as durable hydrophilic coating.
- Guidewire diameter: .014"
- Nominal pressure: 14 atm
- Rated burst pressure: 20 atm, 1/4 size at 26 atm
- Usable length: 139 cm
- Units per package: 1

1. Formed Soft Tip – Yellow
2. Balloon
3. Radiopaque Marker Bands
4. Adhesive Loctite 4041
5. Inner Member Tubing - Black
6. Outer Member Tubing - Natural .029"/.035" + /- .001"
7. Intermediate Shaft - Natural .029"/.035" + /- .001"
8. Hydrophilic Coating
9. Skived Proximal Shaft
10. Strain Relief - White
11. Molded Luer Hub
12. Ink - White



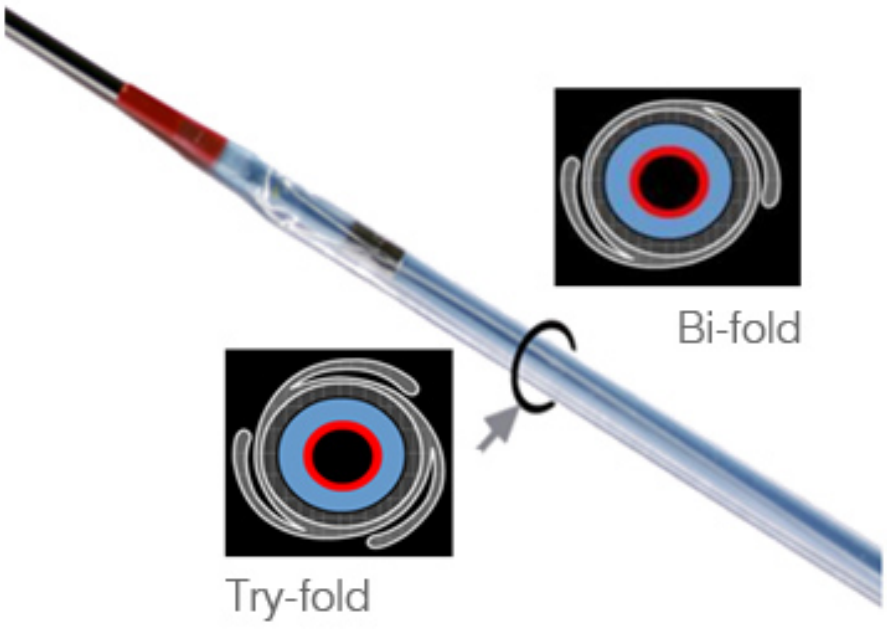


Balloon Diameter	Balloon Length						
	6	10	12	15	20	25	30
2.00	75R06200N	75R10200N	75R12200N	75R15200N	75R20200N		
2.25		75R10225N	75R12225N	75R15225N	75R20225N		
2.50	75R06250N	75R10250N	75R12250N	75R15250N	75R20250N	75R25250N	75R30250N
2.75		75R10275N	75R12275N	75R15275N	75R20275N		
3.00	75R06300N	75R10300N	75R12300N	75R15300N	75R20300N	75R25300N	75R30300N
3.25		75R10325N	75R12325N	75R15325N	75R20325N		
3.50	75R06350N	75R10350N	75R12350N	75R15350N	75R20350N	75R25350N	75R30350N
3.75		75R10375N	75R12375N	75R15375N	75R20375N		
4.00		75R10400N	75R12400N	75R15400N	75R20400N		

For information on indications, contraindications, warnings, and precautions, see page 76.

MOZEC™ PTCA Balloon Dilatation Catheter

With the longest PTCA balloons on the U.S. market, the MOZEC™ PTCA Balloon Dilatation Catheter is available in a broad range of sizes for greater cath lab efficiency. The durable, flexible FeatherLite™ catheter construction transfers more push from hub to tip*. The proprietary memory shape balloon refolds to original size following deflation for excellent rewrap** and the elongated tip and seamless transition facilitates access through tight lesions.



Key features

- MeriGlide™ hydrophilic coating from distal balloon neck up to Rx port for smooth navigation
- Excellent pushability to deliver through complexity
- Tight rewrap after repeat dilatations
- Catheter System: Rapid Exchange (RX)
- Guide Catheter Compatibility: 5 F (min I. D. 0.056" / 1.42 mm)
- Guidewire diameter: .014"
- Nominal pressure: 7 atm for all diameters
- Rated burst pressure: 16 atm for 1.50 to 4.00 mm, 14 atm for 4.50 mm
- Catheter shaft length: 142 cm
- Wrap: 2 folds for 1.50 and 2.00 mm, 3 folds for 2.25 to 4.50 mm
- Units per package: 1

Balloon Diameter	Balloon Length									
	9	12	14	15	17	20	25	30	38	41
1.5	MOZ15009	MOZ15012		MOZ15015						
2	MOZ20009	MOZ20012		MOZ20015						
2.25	MOZ22509		MOZ22514		MOZ22517	MOZ22520	MOZ22525	MOZ22530	MOZ22538	
2.5	MOZ25009		MOZ25014		MOZ25017	MOZ25020	MOZ25025	MOZ25030	MOZ25038	MOZ25041
2.75	MOZ27509		MOZ27514		MOZ27517	MOZ27520	MOZ27525	MOZ27530	MOZ27538	
3	MOZ30009		MOZ30014		MOZ30017	MOZ30020	MOZ30025	MOZ30030	MOZ30038	MOZ30041
3.5	MOZ35009		MOZ35014		MOZ35017	MOZ35020	MOZ35025	MOZ35030	MOZ35038	
4	MOZ40009		MOZ40014		MOZ40017	MOZ40020	MOZ40025	MOZ40030		
4.5	MOZ45009		MOZ45014		MOZ45017	MOZ45020	MOZ45025	MOZ45030		

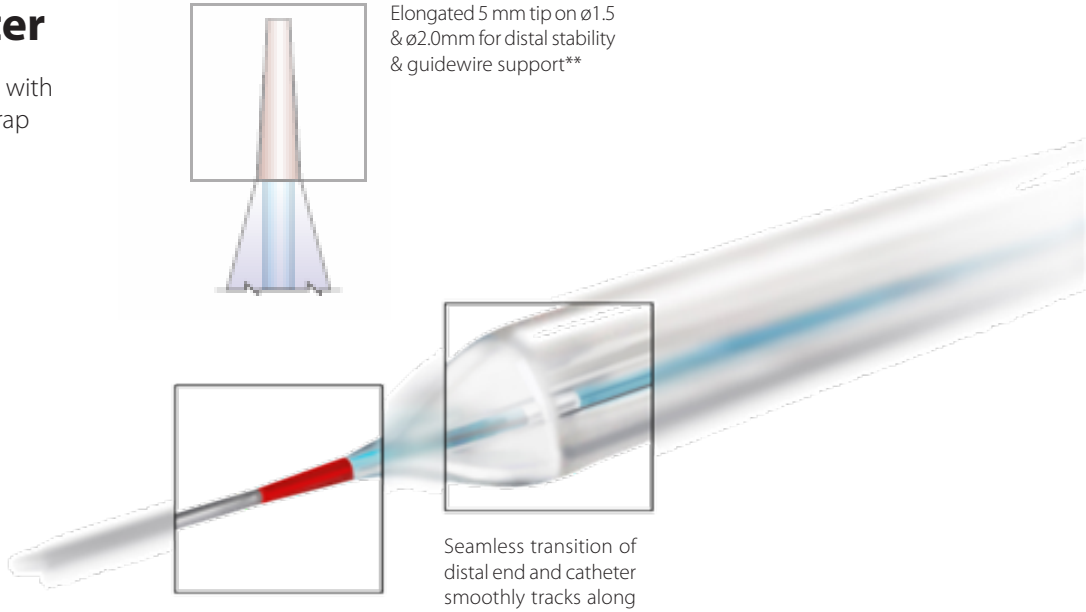
*FeatherLite™ catherter construction applicable for MOZEC™ PTCA balloon dilatation catheter only.
**Nylon balloon material applicable for MOZEC™ PTCA balloon dilatation catheter only.

MOZEC™ NC PTCA Balloon Dilatation Catheter

The MOZEC™ NC PTCA Balloon Dilatation Catheter combines controlled balloon growth with the longest lengths on on the U.S. market. It offers excellent cross and recross, tight rewrap and exceptionally low tracking forces.*

Key features

- MeriGlide™ hydrophilic coating from distal balloon neck up to Rx port for smooth navigation
- Excellent pushability to deliver through complexity
- Tight rewrap after repeat dilatations
- Catheter System: Rapid Exchange (RX)
- Balloon Material: Nylon Non-Compliant
- Guide Catheter Compatibility: 5 F (min I. D. 0.056" / 1.42 mm)
- Guidewire diameter: .014"
- Nominal pressure: 12 atm for all diameters
- Rated burst pressure: 20 atm
- Catheter shaft length: 142 cm
- Wrap: 2 folds for 2.00 mm, 3 folds for 2.25 to 4.50 mm
- Units per package: 1



Balloon Diameter	Balloon Length							
	8	10	13	15	18	23	28	35
2.00	MNC20008	MNC20010	MNC20013	MNC20015	MNC20018	MNC20023	MNC20028	MNC20035
2.25	MNC22508	MNC22510	MNC22513	MNC22515	MNC22518	MNC22523	MNC22528	MNC22535
2.50	MNC25008	MNC25010	MNC25013	MNC25015	MNC25018	MNC25023	MNC25028	MNC25035
2.75	MNC27508	MNC27510	MNC27513	MNC27515	MNC27518	MNC27523	MNC27528	MNC27535
3.00	MNC30008	MNC30010	MNC30013	MNC30015	MNC30018	MNC30023	MNC30028	MNC30035
3.50	MNC35008	MNC35010	MNC35013	MNC35015	MNC35018	MNC35023	MNC35028	MNC35035
4.00	MNC40008	MNC40010	MNC40013	MNC40015	MNC40018	MNC40023	MNC40028	MNC40035
4.50	MNC45008	MNC45010	MNC45013	MNC45015	MNC45018	MNC45023	MNC45028	MNC45035

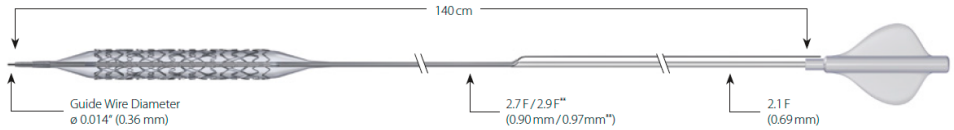
*Data on file at Meril Life Sciences Pvt. Ltd.
** Elongated Tip applicable for Mozec™ PTCA balloon dilatation catheter only.

