Calcified lesions



Prevalence of calcified coronary lesions





Moderate-severe calcification in 13 DES studies

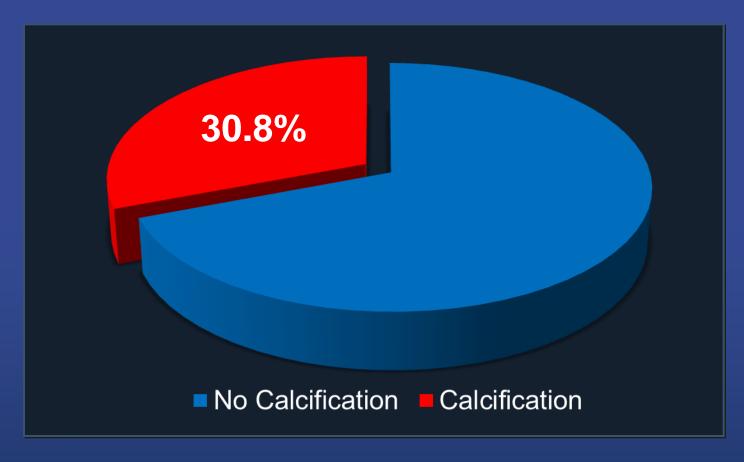
| RAVEL | 23.3% (27/116) |
|--------------|---------------------|
| SIRIUS | 17.1% (91/531) |
| E-SIRIUS | 16.1% (28/174) |
| C-SIRIUS | 12.0% (6/50) |
| TAXUS IV | 18.3% (121/660) |
| TAXUS V | 32.5% (185/570) |
| TAXUS VI | 29.7% (65/219) |
| ENDEAVOR II | 23.7% (140/590) |
| ENDEAVOR III | 17.9% (78/436) |
| ENDEAVOR IV | 33.2% (513/1546) |
| SPIRIT II | 31.4% (91/290) |
| SPIRIT III | 27.8% (277/997) |
| COMPARE | 38.5% (693/1799) |
| Total | 29.0% (2,315/7,978) |





ADAPT-DES (11 center all-comers registry): Mod-Sev Calcification

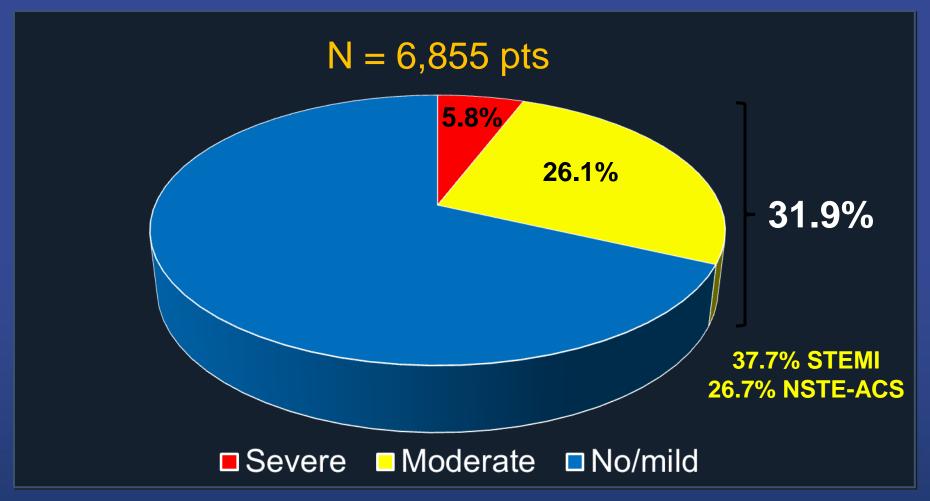
N = 8,582 pts







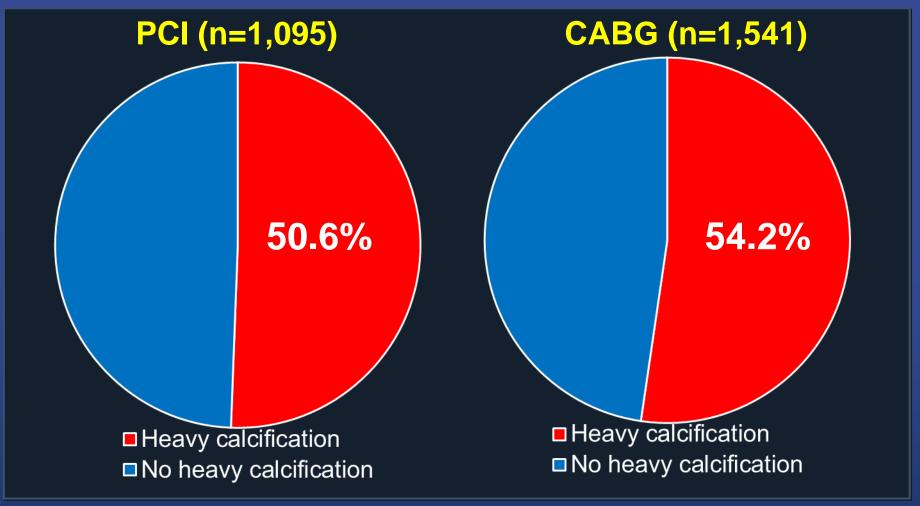
Frequency of Mod-Sev Calcification in NSTE-ACS and STEMI PCI population: (ACUITY and HORIZONS-AMI)





