- DCB Bench testing performed by Boston Scientific Corporation. Data on file at Boston Scientific. Tip profile data on file for 3.00 mm DCBs. Flexibility comparison data on file for 3.50 mm DCBs excluding Selution, data on file for 3.00 mm DCB.
- Pre-clinical pharmacokinetic studies, data on file at Boston Scientific for AGENT, SeQuent Please and Selution. Magic Touch data derived from product brochure (TCT 2020).
- 3. Blagosklonny et al. Paclitaxel induces primary and postmitotic G1 arrest in human arterial smooth muscle cells. Cell Cycle. 2004 Aug;3(8):1050-6. Epub 2004 Aug 24.
- Shahrori ZMF, et al. Drug-Coated Balloons: Recent Evidence and Upcoming Novelties. J Cardiovasc Dev Dis. 2025 May 20;12(5):194. doi: 10.3390/jcdd12050194. PMID: 40422965; PMCID: PMC12112549. Other coronary PTX DCB drug concentrations; SeQuent Please 3 μg/mm², Prevail 3.5 μg/mm², Pantera Lux 3 μg/mm². Also see respective product IFU's.
- 5. Market data on file at BSC as of July 2023.
- Hamm, C, et all, AGENT ISR study, Eurointervention, 2020, 3-year comparative clinical study results on file at Boston Scientific Corporation: Data available upon request.
- 7. Nakamura, M, et all, Drug-coated balloon for the treatment of small vessel coronary artery disease, Circulation Japan 2022.
- 8. Yeh RW, Shlofmitz R, Moses J, et al. Paclitaxel-Coated Balloon vs Uncoated Balloon for Coronary In-Stent Restenosis: The AGENT IDE Randomized Clinical Trial, JAMA. 2024;33(12):1015–1024. doi:10.1001/jama.2024.1361

Please review the full IFU for indications for use and operating instructions.

CAUTION: The law restricts these devices to sale by or on the order of a physician. Indications, contraindications, warnings, and instructions for use can be found in the product labelling supplied with each device or at www.IFU-BSCI.com. Products shown for INFORMATION purposes only and may not be approved or for sale in certain countries. This material not intended for use in France.



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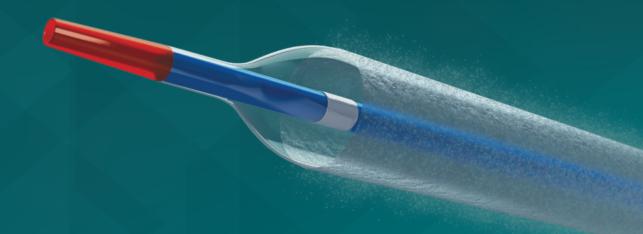
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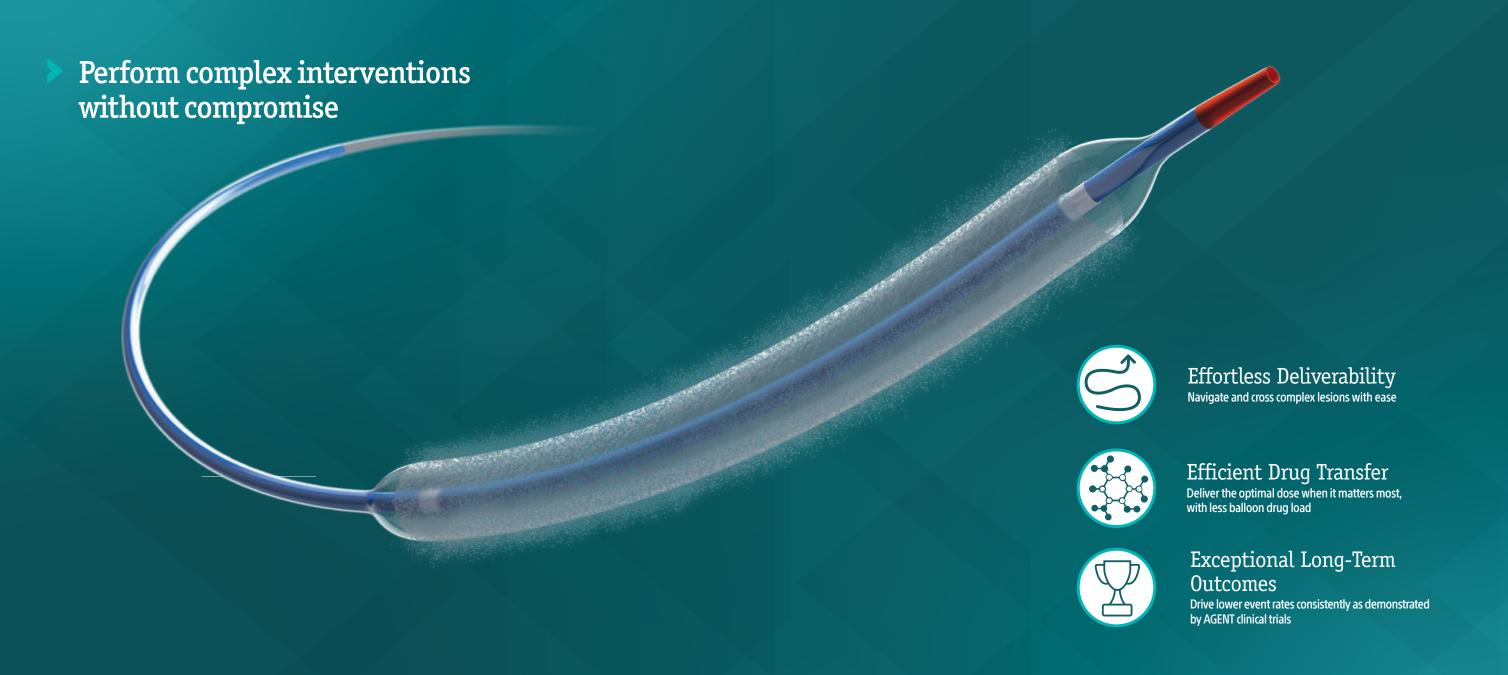