NIRxcell™ CoCr Coronary Stent System

NIRxcell™ Stent System is indicated for improving coronary luminal diameter in patients with symptomatic ischemic heart disease associated with stenotic lesions in de novo native coronary arteries (length ≤ 30 mm) with a reference vessel diameter of 2.50 mm to 4.00 mm.

Key features

- Superior Deliverability with Novel Spring Tip
- Narrowest Strut Width from Dual-Pattern Strut Design
- Exceptional Clinical Outcomes with TLR rate of 5.1% at 9 months
- Thermo-Treated Reinforced Hypotube for improved hub-to-tip force transfer
- Best-in-class Conformability with the WiZeCell™ Stent Design
- Catheter System: Rapid Exchange (RX)
- Guide Catheter Compatibility (ID): ≥ 5 F (0.056")
- Guidewire diameter: .014"
- Radiopaque Markers: 2 - Platinum / Iridium
- Proximal Shaft Diameter: 2.1 F (0.69 mm)
- Distal Shaft Diameter: 2.7 F (0.9 mm) for stent length 8 - 28 mm, 2.9 (0.97 mm) for stent length 33 mm
- Nominal Pressure (NP): 12 atm
- Rated Burst Pressure (RBP): 18 atm



Balloon Diameter	Balloon Length						
	8	12	17	20	24	28	33
2.5	NXL25008US	NXL25012US	NXL25017US	NXL25020US	–	–	–
2.75	NXL27508US	NXL27512US	NXL27517US	NXL27520US	NXL27524US	NXL27528US	NXL27533US
3	NXL30008US	NXL30012US	NXL30017US	NXL30020US	NXL30024US	NXL30028US	NXL30033US
3.5	NXL35008US	NXL35012US	NXL35017US	NXL35020US	NXL35024US	NXL35028US	NXL35033US
4	NXL40008US	NXL40012US	NXL40017US	NXL40020US	NXL40024US	NXL40028US	NXL40033US

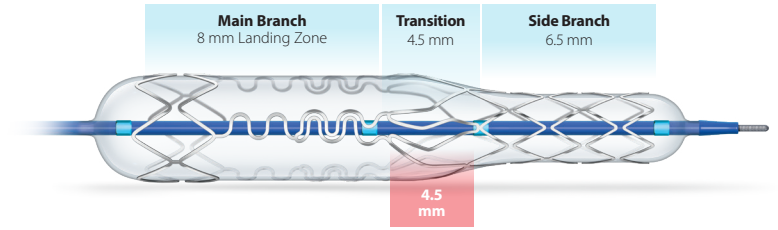
TRYTON Side Branch Stent

The TRYTON Side Branch Stent is indicated for improving the side branch luminal diameter of de novo native coronary artery bifurcation lesions (Medina Classification 1.1.1; 0.1.1; 1.0.1) with a side branch diameter stenosis of ≥50% and a lesion length ≤5.0 mm, along with reference vessel diameters ≥2.5 mm to ≤3.5 mm in the side branch and ≥2.5 mm to ≤4.0 mm in the main branch. The device is intended for use in conjunction with commercially available balloon expandable drug-eluting coronary stents in the main branch.

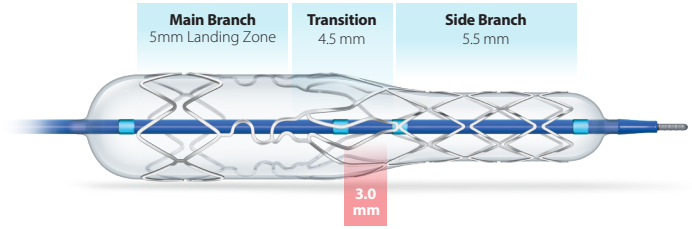
Technical Specifications

- Catheter System: Rapid Exchange (RX)
- Stent Material: Cobalt Chromium
- Strut Thickness: 85 µm
- Balloon Material: Semi-Compliant, Nylon
- Guidewire diameter: .014"
- Tip Profile: 0.020"
- Recommended Guide Catheter : 6 F (min I.D. 0.068" kissing balloon technique)
- Markers on proximal shaft: Brachial and femoral markers 90 cm and 100 cm from distal tip
- Balloon Radiopaque Markers: 4 - Platinum/Iridium
- Normal Pressure: 8 to 10 atm
- Radial Burst Pressure: 14 atm

2.5 mm Side Branch (19 mm)



3.0 - 3.5 mm Side Branch (15 mm)



Ordering Information

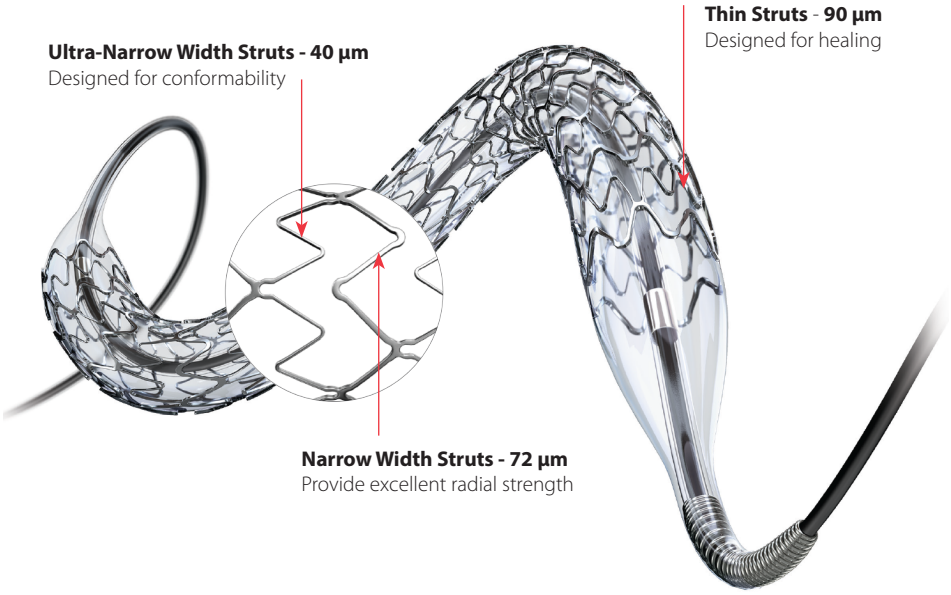
Product Codes	Diameter SB - MB (mm)	Length (mm)	Main Branch Landing Zone (mm)	Minimum Guiding Catheter Diameter	Maximum Post-expansion Diameter SB - MB (mm)	Nominal Pressure (atm)	RBP (atm)
2.5 mm Side Branch							
T52525191US	2.5 - 2.5	19	8	5 F	3.0 - 4.0	8	14
T52530191US	2.5 - 3.0	19	8	5 F	3.0 - 4.0	10	14
T52535191US	2.5 - 3.5	19	8	5 F	3.0 - 4.0	10	14
3.0 - 3.5 mm Side Branch							
T53035151US	3.0 - 3.5	15	5	6F	4.0 - 4.5	8	14
T53540151US	3.5 - 4.0	15	5	6F	4.0 - 4.5	10	14

EluNIR® Ridaforolimus Eluting Coronary Stent System

The EluNIR® DES was designed to help you navigate with ease, even in highly complex anatomies. The unique metallic spring tip, narrow width struts and reinforced hypotube offer excellent pushability, agility and flexibility.

Technical Specifications

- Catheter System: Rapid Exchange (RX)
- Balloon Material: Nylon 12
- Stent Material: CoCr alloy
- Strut Thickness: 90 µm
- Strut Width: Ultra-Narrow 40 µm, Narrow 72 µm
- Drug: Ridaforolimus
- Coating: Poly n-Butyl Methacrylate (PBMA) polymer, Carbosil® 20 55D
- Guidewire diameter: .014"
- Guide Catheter Compatibility: ≥ 5F
- Catheter Working Length: 140 cm
- Proximal Shaft Diameter: 2.1 F (0.69 mm) for all diameters
- Distal Shaft Diameter: 2.7 F (0.9 mm) for stent length 8 - 28 mm, 2.9 F (0.97 mm) for stent length 33 mm
- Nominal Pressure (NP): 10 atm for all diameters
- Rated Burst Pressure (RBP): 18 atm for all diameters
- Tip Characteristics: Stainless Steel Spring Tip
- Tip Entry Profile: 0.59 mm



Balloon Diameter	Balloon Length							
	8	12	15	17	20	24	28	33
2.5	LUN250R08US	LUN250R12US	LUN250R15US	LUN250R17US	LUN250R20US	**LUN250R24US**	**LUN250R28US**	**LUN250R33US**
2.75	LUN275R08US	LUN275R12US	LUN275R15US	LUN275R17US	LUN275R20US	LUN275R24US	LUN275R28US	LUN275R33US
3	LUN300R08US	LUN300R12US	LUN300R15US	LUN300R17US	LUN300R20US	LUN300R24US	LUN300R28US	LUN300R33US
3.5	LUN350R08US	LUN350R12US	LUN350R15US	LUN350R17US	LUN350R20US	LUN350R24US	LUN350R28US	LUN350R33US
4	LUN400R08US	LUN400R12US	LUN400R15US	LUN400R17US	LUN400R20US	LUN400R24US	LUN400R28US	LUN400R33US

Standard Biopsy Forceps

Key Features

- Choice of two forceps diameters (for taking samples adapted to a wide range of clinical situations)
  - 2.46 mm^3 of tissue sample using the 5.5F forceps
  - 5.20 mm^3 of tissue sample using the 7F forceps
- Jugular or femoral access possible with two lengths of forceps
- Units per package: 1

Shaft OD (French)	Shaft OD (mm)	Length (cm)	Description	Product Code
5.5	1.85	104	For femoral approach	504300
5.5	1.85	50	For internal jugular approach	504302
7	2.3	104	For femoral approach	504300L
7	2.3	50	For internal jugular approach	504302L

For information on indications, contraindications, warnings, and precautions, see page 82.