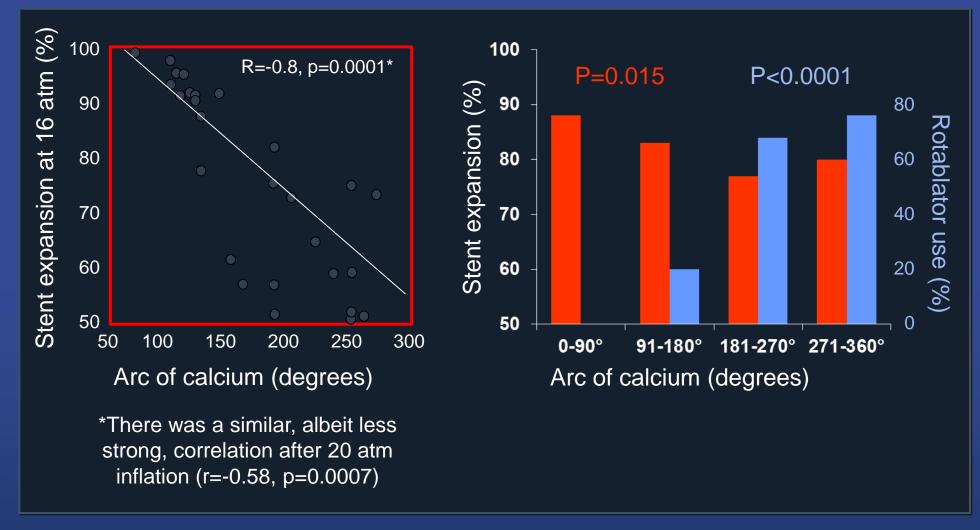


Frequency of "heavy" calcification in the SYNTAX trial: Randomized + Registry N=2,636 pts with LM or 3VD