AGENT ™ Drug-Coated Balloon





Leave the Right **AGENT** Behind



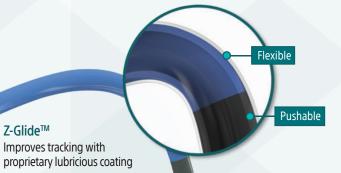
Effortless DeliverabilityNavigate and cross complex lesions with ease

> Z-Glide™



▶ Laser Bonded Tip Improves crossability and reduces tip catch

➤ Bi-segment Inner Shaft Maximizes flexibility without compromising push for reduced resistance in tortuous anatomy



The AGENT Difference Ultra-Low Tip Profile¹

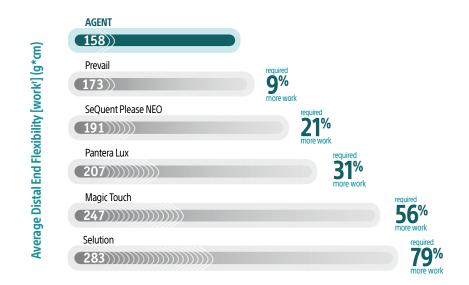






LOWER IS BETTER

Uncompromised Flexibility¹



Distal End Flexibility

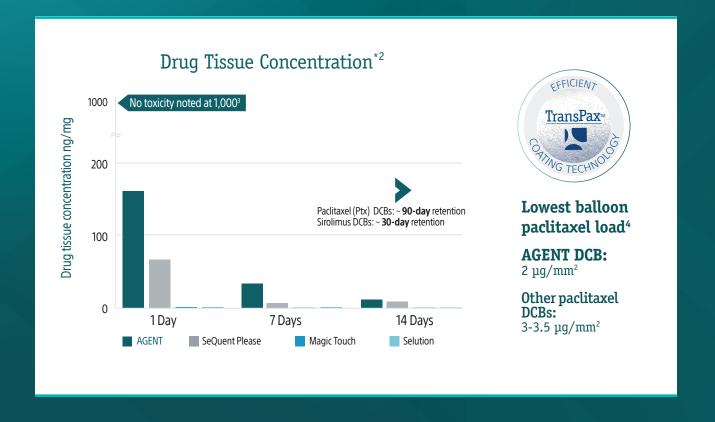
Overcoming the rigidity typically introduced by drugcoating matrices (drug + excipient/carrier) is key to optimal distal end flexibility. AGENT DCB demonstrated bestin-class distal end flexibility, minimizing resistance and enhancing deliverability in tortuous anatomy.



† Measure of how much work is required to deliver the distal end of the device through a curved simulated artery – how easily does the distal assembly flex.

Efficient Drug Transfer

Deliver the optimal dose when it matters most, with less balloon drug load. Due to the efficiency of AGENTTM Drug-Coated Balloon's proprietary TransPaxTM Coating Technology, AGENT is able to use the lowest paclitaxel dosage compared to other paclitaxel-coated balloons, while still resulting in the highest concentration of drug in the tissue.



➤ TransPax[™] coating optimizes the three phases of drug delivery

AGENT DCB is designed with a novel excipient, sharp-edge structure, and uniform crystalline formulation. As a result, AGENT transfers more drug to the tissue and less of it downstream.