BI-PAL® Biopsy Forceps

Key Features

- Disposable and torquable (PTFE sheath for curving the distal section of the forceps into the desired shape and to direct it towards the ventricular wall)
  - 5.03mm^3 of tissue sample using the 7F forceps
- Jugular or femoral access possible with two lengths of forceps

Shaft OD (French)	Shaft OD (mm)	Length (cm)	Description	Product Code
7	2.3	104	Straight tip for femoral approach	502400B
7	2.3	50	Radial tip shape for internal jugular approach	502402B
7	2.3	50	Multipurpose tip shape for internal jugular approach	502402M

Biopsy Forceps Catheter Sheath Introducers and Sheath Sets

For Use With Catheter French (F)	Sheath Length (cm)	Description	Product Code
7	45	Sheath only, straight tip	501611
7	98	Sheath only, straight tip	501613
7	45	Sheath only, multipurpose curve	501611A
7	98	Sheath only, multipurpose curve	501613A
7	45	Sheath only, multipurpose curve, 50 cm 7F multipurpose A2 catheter, 8F vessel dilator	501616A
7	98	Sheath only, straight tip, 110 cm 7F pigtail HF catheter, 8F vessel dilator	501617

For information on indications, contraindications, warnings, and precautions, see page 76.

CORDIS® Closure Portfolio

Our closure portfolio includes the MYNX ACE®, MYNXGRIP®, and EXOSEAL® Vascular Closure Devices. MYNX ACE® and MYNXGRIP® Vascular Closure Devices utilize the proprietary GRIP™ sealant to seal the arteriotomy. The GRIP™ sealant, comprised of Polyethylene Glycol (PEG), grips the artery, providing a secure close. The sealant dissolves within 30 days, leaving nothing permanently behind but a healed artery. MYNX® Closure Devices treat a wide range of patients and clinical scenarios including punctures at or below the bifurcation and antegrade punctures. The versatile design provides options in challenging anatomies.

MYNXGRIP® Vascular Closure Device

The MYNXGRIP® Closure Device offers a patient-friendly closure option with no cinching, suturing, or metal implants. Learn more on page <OT>.

MYNX ACE® Vascular Closure Device

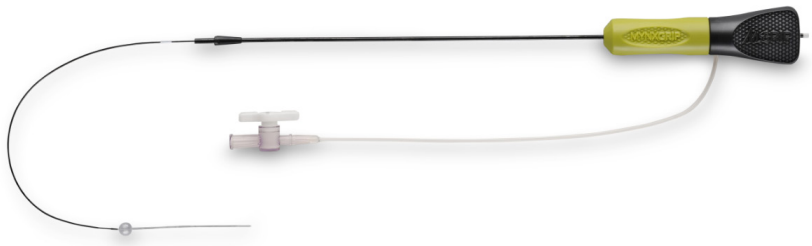
The MYNX ACE® Closure Device combines the reliability of an easy to use deployment system with the security of mechanical closure. Learn more on page 72.

EXOSEAL® Vascular Closure Device

Combining safety and ease-of-use, the Cordis EXOSEAL® Vascular Closure Device means a confident close and excellent patient outcomes. Learn more on page 73.

MYNXGRIP® Vascular Closure Device

The MYNXGRIP® Device provides secure mechanical closure with the safety of an extravascular sealant. The MYNXGRIP® Device contains the proprietary GRIP™ Sealant which actively adheres to and seals the arteriotomy or venotomy while expanding to fill the tissue tract. The MYNXGRIP® Device offers a patient-friendly closure option with no cinching, suturing, or metal implants. The GRIP™ sealant dissolves within 30 days leaving nothing permanently behind but a healed artery. The MYNXGRIP® Device is indicated to close femoral arterial and venous access sites utilizing a 5F, 6F, or 7F procedural sheath.



Ordering Information

The MYNXGRIP® Vascular Closure Device includes:

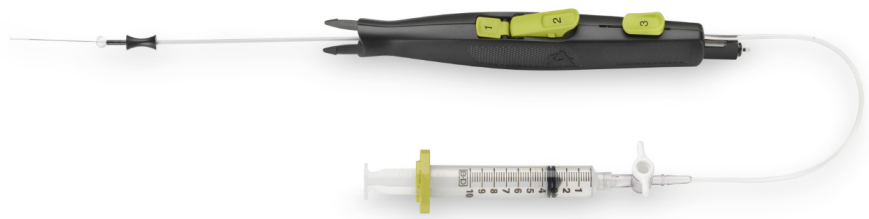
- Balloon catheter with integrated sealant
- 10 ml locking syringe
- Units per package: 10

Product	Size	Product Code
MYNXGRIP® Vascular Closure Device	5F	MX5021
MYNXGRIP® Vascular Closure Device	6F/7F	MX6721

MYNXGRIP® Vascular Closure Devices are manufactured by Cardinal Health and are part of the Cordis portfolio.
For information on indications, contraindications, warnings, and precautions, see page 81.

MYNX ACE® Vascular Closure Device

The MYNX ACE® Device is the latest addition to the MYNX® Product Family of Vascular Closure Devices. Combining the reliability of an easy to use deployment system with the security of mechanical closure and safety of an extravascular sealant, the MYNX ACE® device provides closure you can count on. The GRIP™ sealant securely adheres to the arteriotomy and dissolves within 30 days, leaving nothing permanently behind but a healed artery. The easy deployment and safety features of the MYNX ACE® Device help ensure proper use for a consistent close.* The MYNX ACE® Device is indicated to close femoral arterial access sites utilizing a 5F, 6F, or 7F procedural sheath.



Ordering Information

The MYNX ACE® Vascular Closure Device includes:

- (1) MYNX ACE® Device including balloon catheter and integrated sealant
- (1) 10 ml locking syringe
- (1) Introducer
- (1) Dilator
- Units per package: 10

Product	Size	Order Number
MYNX ACE® Vascular Closure Device	5F/6F/7F	MX6740

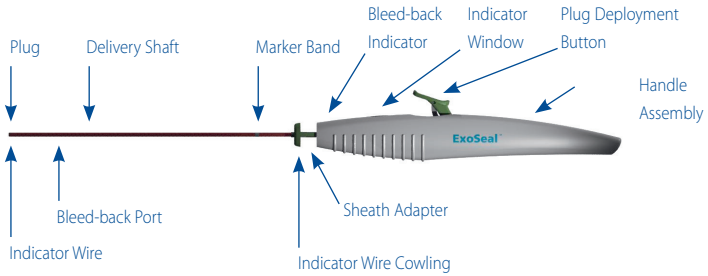
* Data on file at Cardinal Health. MYNX ACE® Vascular Closure Devices are manufactured by Cardinal Health and are part of the Cordis portfolio.
For information on indications, contraindications, warnings, and precautions, see page 84.

EXOSEAL® Vascular Closure Device

The EXOSEAL® Vascular Closure Device is indicated for femoral artery puncture site closure, reducing times to hemostasis and ambulation in patients who have undergone diagnostic or interventional catheterization procedures using a standard 5F, 6F, or 7F vascular sheath introducer with up to a 12-cm working length. The EXOSEAL® Vascular Closure Device is designed for a safe, simple, and secure close.

Key Features

- No anchor left inside the artery
- Two unique visual indicators enable precise positioning
- Easy-to-learn deployment helps efficiently achieve procedural success
- Simple 3-step procedure
- Available in 3 French sizes



Product	Size	Order Number
EXOSEAL® Vascular Closure Device	5F	EX500
EXOSEAL® Vascular Closure Device	6F	EX600
EXOSEAL® Vascular Closure Device	7F	EX700

For information on indications, contraindications, warnings, and precautions, see page 78.

ADROIT® Guiding Catheter

Indications
The guiding catheter is intended for use for intravascular introduction of interventional/diagnostic devices into the coronary or peripheral vascular systems.

Contraindications
None known for guiding catheters.

Warnings
Risk of reuse: This product is designed and intended for single use. It is not designed to undergo reprocessing and

re-sterilization after initial use. Reuse of this product, including after reprocessing and/or re-sterilization, may cause a loss of structural integrity which could lead to a failure of the device to perform as intended and may lead to a loss of critical labeling/use information all of which present a potential risk to patient safety.

Do not use with Ethiodol™ or Lipiodol™ contrast media, or other such contrast media which incorporates the components of these agents.

Precautions
- Store in a cool, dark, dry place.
- Do not use open or damaged packages.
- Use prior to the "Use By" date.
- Do not resterilize.
- Do not expose to organic solvents.
- Inspect the guiding catheter before use to verify that its size, shape and condition are suitable for the specific procedure.
- If strong resistance is met during manipulation, discontinue the procedure and determine the cause of the resistance before proceeding. If the cause of the resistance can not be determined, withdraw the catheter.
- Torquing the guiding catheter excessively while kinked may cause damage which could result in possible separation along the catheter shaft. Should the guiding catheter shaft become severely kinked, withdraw the entire system (guiding catheter, guidewire and catheter sheath introducer).
- Advancement, manipulation and withdrawal of the guiding catheter should always be performed under fluoroscopic guidance.
- Extreme care must be taken to avoid damage to the vasculature through which the guiding catheter passes. The guiding catheter may occlude smaller vessels. Care must be taken to avoid complete blood flow blockage.
- Large internal lumen guiding catheters require less force on the syringe during injection.

Complications
Procedures requiring percutaneous catheter introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at anytime during or after the procedure.

Possible complications include, but are not limited to the following:
- air embolism
- hematoma at the puncture site
- infection
- perforation of the heart
- vessel damage, dissection or perforation
- vasospasm
- ischemia
- hemorrhage
- arrhythmia
- reaction to contrast media
- death

Please refer to the Instructions for Use for complete information, including Adverse Events.

ATW™ Steerable Guidewire, ATW™ Marker Wire, STABILIZER® Plus Steerable Guidewire, STABILIZER® XS Steerable Guidewire, SHINOBI® Steerable Guidewire, SHINOBI® Plus Steerable Guidewire, REFLEX® Steerable Guidewire, WIZDOM™ Steerable Guidewire

Indications
Cordis Steerable Guidewires are intended for use in angiographic procedures to introduce and position catheters and interventional devices within the vasculature.

Contraindications
Cordis Steerable Guidewires are contraindicated for use in chronic total occlusions.

Contraindications for interventional devices are described in the instructions supplied with the respective device.

Warnings
Do not reuse. Discard after one procedure. Structural integrity and/or function may be impaired through reuse or cleaning. All parts are extremely difficult to clean after exposure to biological materials and may cause adverse patient reactions if reused. Cordis Corporation will not be responsible for any direct, incidental or consequential damages resulting from reuse of the product.

Guidewires are delicate instruments and should be handled carefully. Prior to use and when possible during the procedure, inspect the guidewire carefully for coil separation, bends, or kinks. Do not use a guidewire that shows signs of damage. Damage will prevent the guidewire from performing with accurate torque response and control.

- Guidewire manipulation/torquing should always be performed under fluoroscopic guidance.
- Never push, auger, withdraw, or torque a guidewire that meets resistance. First, using fluoroscopy, determine the cause of resistance and take any necessary remedial action. Torquing or pushing a guidewire against resistance may cause guidewire damage, and/or guidewire tip separation, or direct damage to the vessel. Resistance may be felt and/or observed (via fluoroscopy) by noting any buckling of the guidewire tip. If guidewire tip prolapse is observed, DO NOT allow the tip to remain in a prolapsed position; otherwise damage to the guidewire may occur.
- If any resistance is felt, i.e., due to vessel spasm, bent guidewire, or guidewire entrapment, while manipulating or removing the guidewire in the blood vessel: STOP the procedure. DO NOT move or torque the guidewire. Using fluoroscopy, first determine the cause of the resistance, then take appropriate remedial action. If the guidewire is moved excessively, it may break or become damaged. This may cause blood vessel injury or result in fragments being left inside the vessel.
- Should torque control/tip response be compromised during use, confirm tip integrity using fluoroscopy. LOSS OF TORQUE CONTROL MAY BE DUE TO CORE WIRE FRACTURE. Under fluoroscopic guidance, advance the balloon catheter to the distal end of the guidewire and remove the balloon catheter/guidewire system as a unit.

Precautions
- Store in a cool, dark, dry place.
- Do not use open or damaged packages.
- Use prior to the "Use By" date.
- Exposure to temperatures above 54°C (130°F) may damage the guidewire.
- Do not expose to organic solvents.
- Movement of torque device or metal insertion tool on a guidewire's coating may compromise the integrity of the coating.

Complications
Procedures requiring percutaneous guidewire introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at anytime during or after the procedure.

Possible complications include, but are not limited to:
- Air embolism
- Hematoma at the puncture site
- Infection
- Perforation of the heart or vessel wall
- Tip fractures have been reported in procedures involving guidewire entrapment, total occlusions, highly tortuous vasculature, and small side branches. For guidewire tip retrieval, please refer to the referenced publications for recommended techniques

Please refer to the Instructions for Use for complete information, including Adverse Events.

AVANTI®+ Sheath Introducer

Indications
The CSI is indicated for use in arterial and venous procedures requiring percutaneous introduction of intravascular devices.

Contraindications
None known.

Warnings
For one use only. Do not resterilize or reuse. Structural integrity and/or function may be impaired through cleaning, resterilization or reuse and may cause adverse patient reactions. Accordingly, Cordis will not be responsible for any direct or consequential damages or expenses resulting from reuse of the CSI.

Do not use with Ethiodol™ or Lipiodol™contrast media, or other such contrast media, which incorporate the components of these agents.

Do not leave a CSI in place for extended periods of time without a catheter or an obturator to support the cannula wall.

Precautions
- Store in a dry, dark, cool place.
- Do not use if package is open or damaged.
- Note "Use Before" or "Use By" date prior to using product.
- Do not resterilize. Exposure to temperatures above 54°C (130°F) may damage the catheter sheath and components.
- Do not expose to organic solvents, e.g. alcohol.
- If increased resistance is felt upon insertion of the CSI, investigate the cause before continuing. If the cause of the resistance cannot be determined and corrected, discontinue the procedure and withdraw the CSI.

Complications
Possible complications include, but are not limited to:
- air embolism
- infection
- intimal tear
- hematoma at the puncture site
- perforation of the vessel wall
- thrombus formation

Please refer to the Instructions for Use for complete information, including Adverse Events.

BI-PAL® Biopsy Forceps

Indications
The Cordis biopsy forceps are designed for endomyocardial biopsies.

Contraindications
None known.

Clinical History
These instructions are based on experience gathered to date. The physician may wish to vary the procedure in accordance with clinical judgement.

Warnings
Discard the forceps after completing one procedure. Structural integrity and/or function may be impaired through reuse or cleaning.

Forceps are extremely difficult to clean after exposure to biological material and may cause adverse patient reactions if reused.

Precautions
- The forceps should be thoroughly rinsed with heparinized saline before and after each biopsy during the procedure.
- The heart should be routinely monitored by ECG during the procedure.
- Use prior to the "Use By" date.
- Do not use if the inner package is open or damaged.
- Consider the use of systemic heparinization.
- If strong resistance is met during manipulation, discontinue the procedure and determine the cause of resistance before proceeding. If the cause of resistance cannot be determined, withdraw the forceps.

Complications
Procedures requiring biopsy forceps should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at any time during or after the procedure.

Possible complications include, but are not limited to:
- hematoma at the puncture site
- infection
- perforation of the vessel wall or the myocardium
- vessel trauma
- embolism
- death

Please refer to the Instructions for Use for complete information, including Adverse Events.

EluNIR® Ridaforolimus Eluting Coronary Stent System

Indications
The EluNIR® Ridaforolimus Eluting Coronary Stent System is indicated for improving coronary luminal diameter in patients with symptomatic heart disease due to de novo lesions ≤30mm in length in native coronary arteries with reference diameters of 2.50mm to 4.25mm.

Contraindications
Coronary artery stenting is generally contraindicated in the following patient types:
- Patients who cannot receive recommended antiplatelet and/or anticoagulation therapy.
- Patients judged to have a lesion which prevents complete inflation of an angioplasty balloon or proper placement of the stent or delivery system.
- Patients with hypersensitivity or allergies to aspirin, heparin, clopidogrel, ticlopidine, drugs such as ridaforolimus or similar drugs, the polymer or its individual components CarboSil® 20 55D (Thermoplastic Silicone-Polycarbonate-urethane) and Poly n-Butyl Methacrylate (PBMA), cobalt, chromium, nickel, molybdenum, or contrast media.

Warnings
- Please ensure that the inner package has not been opened or damaged as this would indicate that the sterile barrier has been breached.
- The use of this device carries the associated risks of thrombosis, vascular complications and/or bleeding events.
- This product should not be used in patients who are not likely to comply with the recommended antiplatelet therapy.

Precautions
- Stent implantation should only be performed by physicians who have received appropriate training.
- Subsequent restenosis may require repeat dilatation of the arterial segment containing the stent. Long-term outcomes following repeat dilatation of the stent is presently not well characterized.
- Risks and benefits should be considered in patients with severe reaction to contrast agent.
- Patients with known hypersensitivity to the product components (stent substrate, polymer(s), drug substance) may suffer an allergic reaction to this implant.
- Do not expose or wipe the product with organic solvents such as alcohol.
- Care should be taken to control the guiding catheter tip during stent delivery, deployment, and balloon withdrawal. Before withdrawing the stent delivery system, visually confirm complete balloon deflation by fluoroscopy to avoid guiding catheter movement into the vessel and subsequent arterial damage.
- Stent thrombosis is a low-frequency event that is frequently associated with myocardial infarction (MI) or death.
- When DES are used outside the specified Indications for Use, patient outcomes may differ from the results observed in the EluNIR® clinical trials.

- Compared to use within the specified Indications for Use, the use of DES in patients and lesions outside of the labeled indications, including more tortuous anatomy, may have an increased risk of adverse events, including stent thrombosis, stent embolization, MI, or death.

Complications
Complications may include bleeding, hematoma, pseudoaneurysm, or vessel perforation.

Please refer to the Instructions for Use for complete information, including Adverse Events.

EMERALD® Diagnostic Guidewire

Indications
Cordis Guidewires are intended for use in the percutaneous introduction of catheters.

Contraindications
None known.

Warnings
Do not reuse. Discard after one procedure. Structural integrity and/or function may be impaired through reuse or cleaning. All parts are extremely difficult to clean after exposure to biological materials and may cause adverse patient reactions if reused. Accordingly, Cordis Corporation will not be responsible for any direct, incidental or consequential damages resulting from reuse of the product.

Precautions
- Store in a cool, dark, dry place.
- Do not use if package is open or damaged.
- Use prior to "Use By" date.
- Do not expose to organic solvents.
- Exposure to temperatures above 54°C (130°F) may damage the components.
- Do not withdraw a PTFE coated guidewire through a metal-cannula needle. Withdrawal may damage the guidewire coating.
- If strong resistance is met during manipulation, discontinue the procedure and determine the cause of resistance before proceeding. If the cause of the resistance cannot be determined, withdraw the catheter and guidewire.

Complications
Procedures requiring percutaneous catheter/guidewire introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at any time during or after the procedure.

Possible complications include, but are not limited to:
- air embolism
- hematoma at the puncture site
- infection
- perforation of the vessel wall

Please refer to the Instructions for Use for complete information, including Adverse Events.

EMPIRA® Balloon Catheter

Indications
The Cordis EMPIRA® RX PTCA Dilation Catheter and EMPIRA NC® RX PTCA Dilatation Catheter are indicated for balloon dilatation of the stenotic portion of a coronary artery or bypass graft stenosis for the purpose of improving myocardial perfusion.

The Cordis EMPIRA NC® RX PTCA Dilatation Catheter is also indicated for post-delivery expansion of balloon expandable stents.

Note: In vitro testing was performed with the EMPIRA NC® RX PTCA

Dilatation Catheter and with commercially available Cordis balloonexpandable stents. Caution should be taken when using this device with stents of other manufacturers, due to stent design differences.

Contraindications
- Unprotected left main coronary artery lesions.
- Coronary artery spasm in the absence of a significant stenosis.

Warnings
- Use extreme caution and careful judgment in patients for whom anticoagulation is contra-indicated.
- The catheter is supplied STERILE. Do not use if the sterile barrier is damaged.
- For single patient use only. Do not reuse, reprocess or resterilize. Reuse, reprocessing or resterilization may compromise the structural integrity of the device and/or lead to device failure, which in turn, may result in increased risk of cross contamination, patient injury, illness or death.
- To reduce the potential for vessel damage, the inflated diameter of the balloon should approximate the diameter of the vessel just proximal and distal to the stenosis.
- PTCA in patients who are not acceptable candidates for coronary artery bypass graft surgery requires careful consideration, including possible hemodynamic support during the procedure, as treatment of this patient population carries special risk.
- When the catheter is exposed to the vascular system, it should be manipulated while using high-quality fluoroscopy. Do not advance or retract the catheter unless the balloon is fully deflated by vacuum. If resistance is felt during manipulation, determine the cause of the resistance before proceeding.