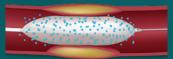
▶ Design Requirement

Targeted Transfer
Balloon to vessel wall



• •





Sustained Retention
Prolonged tissue concentration



➤ Paclitaxel (PTX) Drug Choice

Hydrophobic
Paclitaxel is durable under hydration



Lipophilic

Paclitaxel has a high affinity for fatty tissue



Chemically Stable

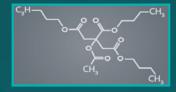
Paclitaxel has a long half-life



▶ AGENT Coating Design

Optimized Excipient

Acetyl tributyl citrate (ATBC) maximizes balloon-to-vessel wall transfer



Sharp-Edge Structure

Needle-like coating improves tissue penetration



Crystalline Formulation

Crystallized coating maintains therapeutic effect



Exceptional Long-Term Outcomes

Drive lower event rates consistently as demonstrated by AGENT clinical trials. **Over 4,400 patients** have been evaluated or are undergoing evaluation with AGENT DCB.⁵ This includes studies in ISR, de novo lesions, small vessels, large vessels, bifurcations, complex lesions, and patient-specific cases.

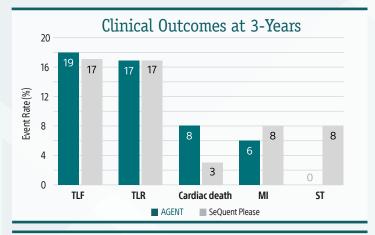
AGENT ISR Study⁶

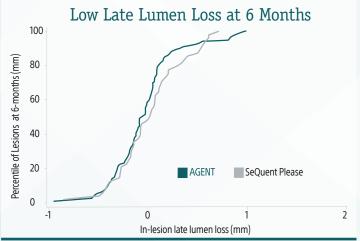
AGENT demonstrated comparable event rates at 3-years compared to SeQuent Please

AGENT Japan SV Study⁷

AGENT demonstrated numerically lower levels of late lumen loss compared to SeQuent Please







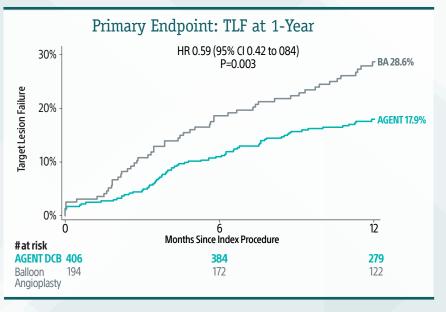
Proven clinical performance

AGENT US IDE Study⁸

This prospective, multicenter, 2:1 randomized controlled trial showed statistically lower event rates for AGENT DCB at one year.







At one-year, AGENT DCB also demonstrated statistically lower event rates:



(13.0% vs. 24.7%, P=0.0005)



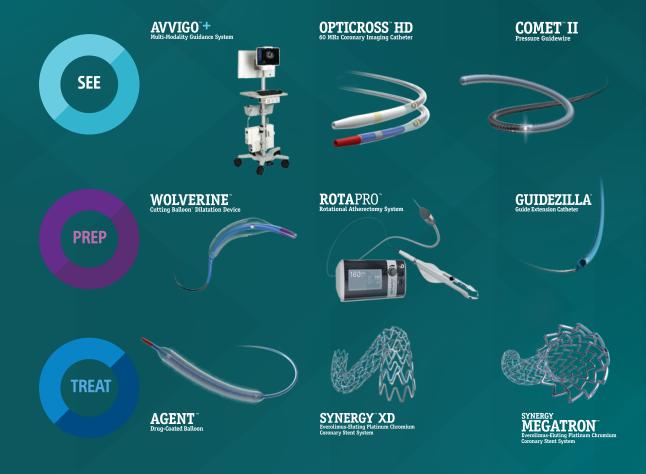


(0.0% vs. 3.2%, P=0.0004)

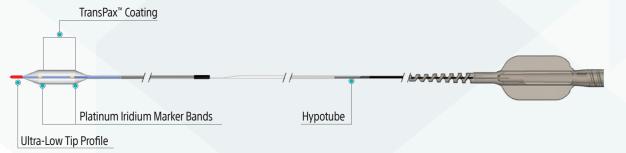
*Number Needed to Treat to prevent 1 TLF

Modern PCI Portfolio

With the most comprehensive Modern PCI portfolio on the market, Boston Scientific has the tools needed to See, Prep, and Treat each lesion.







One platform. Expanded potential. AGENT DCB's 40 mm balloon is here.



Drug-Coated Balloon Length (mm)

Balloon Length Balloon Diameter	12mm	15mm	20mm	30mm	40mm
2.00	H74939222201210	H74939222201510	H74939222202010	H74939222203010	H74939222204010
2.25	H74939222221210	H74939222221510	H7493922222010	H7493922223010	-
2.50	H74939222251210	H74939222251510	H74939222252010	H74939222253010	H74939222254010
2.75	H74939222271210	H74939222271510	H74939222272010	H74939222273010	-
3.00	H74939222301210	H74939222301510	H74939222302010	H74939222303010	H74939222304010
3.50	H74939222351210	H74939222351510	H74939222352010	H74939222353010	H74939222354010
4.00	H74939222401210	H74939222401510	H74939222402010	H74939222403010	H74939222404010