- Balloon pressure should not exceed the rated burst pressure. The rated burst pressure is based on the results of in vitro testing. At least 99.9% of the balloons (with a 95% confidence), will not burst at or below their rated burst pressure. Use of a pressure monitoring device is recommended to prevent over pressurization.
- PTCA should only be performed at hospitals where emergency coronary artery bypass graft surgery can be quickly performed in the event of a potentially injurious or life-threatening complication.
- To reduce the potential for air embolus into the vessel, use 50% solution of contrast medium diluted with sterile heparinized-saline.
- During withdrawal of the PTCA catheter, hold a heparinizedsaline soaked gauze around the exposed catheter shaft and pull the catheter through the gauze to remove blood or any other residues.
- Care should be taken when handling the distal part of the catheter (including the balloon) to prevent damages and prematurely removing balloon cover.

Precautions

- Prior to angioplasty, the catheter should be examined to verify functionality and ensure that its size and shape are suitable for the specific procedure for which it is to be used.
- The compatibility of the device has not been evaluated for the delivery of materials (e.g., drugs, alcohol or stem cells) through the guidewire lumen, other than those required for normal use.
- The catheter system should be used only by physicians trained in the performance of PTCA.
- Appropriate anticoagulant/antiplatelet therapy should be used during this procedure.
- Prior to insertion or withdrawal of the PTCA catheter, wipe the guidewire with heparinized-saline soaked gauze to remove blood or residues thus providing better catheter movement over the guidewire.
- Care should be taken to control the position of the guide catheter tip during manipulation of the balloon catheter.
- Use the catheter before the "Use By" date specified on the package.
- The safety and effectiveness of this PTCA balloon catheter for the treatment of in-stent restenosis (ISR) has not been established.

Please refer to the Instructions for Use for complete information, including Adverse Events.

EXOSEAL® Vascular Closure Device

Indication for Use

The EXOSEAL® Vascular Closure Device is indicated for femoral artery puncture site closure, reducing times to hemostasis and ambulation in patients who have undergone diagnostic or interventional catheterization procedures using a standard 5F, 6F, or 7F vascular sheath introducer with up to a 12 cm working length. Additionally, the EXOSEAL® Vascular Closure Device is indicated to reduce times to hemostasis and ambulation in patients who have undergone interventional catheterization procedures, using a standard 6F vascular sheath introducer up to a 12 cm working length, who have received preprocedural and/or intraprocedural glycoprotein (GP) IIb-IIIa inhibitor therapy.

Contraindications

There are no contraindications to the use of this device. Attention is drawn to the Warnings, Precautions, and Special Patient Populations.

Warnings

- Do not use the EXOSEAL® Vascular Closure Device if the package is damaged or any portion of the package has been previously opened.
- Do not use the EXOSEAL® Vascular Closure Device if the device appears damaged or defective in any way.
- Do not use the EXOSEAL® Vascular Closure Device if the sterile field has been broken where bacterial contamination of the sheath or surrounding tissues may have occurred; a broken sterile field may result in infection.
- For SINGLE USE ONLY. Do not resterilize or reuse. Reuse, reprocessing, or resterilization may compromise the structural integrity of the device and/or lead to device failure, which in turn may result in patient injury, illness, or death. Use aseptic technique when handling the product.
- Do not use the EXOSEAL® Vascular Closure Device in patients with known allergy to polyglycolic acid.

Precautions

- Serious adverse events might result with the use of the EXOSEAL® Vascular Closure Device in vessels not suitable for the use of the device. Avoid the use of the EXOSEAL® Vascular Closure Device in patients with arteriotomies created in areas of calcified plaque or in vessels with diameters < 5mm.
- With antegrade puncture (restricted to peripheral vascular catheterization procedures), the ability to accurately assess vessel size or extraluminal device position may be limited.
- The EXOSEAL® Vascular Closure Device procedure should be performed by physicians who have expertise in the techniques of vascular catheterization (or other health care professionals authorized by, or under the direction of, such physicians) and possess adequate training in the use of the device, eg, participation in an EXOSEAL® Vascular Closure Device training program.
- Observe sterile technique at all times when using the EXOSEAL® Vascular Closure Device. Employ proper groin management postprocedure and posthospital discharge to prevent infection.

- The vascular sheath introducer and/or EXOSEAL® Vascular Closure Device should not be advanced or withdrawn when resistance is met without first determining the cause by fluoroscopic examination. Using excessive force to advance or torque the EXOSEAL® Vascular Closure Device may lead to arterial damage and/or breakage of the device, which may necessitate interventional and/or surgical removal of the device and arterial repair.
- If for any reason it is desired to abort the procedure once the EXOSEAL® Vascular Closure Device has been introduced into the bloodstream, remove the EXOSEAL® Vascular Closure Device and vascular sheath introducer as a unit. Do not attempt to withdraw the EXOSEAL® Vascular Closure Device from the vascular sheath introducer, as Plug dislodgement may occur.
- Pulsatile flow is necessary for proper positioning. If pulsatile flow is not observed from the Bleed-Back Indicator, discontinue the procedure.
- Do not remove the EXOSEAL® Vascular Closure Device from the vascular sheath introducer after removal from the patient; discard the EXOSEAL® Vascular Closure Device with the Delivery Shaft still locked inside the vascular sheath introducer.
- In patients undergoing interventional endovascular procedures, ambulation less than 2 hours after EXOSEAL® Vascular Closure Device use increases the risk of oozing or rebleeding after initial hemostasis and should be done only after all clinical factors have been considered.

Special Patient Population

The safety and effectiveness of the EXOSEAL® Vascular Closure Device has not been established in the following patient populations:

- Patients with acure ST-elevation myocardial infarction ≤ 48 hours prior to the cardiac or peripheral catheterization
- Patients with uncontrolled hypertension at time of closure (BP ≥180/110 mmHg)
- Patients who bruise or bleed easily or with a history of significant bleeding or platelet disorders, such as thrombocytopenia (with < 100,000 platelet count), Von Willebrand's disease, anemis (Hgb < 10 g/dL, Hct < 30%), thrombasthenia, decreased fibrinogen (<200 mg/dl), and Factory V deficiency
- Patients with prior femoral vascular surgery or vascular graft in region of access site
- Patients with pre-existing systemic or cutaneous infection
- Patients who are known to be pregnant or who are lactating
- Patients on thrombolytic (e.g. streptokinase, urokinase, t-PA) ≤ 24 hours prior to the catheterization procedure
- Patients on Angiomax (bivalirudin) or other thrombin-specific anticoagulants or low molecular weight heparin ≤ 24 hours prior to the cardiac or peripheral catheterization procedure
- Patients with a BMI > 40 Kg/m2
- Patients with symptomatic leg ischemia in the target vessel limb including severe claudication (30.48 meters / < 100 feet) or weak/absent pulse
- Patients with planned arterial access at the same access site ≤ 30 days following the femoral artery closure procedure
- Patients undergoing arterial puncture in the femoral artery or both legs
- Patients with prior target artery closure with any closure device, or closure with manual compression ≤ 30 days prior to the cardia or peripheral catheterization procedure
- Patients with prior or recent use of an intra-aortic balloon pump through the arterial access site
- Patients with evidence of a preexisting hematoma, arteriovenous fistula, or pseudoaneurysm at the access site prior to start of femoral artery closure procedure
- Patients with a tortuous targeted femoral artery
- Patients who within ≤ 1 cm of the puncture site have fluoroscopically visible calcium, atherosclerotic disease, or a stent
- Patients with a targeted femoral artery diameter stenosis ≥ 50%
- Patients with arteriotomies in vessels with diameters < 5mm
- Patients where there is difficulty in obtaining vascular access resulting in multiple arterial punctures and/or posterior arterial puncture
- Patients with antegrade puncture
- Heparinized patients with elevated pre-closure ACT level: 250 seconds with GP IIb/IIIa inhibitor, > 300 seconds without GP IIb/IIIa inhibitor
- Patients experiencing cardiogenic shock (hemodynamic instability requiring intravenous medications or mechanical support) during or immediately post-catheterization

Please refer to the Instructions for Use for complete information, including Adverse Events.

7F HIGHFLOW™ Diagnostic Catheter, INFINITI® Diagnostic Catheter, TEMPO AQUA® Diagnostic Catheter

Indications

Cordis catheters are designed to deliver radiopaque contrast medium to selected sites in the vascular system.

Contraindications

None known.

Warnings

- Discard catheters after one procedure. Structural integrity and/or function may be impaired through reuse or cleaning. All parts are extremely difficult to

clean after exposure to biological materials and may cause adverse patient reactions if reused.
- Do not expose to organic solvents.
- Do not use with Ethiodol™ or Lipiodol™ contrast media, or other such contrast media which incorporates the components of these agents.
- Do not exceed maximum pressure rating printed on product label and hub.

Precautions
- Store in a cool, dark, dry place.
- Do not use if package is open or damaged.
- Do not use the catheter if the "Use By" date on the package label has expired.
- Do not resterilize.
- Exposure to temperatures above 54°C (130°F) may damage the catheter.
- To prevent damage to the catheter tip during removal from the package, grasp the hub and withdraw the catheter.
- Exercise care when removing guidewires from multiple-curve catheters.
- To prevent kinking of 5F (1.65 mm) and smaller angiographic catheters, and specifically the 4F (1.35 mm) INFINITI® pigtail catheters:
  - Straighten the pigtail catheter tip only with a diagnostic guidewire or, if applicable, with a tip straightener. Do not straighten by hand.
  - Use a guidewire when introducing the catheter through the catheter sheath introducer (CSI) and into the left ventricle.
- Treat all 4F (1.35 mm) catheters and smaller French sizes with ultimate care. The performance of these products may be impaired if not properly and cautiously handled during unpacking and preparation.
- Before use, flush all devices entering a blood vessel with sterile heparinized saline or a similar isotonic solution.
- Keep the catheter filled with either flushing solution or contrast medium while the catheter is in the vascular system and consider the use of systemic heparinization.
- Forcibly aspirate and flush the catheter with heparinized saline solution at least once every two minutes.

Complications
Procedures requiring percutaneous catheter introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at any time during or after the procedure.

Possible complications include, but are not limited to the following:
- air embolism
- hematoma at the puncture site
- infection
- perforation of the vessel wall

MOZEC™ RX PTCA Balloon Dilatation Catheter

Indications
The MOZEC™ Rx PTCA Balloon Dilatation Catheter is indicated for balloon dilatation of the stenotic portion of a coronary artery or bypass graft stenosis for the purpose of improving myocardial perfusion.

The MOZEC™ Rx PTCA Balloon Dilatation Catheter (balloon models 2.25nun to 4.50nun) is also indicated for post-delivery expansion of balloon expandable stents.

Note: Bench testing was conducted with the MOZEC™ Rx PTCA Balloon Dilatation Catheter and marketed balloon expandable stents (viz. Medtronic Integrity and Abbott's Multi Link Stents). Consideration should be taken when this device is used with different manufacturer's stents due to differences in stent design. All stents should be deployed in accordance with the manufacturer's indications and instructions for use.

Contraindications
Unprotected left main coronary artery lesion. Coronary artery spasm in the absence of a significant stenosis. Patients with a contraindication for anti-platelet/ anticoagulant therapy.

Warnings
- For single patient use only. Do not reuse, reprocess or resterilize. Reuse, reprocessing or re-sterilization may compromise the structural integrity of the device and/ or lead to device failure which, in tum, may result in patient injury, illness or death.
- Reuse, reprocessing or re-sterilization may also create a risk of contamination of the device and/ or cause patient infection or cross-infection, including, but not limited to the transmission of infectious disease(s) from one patient to another. Contamination of the device may lead to injury, illness or death of the patient.
- PTCA should only be performed at hospitals where emergency coronary artery bypass graft surgery can be quickly performed in the event of a potentially injurious or life-threatening complication.
- Since use of this device carries the associated risk of sub-acute thrombosis, vascular complications and/ or bleeding events, judicious selection of patients is necessary.

Note: Animal testing in canines showed thrombus formation along the catheter length when anticoagulation was not used.

- PTCA in patients who are not acceptable candidates for coronary artery bypass graft surgery requires careful consideration, including potential haemodynamic support during the procedure, as treatment of this patient population carries special risk.

- Guiding catheters used must have lumen sizes that are suitable to accommodate the introduction of MOZEC™ Rx PTCA Balloon Dilatation Catheter.
- Do not use if the inner package is open or damaged. Carefully remove the PTCA catheter from the pouch to prevent damage and premature removal of the balloon cover.
- To reduce the potential for air embolus into the vessel, use only the recommended balloon inflation medium. Never use air or any gaseous medium to inflate the balloon.
- When the catheter is exposed to the vascular system, it should be manipulated while under high-quality fluoroscopic observation. Do not advance or retract the catheter unless the balloon is fully deflated under vacuum. If resistance is felt during manipulation, determine the cause of the resistance before proceeding.
- Balloon pressure should not exceed the rated burst pressure. The rated burst pressure is based on the results of in vitro testing. At least 99.9 percent of the balloons, (with a 95 percent confidence) will not burst at or below their rated burst pressure. Use of a pressure monitoring device is recommended to prevent over pressurization.
- To reduce the potential for vessel damage, the inflated diameter of the balloon should approximate the diameter of the vessel just proximal and distal to the stenosis.

Precautions

- Prior to angioplasty, examine the PTCA catheter to verify functionality. Ensure that its size and shape are suitable for the specific procedure for which it is to be used.
- Only the physicians trained in the performance of PTCA should use the catheter system. Appropriate anticoagulant/antiplatelet and vasodilator therapy should be used during the procedure.
- Prior to reinsertion or withdrawal of the PTCA catheter, wipe the guide wire with saline-soaked gauze to remove blood or other residues.
- After the procedure, anticoagulant therapy should be continued as recommended by the physician. Use the catheter before the "Use by" date specified on the package.

Adverse Events

Potential adverse events, which may be associated with the use of the MOZEC™ Rx PTCA Balloon Dilatation Catheter, include but are not limited to:

- Acute myocardial infarction Acute vessel closure
- Allergic reactions to anti-coagulant and /or antithrombotic therapy/ contrast medium Arrhythmia, including ventricular fibrillation (VF)
- Arteriovenous fistula Coronary embolism Coronary artery spasm Coronary aneurysm
- Coronary vessel dissection/injury/perforation/rupture Death
- Emergency or non-emergent Coronary Artery Bypass Graft Surgery Hematoma or Hemorrhage
- Hypotension I Hypertension
- Infection and / or pain at the access site Restenosis of treated segment
- Total occlusion of coronary artery/bypass graft Unstable angina pectoris
- Stroke, air embolism and embolization or fragmentation of thrombotic or atherosclerotic material

Please refer to the Instructions for Use for complete information.

MOZEC™ NC RX PTCA Balloon Dilatation Catheter

Indications

The MOZEC™ NC Rx PTCA Balloon Dilatation Catheter is indicated for balloon dilatation of the stenotic portion of a coronary artery or bypass graft stenosis for the purpose of improving myocardial perfusion. The MOZEC ™ NC Rx PTCA Balloon Dilatation Catheter is also indicated for post-delivery expansion of balloon expandable stents.

Note: Bench testing was conducted with the MOZEC™ NC Rx PTCA Balloon Dilatation Catheter and marketed balloon expandable stents (viz. Medtronic Integrity, Boston Scientifics Liberte & Abbott's Multi Link Stents). Consideration should be taken when this device is used with different manufacturer's stents due to differences in stent design. All stents should be deployed in accordance with the manufacturer's indications and instructions for use.

Contraindications

- Unprotected left main coronary artery lesion.
- Coronary artery spasm in the absence of a significant stenosis.

Warnings

- For single patient use only. Do not reuse, reprocess or resterilize. Reuse, reprocessing or re-sterilization may compromise the structural integrity of the device and/ or lead to device failure which, in turn, may result in patient injury, illness or death.
- Reuse, reprocessing or re-sterilization may also create a risk of contamination of the device and/ or cause patient infection or cross-infection, including, but not limited to the transmission of infectious disease(s) from one patient to another. Contamination of the device may lead to injury, illness or death of the patient.

- PTCA should only be performed at hospitals where emergency coronary artery bypass graft surgery can be quickly performed in the event of a potentially injurious or life-threatening complication.
- PTCA in patients who are not acceptable candidates for coronary artery bypass graft surgery requires careful consideration, including potential haemodynamic support during the procedure, as treatment of this patient population carries special risk.
- Guiding catheters used must have lumen sizes that are suitable to accommodate the introduction of MOZECTM NC Rx PTCA Balloon Dilatation Catheter.
- Do not use if the inner package is open or damaged. Carefully remove the PTCA catheter from the pouch to prevent damage and premature removal of the balloon cover.
- To reduce the potential for air embolus into the vessel, use only the recommended balloon inflation medium. Never use air or any gaseous medium to inflate the balloon.
- When the catheter is exposed to the vascular system, it should be manipulated while under high-quality fluoroscopic observation. Do not advance or retract the catheter unless the balloon is fully deflated under vacuum. If resistance is felt during manipulation, determine the cause of the resistance before proceeding.
- Balloon pressure should not exceed the rated burst pressure. The rated burst pressure is based on the results of in vitro testing.
- At least 99.9 percent of the balloons, (with a 95 percent confidence) will not burst at or below their rated burst pressure. Use of a pressure monitoring device is recommended to prevent over pressurization.
- To reduce the potential for vessel damage, the inflated diameter of the balloon should approximate the diameter of the vessel just proximal and distal to the stenosis.

Precautions

- Prior to angioplasty, examine the PTCA catheter to verify functionality. Ensure that its size and shape are suitable for the specific procedure for which it is to be used.
- Only the physicians trained in the performance of PTCA should use the catheter system.
- Appropriate anticoagulant/antiplatelet and vasodilator therapy should be used during the procedure.
- The safety and effectiveness of this PTCA balloon catheter for the treatment of ISR has not been established.
- Prior to reinsertion or withdrawal of the PTCA catheter, wipe the guide wire with saline-soaked gauze to remove blood or other residues.
- After the procedure, anticoagulant therapy should be continued as recommended by the physician.
- Use the catheter before the "Use by" date specified on the package.

Adverse Events

- Potential adverse events, which may be associated with the use of the Mozec™ NC - Rx PTCA Balloon Dilatation Catheter, include but are not limited to:
- Acute myocardial infarction
- Acute vessel closure
- Allergic reactions to anti-coagulant and /or antithrombotic therapy/ contrast medium
- Arrhythmia, including ventricular fibrillation (VF)
- Arteriovenous fistula
- Coronary embolism
- Coronary artery spasm
- Coronary aneurysm
- Coronary vessel dissection/injury/perforation/rupture
- Death
- Emergency or non-emergent Coronary Artery Bypass Graft Surgery
- Hematoma or Hemorrhage
- Hypotension / Hypertension
- Infection and / or pain at the access site
- Restenosis of treated segment
- Total occlusion of coronary artery/bypass graft
- Unstable angina pectoris
- Stroke, air embolism and embolization or fragmentation of thrombotic or atherosclerotic material

Please refer to the Instructions for Use for complete information.

MYNX ACE® Vascular Closure Device

Indications for Use:
The MYNX ACE® Device is indicated for use to seal femoral arterial access sites while reducing times to hemostasis and ambulation in patients who have undergone diagnostic or interventional endovascular procedures utilizing a 5F, 6F or 7F procedural sheath.

Precautions
The MYNX ACE® Device should only be used by a trained licensed physician or healthcare professional. The MYNX ACE® Device should not be used in patients with a known allergy to PEG.

Warnings
Do not use if components or packaging appear to be damaged or defective or if any portion of the packaging has been previously opened. DO NOT REUSE OR RESTERILIZE. The MYNX ACE® Device is for single use only. The catheter is loaded with a single hydrogel sealant. Reuse of the device would result in no delivery of hydrogel sealant. Do not use the MYNX ACE® Device if the puncture site is located above the most inferior border of the inferior epigastric artery (IEA) and/or above the inguinal ligament based upon bony landmarks, since such a puncture site may result in a retroperitoneal hematoma/ bleed. Perform a femoral angiogram to verify the location of the puncture site. Do not use the MYNX ACE® Device

if the puncture is through the posterior wall or if there are multiple punctures, as such punctures may result in a retroperitoneal hematoma/bleed.

Please refer to the Instructions for Use for complete information, including Adverse Events.

MYNXGRIP® Vascular Closure Device

Indications For Use
The MYNXGRIP® Device is indicated for use to seal femoral arterial and femoral venous access sites while reducing times to hemostasis and ambulation in patients who have undergone diagnostic or interventional endovascular procedures utilizing a 5F, 6F or 7F procedural sheath.

Precautions
The MYNXGRIP® Device should only be used by a trained licensed physician or healthcare professional. The MYNXGRIP® Device should not be used in patients with a known allergy to PEG.

Warnings
Do not use if components or packaging appear to be damaged or defective or if any portion of the packaging has been previously opened. DO NOT REUSE OR RESTERILIZE. The MYNXGRIP® Device is for single use only. The balloon catheter is loaded with a single hydrogel sealant. Reuse of the device would result in no delivery of hydrogel sealant. Do not use the MYNXGRIP® Device if the puncture site is located above the most inferior border of the inferior epigastric artery (IEA) (for arterial application) and/or above the inguinal ligament based upon osseus landmarks, since such a puncture site may result in a retroperitoneal hematoma/bleed. Perform a femoral angiogram or venogram to verify the location of the puncture site. Do not use the MYNXGRIP® Device if the puncture is through the posterior wall or if there are multiple punctures, as such punctures may result in a retroperitoneal hematoma/bleed.

Potential Adverse Events
In addition to the complications noted in the MYNX® Device clinical trial, the following potential complications, which may be related to the endovascular procedure or the vascular closure, may occur:
- allergic reaction
- ecchymosis
- superficial vein thrombosis
- foreign body/local reaction
- retroperitoneal bleed
- vessel occlusion
- pulmonary embolism
- death.

Please refer to the Instructions for Use for complete information, including Adverse Events.

NIRxcell™ CoCr Coronary Stent System

Indications
NIRxcell Stent System is indicated for improving coronary luminal diameter in patients with symptomatic ischemic heart disease associated with stenotic lesions in de novo native coronary arteries (length ≤ 30mm) with a reference vessel diameter of 2.50mm to 4.00mm.

Contraindications
Coronary artery stenting is generally contraindicated in the following patient types:

- Patients for whom antiplatelet and/or anticoagulation therapy is contraindicated
- Patients judged to have a lesion which prevents complete inflation of an angioplasty balloon or proper placement of the stent or delivery system

Warnings
- Since the use of this device carries the associated risks of thrombosis, vascular complications, and/or bleeding events, judicious selection of patients is necessary.
- Persons allergic to L-605 cobalt chromium alloy may suffer an allergic response to this implant.

Precautions
- Do not use with Ethiodol or Lipiodol contrast media*.
- Do not expose the delivery system to organic solvents such as alcohol or detergents.
- Only physicians who have received appropriate training should perform implantation of the stent.
- Stent placement should be performed only at hospitals where emergency coronary artery bypass graft surgery can be readily performed.
- Subsequent restenosis may require repeat dilatation of the arterial segment containing the stent. The long-term outcome following repeat dilatation of endothelialized coronary stents is not well characterized.
- When multiple stents are required, stent materials should be of similar composition. Placing multiple stents. of different materials in contact with each other may increase the potential for corrosion.
- The device should be manipulated while under high-quality fluoroscopic observation.
- Do not advance or retract the catheter unless the balloon is fully deflated. If resistance is met during manipulation, determine the cause of resistance under fluoroscopy before proceeding.
- Do not try to straighten a kinked hypotube. Straightening a kinked metal shaft may result in breakage of the shaft.

The risks and benefits should be considered for each patient before use of NIRxcell™ Stent System. Patient selection factors to be assessed should include a judgment regarding risk of antiplatelet therapy. Special

consideration should be given to those patients with recently active gastritis or peptic ulcer disease. Co-morbidities that increase the risk of poor initial results or the risks of emergency referral for bypass surgery (diabetes mellitus, renal failure, and severe obesity) should be reviewed.

Thrombosis following stent implantation is affected by several baseline angiographic and procedural factors. These include vessel diameter less than 3mm, intra-procedural thrombus, and dissection following stent implantation.

Complications
Complications may include bleeding, hematoma, pseudoaneurysm, or vessel perforation.

Please refer to the Instructions for Use for complete information, including Adverse Events.

Standard Biopsy Forceps

Indications
The Cordis biopsy forceps are designed for endomyocardial biopsies.

Contraindications
None known.

Clinical History
These instructions are based on experience gathered to date.

The physician may wish to vary the procedure in accordance with clinical judgement.

Warnings
Discard forceps after completing one procedure. Structural integrity and/or function may be impaired through reuse or cleaning. Forceps are extremely difficult to clean after exposure to biological material and may cause adverse patient reactions if reused.

Precautions
- The forceps should be thoroughly rinsed with heparinized saline before and after each biopsy during the procedure.
- The heart should be routinely monitored by ECG during the procedure.
- Use prior to the "Use By" date.
- Do not use if the inner package is open or damaged.
- Consider the use of systemic heparinization.
- If strong resistance is met during manipulation, discontinue the procedure and determine the cause of resistance before proceeding. If the cause of resistance cannot be determined, withdraw the forceps.
- Store in a cool, dark, dry place.

Complications
Procedures requiring biopsy forceps should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at any time during or after the procedure.

Possible complications include, but are not limited to:
- hematoma at the puncture site
- infection
- perforation of the vessel wall or the myocardium
- vessel trauma
- embolism
- death

Please refer to the Instructions for Use for complete information, including Adverse Events.

SUPER TORQUE® Diagnostic Catheter

Indications
Cordis Angiographic Catheters with Marker Bands are designed to provide angiographic visualization and linear measurement of the vasculature when combined with the delivery of radiopaque contrast media to selected sites in the vascular system.

Contraindications
None known.

Warnings
- Failure to observe these instructions may result in damage, breakage or separation of the catheter or the markerbands, which may necessitate additional intervention.
- Manipulation of the catheter under excessive friction due to interaction with other devices or while trapped in the vasculature, can lead to stretching or elongation of the catheter.
- Stretching or elongation of the catheter during endovascular procedures could result in the marker bands moving along the catheter. In extreme cases, marker bands may come off the catheter and dislodge into the vascular system.
- This product is designed and intended for single use. It is not designed to undergo reprocessing and re-sterilization after initial use. Reuse of this product, including after reprocessing and/or re-sterilization, may cause a loss of structural integrity which could lead to a failure of the device to perform as intended and may lead to a loss of critical labeling/use information all of which present a potential risk to patient safety.
- Do not expose to organic solvents.
- Do not exceed maximum pressure rating printed on label and hub.

Precautions

- Avoid entrapment of the catheter between other endovascular devices and the vessel wall.
- Avoid excessive friction on the catheter; avoid simultaneous introduction of the catheter and aortic graft devices through the same sheath.
- Store in cool, dark, dry place.
- Do not use if the package is open or damaged.
- Do not use the catheter if the "Use By" date on the package label has expired.
- Do not resterilize.
- Exposure to temperatures above 54°C (130°F) may damage the catheter.

Complications

Procedures requiring percutaneous catheter introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at any time during or after the procedure. Possible complications include, but are not limited to, the following: air embolism, hematoma at the puncture site, infection, thrombosis, hemorrhage, dissection, perforation or other damage of the vessel wall. Movement of the marker bands along the catheter can result in inaccurate reference and device sizing. Dislodgement of the marker bands into the vascular system can result in additional intervention, embolism, thrombosis or other vascular complications.

Please refer to the Instructions for Use for complete information, including Adverse Events.

TRYTON Side Branch Stent

Indications

The TRYTON Side Branch Stent is indicated for improving the side branch luminal diameter of de novo native coronary artery bifurcation lesions (Medina Classification 1.1.1; 0.1.1; 1.0.1) with a side branch diameter stenosis of ≥50% and a lesion length ≤5.0 mm, along with reference vessel diameters ≥2.5 mm to ≤3.5 mm in the side branch and ≥2.5 mm to ≤4.0 mm in the main branch.

The device is intended for use in conjunction with commercially available balloon expandable drug-eluting coronary stents in the main branch.

Contraindications

The TRYTON Side Branch Stent is contraindicated in the following conditions or uses:

- Vessels that are totally occluded
- Vessels that have moderate to severe calcification
- Target lesions that have excessive tortuosity unsuitable for stent delivery and deployment
- Angiographic evidence of thrombus in the target vessel
- Lesions in which complete angioplasty balloon inflation cannot be achieved during pre-dilatation
- TRYTON Stent placement without angioplasty pre-dilatation of the main branch and side branch (i.e., direct stenting is contraindicated)
- TRYTON Stent placement alone, without implantation of a main branch stent
- An untreated significant (> 50%) stenosis proximal or distal to the main branch or side branch target lesion
- Impaired runoff in the treatment vessel with diffuse distal disease
- Ejection fraction ≤ 30%
- Impaired renal function (creatinine >2.0 mg/dl or 150mmol/l)
- Platelet count <100,000 cells/mm3 or >700,000 cells/mm3, a WBC of <3,000 cells/mm3, or documented or suspected liver disease (including laboratory evidence of hepatitis)
- Presence of a heart transplant
- Known allergy to cobalt chromium
- Hypersensitivity or contraindication to cobalt-chromium or structurally-related compounds, cobalt, chromium, nickel, or tungsten
- Anticipated use of rotational atherectomy
- Patients in whom the use of a drug eluting stent is contraindicated, e.g., who cannot receive the recommended dual anti-platelet (aspirin and an approved P2Y12 Inhibitor) and/or anticoagulation therapy

Warnings

- Use of the TRYTON Side Branch Stent in appropriately sized main vessels and side branches is required for safe and effective performance of the device.
- Do not use the TRYTON Stent in small side branches [<2.50 mm in diameter by visual assessment or <2.25 mm in diameter by quantitative coronary angiography (QCA)], as its use may lead to an increased risk of adverse cardiac events such as myocardial infarction and the need for repeat revascularization. To confirm appropriately-sized side branch diameters, the diameter of the pre-dilation balloon inflated to nominal pressure may be used as a reference. Alternatively, the use of quantitative imaging methods such as on-line quantitative coronary angiography, intravascular ultrasound or optimal coherence tomography should be considered.

Use of the TRYTON Side Branch Stent, as with percutaneous coronary stent implantation procedures in general, is known to be associated with the following risks:

- Vessel thrombosis
- Increased length of hospital stay relative to those of coronary balloon angioplasty alone. Judicious selection of patients to receive this device rather than balloon angioplasty alone is strongly advised.
- Infection secondary to contamination of the stent may lead to thrombosis, pseudoaneurysm or rupture.
- The stent may cause spasm, distal embolization, thrombus, or could migrate from the site of implantation. Excessive dilatation of the artery may cause vessel rupture and life-threatening bleeding.
- Stents may not be fully expanded during deployment, particularly in resistant lesions.
- Stent dislodgment from the balloon surface during deployment and/or dislodgment from the target site post-deployment can occur.
- Major bleeding.

Precautions

- Side branch pre-dilatation is required and should only be performed with an angioplasty balloon appropriate for a vessel ≥2.5 mm in diameter by visual assessment or ≥2.25 mm in diameter by QCA, inflated to nominal pressure.
- Following pre-dilation, angiography should be performed following the administration of intracoronary nitroglycerin to reassess vessel dimensions with attention to the side branch reference vessel diameter (RVD) to ensure that it is of appropriate size. The side branch RVD should be based on the most angiographic normal-appearing segment distal to the lesion.
- Use of this product should be performed only in hospitals with access to emergency coronary artery bypass graft surgery that can be performed quickly in the event of a potentially injurious or life-threatening complication.
- All TRYTON Side Branch Stent/Stent Delivery Systems are intended for single use only. Under no circumstances should this device or any part thereof be resterilized or reused. Reuse may result in device malfunction and subsequent patient complications and/or adverse events.
- All equipment required for the implantation of this stent must be carefully examined prior to use to verify proper function.
- Special care should be taken not to disrupt the stent on the delivery catheter, particularly during removal from its packaging, placement over guidewire, and advancement through hemostasis valve and guiding catheter.
- When the delivery catheter is exposed to the vascular system, it should be manipulated while under high-quality fluoroscopic observation. If resistance is met during manipulation, determine the cause of the resistance before proceeding. Excessive manipulation may cause dislodgment of the stent from the delivery catheter or vessel damage.
- For deployment of the stent, use a mixture of radiographic contrast media and sterile saline. Do not inflate the delivery system with air or any gaseous media.
- Balloon pressure should not exceed the rated burst pressure of the delivery catheter. Use of a pressure monitoring device is required to prevent over-pressurization.
- Do not attempt to reposition a partially deployed stent. Attempted repositioning may result in severe vessel damage.
- When recrossing a recently implanted stent, care should be taken to assure the guide wire is placed within the lumen and not in between the stent and the vessel wall. Otherwise, inadvertent dislodgment of the stent may occur leading to faulty positioning of the stent.
- Do not attempt to pull an unexpanded stent back into the guiding catheter, as stent damage or stent dislodgement may occur. Movement in and out through the distal end of the guiding catheter should not be performed as the stent may be damaged when retracting the undeployed stent back into the guiding catheter. To withdraw the TRYTON Side Branch Stent system, the entire system with the guiding catheter should be removed as a single unit.
- If a guide catheter extension is utilized to deliver/position the TRYTON Stent and it becomes necessary to withdraw/remove an unexpanded TRYTON Stent/Stent Delivery System, do not withdraw the TRYTON Stent/Stent Delivery System into the guide catheter extension. Withdrawal of the TRYTON Stent/Stent Delivery System into a guide catheter extension may cause dislodgement of the TRYTON Stent from the Stent Delivery System. Refer to procedure step #5 under Use of TRYTON Side Branch Stent/Stent Delivery System.
- Main branch artery preparation including pre-dilatation, stent positioning and deployment should be completed following main branch stent instructions for use.
- Stent retrieval methods (use of additional wires, snares, and/or forceps) may result in additional trauma to the coronary vasculature and/or the vascular access site. Complications may include bleeding, hematoma or pseudoaneurysm.
- The TRYTON Side Branch Stent has not been evaluated in pediatric cases or cases of in-stent restenosis or previously stented lesions.

Pease refer to the Instructions for Use for complete information.

VISTA BRITE TIP® Guiding Catheter

Indications
The guiding catheter is intended for use for intravascular introduction of interventional/diagnostic devices into the coronary or peripheral vascular systems.

Contraindications
None known for guiding catheters.

Warnings
Risk of reuse: This product is designed and intended for single use. It is not designed to undergo reprocessing and resterilization after initial use. Reuse of this product, including after reprocessing and/or re-sterilization, may cause a loss of structural integrity which could lead to a failure of the device to perform as intended and may lead to a loss of critical labeling/use information all of which present a potential risk to patient safety.

Do not use with Ethiodol™ or Lipiodol™ contrast media, or other such contrast media which incorporates the components of these agents.

Precautions
- Store in a cool, dark, dry place.
- Do not use open or damaged packages.
- Use prior to the "Use By" date.
- Do not resterilize.
- Do not expose to organic solvents.
- Inspect the guiding catheter before use to verify that its size, shape and condition are suitable for the specific procedure.
- If strong resistance is met during manipulation, discontinue the procedure and determine the cause of the resistance before proceeding. If the cause of the resistance can not be determined, withdraw the catheter.
- Torquing the guiding catheter excessively while kinked may cause damage which could result in possible separation along the catheter shaft. Should the guiding catheter shaft become severely kinked, withdraw the entire system (guiding catheter, guidewire and catheter sheath introducer).
- Advancement, manipulation and withdrawal of the guiding catheter should always be performed under fluoroscopic guidance.
- Extreme care must be taken to avoid damage to the vasculature through which the guiding catheter passes. The guiding catheter may occlude smaller vessels. Care must be taken to avoid complete blood flow blockage.
- Large internal lumen guiding catheters require less force on the syringe during injection.

Complications
Procedures requiring percutaneous catheter introduction should not be attempted by physicians unfamiliar with the possible complications. Complications may occur at anytime during or after the procedure.

Possible complications include, but are not limited to the following:
- air embolism
- hematoma at the puncture site
- infection
- perforation of the heart
- vessel damage, dissection or perforation
- vasospasm
- ischemia
- hemorrhage
- arrhythmia
- reaction to contrast media
- death

Please refer to the Instructions for Use for complete information, including Adverse Events.

U.S. Headquarters

1820 McCarthy Blvd.
Milpitas CA, 95035
Phone: 408.273.3700

Customer Service Center

Phone: 1.800.327.7714 (toll-free, continental U.S.)
Fax: 800.997.1122

Attn: Customer Contact Center
14201 Northwest 60th Ave
Miami Lakes, FL 33014

Technical Information

1.800.327.7714 (toll-free, continental US)

All orders should be directed to the Customer Service Center by fax, EDI, or telephone. The Customer Service Center is open weekdays (excluding US national holidays) between the hours of 7:30 AM and 9:00 PM EST/EDT. Emergency order service is also available 24 hours a day.

Terms

Net 30 days. Normal delivery second day A.R.O., F.O.B. shipping point with freight prepaid. Standing order shipments are F.O.B. shipping point with freight prepaid. Prices are subject to change without notice.

Disclaimer of warranty and limitation of remedy

There is no express or implied warranty, including any implied warranty of merchantability of fitness for a particular purpose, including the products described in this publication. Under no circumstances shall Cordis be liable for any direct, incidental or consequential damages other than as expressly provided by specific law. No person has the authority to bind Cordis to any representation or warranty except as specifically set forth herein.

Standing Purchase Orders

Cordis encourages standing purchase orders. Standing purchase orders are agreements to purchase specific quantities of product for delivery on a specific, predetermined schedule. To establish a standing purchase order, telephone the Customer Contact Center at 1.800.327.7714 option 1, or contact your local Cordis Sales Representative.

Intended Product Usage/Storage

- **Usage:** Most products in this catalog are supplied sterile and nonpyrogenic. Do not use any products if their sterile package is damaged. Discard catheters and single-use accessories after one procedure. All parts are extremely difficult to clean. DO NOT REUSE OR RESHAPE. DO NOT AUTOCLAVE SINGLE-USE PRODUCTS. Structural integrity and/or function may be impaired through reuse, reshaping, and cleaning. ACCORDINGLY, CORDIS CORPORATION WILL NOT BE RESPONSIBLE FOR ANY DIRECT, INCIDENTAL OR CONSEQUENTIAL DAMAGES RESULTING FROM REUSE OF CATHETERS AND ACCESSORIES. Prior to use, refer to the instructions accompanying the product.
- **Storage:** Store products in a cool, dark, dry place. Use sterile products prior to the "Use By" date. Do not expose to organic solvents.
- **Caution:** Federal (USA) law restricts these devices to sale by or on the order of a physician.

Return Policy

Products on consignment are not returnable for credit under this policy. To return any product on consignment, you must contact your local Sales Representative for instructions.

Return Authorization

To return other Cordis products you must contact the Customer Service Center at 1.800.327.7714 option 2 or your local sales representative for a return goods authorization. You must also provide information about the acquisition method (e.g. purchase, consignment, or evaluation) for the product being returned.

Credits

Full Credit of the invoice price will be issued on products returned within 180 days from the date of invoice. Full Credit of the invoice price, less a 10% restocking charge, will be issued on products returned later than 180 days from the date of invoice with the exception of EXOSEAL® Vascular Closure Device.

Partial credit will be issued for product returned in quantities less than the full 10 pack for EXOSEAL® Vascular Closure Device when returned within 365 days.

Credit will not be issued for products returned greater than 365 days from the date of invoice. Additionally, credit will not be issued for product meeting the following conditions:

- Discontinued from sale
- Modified products ("Specials")
- Packaging is opened
- Modified product/trays
- Modified instruments
- Damaged
- Used
- Less than full shipping unit
- Stickers, markings, or any changes made to blister package or packaging in general

Return Logistics

All products should be returned with freight prepaid by the customer within five (5) business days of receiving Return Authorization approval. All returned products must be accompanied with the Return Authorization Confirmation in its proper protective packaging along with the Return Authorization Number written on the packaging and sent to:

Cordis Returned Goods
8640 Nail Road, Suite 115
Olive Branch, MS 38654

All returns due to Cordis should use the merchandise pick-up process. Contact Customer Service at 1.800.327.7714 option 1, and they will have our Transportation Department arrange for the carrier to pick up the order error for return.

Product Complaints

This policy does not address product complaints. All product complaints should be handled by calling Customer Service at 1.800.327.7714.

Trademarks

The trademarks listed below are trademarks of Cardinal Health and may be registered in the US and/or in other countries. All other marks are the property of their respective owners.

- CORDIS
- The Cordis LOGO
- ADROIT® Guiding Catheter
- ATW™ Steerable Guidewire
- ATW™ Marker Wire
- AVANTI®+ Sheath Introducer
- BI-PAL® Biopsy Forceps
- EMERALD® Diagnostic Guidewire
- EMPIRA® Balloon Catheter
- EXOSEAL® Vascular Closure Device
- 7F HIGHFLOW™ Diagnostic Catheter
- INFINITI® Diagnostic Catheter
- MYNX ACE® Vascular Closure Device
- MYNXGRIP® Vascular Closure Device
- REFLEX® Steerable Guidewire
- SHINOBI® Plus Steerable Guidewire
- SHINOBI® Steerable Guidewire
- STABILIZER® Plus Steerable Guidewire
- STABILIZER® XS Steerable Guidewire
- SUPER TORQUE® Diagnostic Catheter
- VISTA BRITE TIP® Guiding Catheter
- WIZDOM™ Steerable Guidewire

For customer service call 1.800.327.7714. To learn more visit cordis.com.
CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.
Important information: Prior to use, refer to the instructions for use supplied with this device for indications, contraindications, side effects, suggested procedure, warnings and precautions.

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