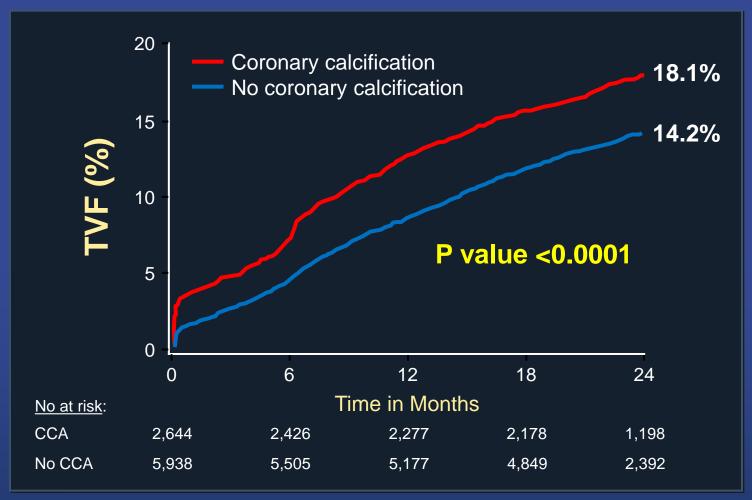


Stent Expansion in Calcified Lesions





ADAPT-DES (N=8,582) Target vessel failure at 2 years





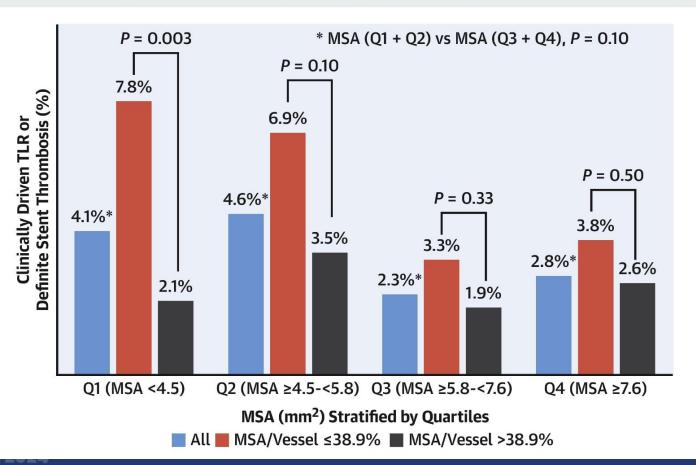


ADAPT-DES (N=8,582): Calcification and 2-year Events

| | Calcification | | Unadjusted | Adjusted | Adjusted |
|---------------------------|-----------------|------------------|------------|-------------------|----------|
| | No (n=5,938) | Yes (n=2,644) | p | HR [95% CI] | p |
| TVF | 14.2% | 18.1% | <0.0001 | 1.23 [1.09, 1.39] | 0.0008 |
| MACE | 5.6% | 8.3% | <0.0001 | 1.47 [1.22, 1.76] | <0.0001 |
| Death | 3.5% | 4.8% | 0.003 | 1.15 [0.90, 1.46] | 0.26 |
| CV death | 2.3% | 2.8% | 0.09 | 1.09 [0.80, 1.48] | 0.60 |
| MI | 4.0% | 6.4% | <0.0001 | 1.61 [1.30, 1.99] | <0.0001 |
| Clinically- driven TVR | 9.5% | 10.4% | 0.16 | 1.10 [0.94, 1.29] | 0.24 |
| Stent thrombosis | 0.9% | 1.1% | 0.32 | 1.49 [0.92, 2.43] | 0.11 |

Stent Expansion Indexes to Predict Clinical Outcomes: An IVUS Substudy From ADAPT-DES

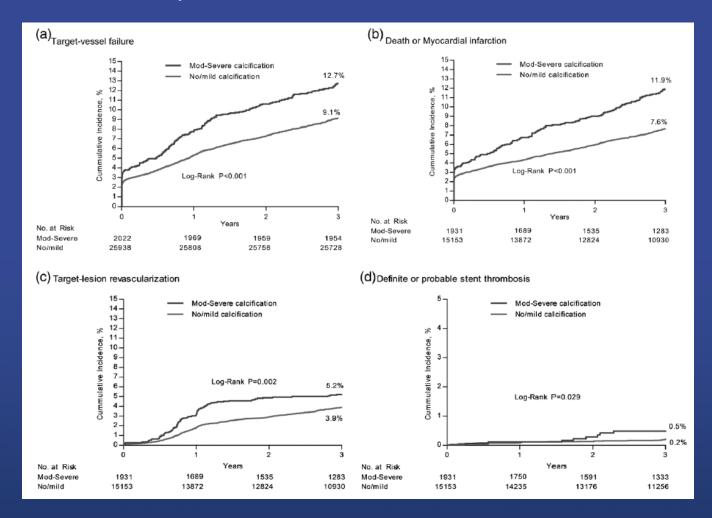
CENTRAL ILLUSTRATION: 2-Year Rate of Clinically Driven Target Lesion Revascularization or Definite Stent Thrombosis Stratified by Minimum Stent Area Quartiles and Minimum Stent Area/Vessel (≤38.9% Versus >38.9%)



Stent/vessel area at the MSA site, an index of relative stent expansion, was superior to absolute MSA and other expansion indexes in predicting 2-year clinically driven TLR or definite stent thrombosis

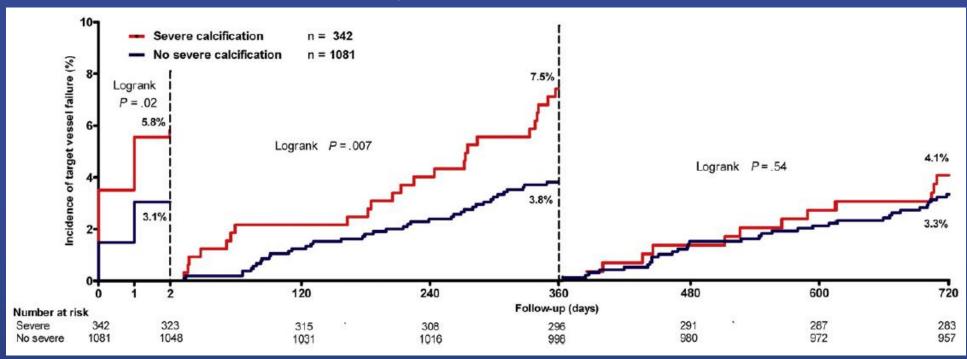
Data from IRIS-DES Registries

17,084 patients who underwent PCI with DES



TWENTE and DUTCH PEERS (TWENTE II): Impact of Severe Calcification with 2nd Generation DES

1,423 pts with stable angina; 342 (24%) with severe calcification



At 2 years, TVF was 16.4% vs. 9.8%, p=0.001 predominantly driven by events in the first 48 hours and up to 1 year



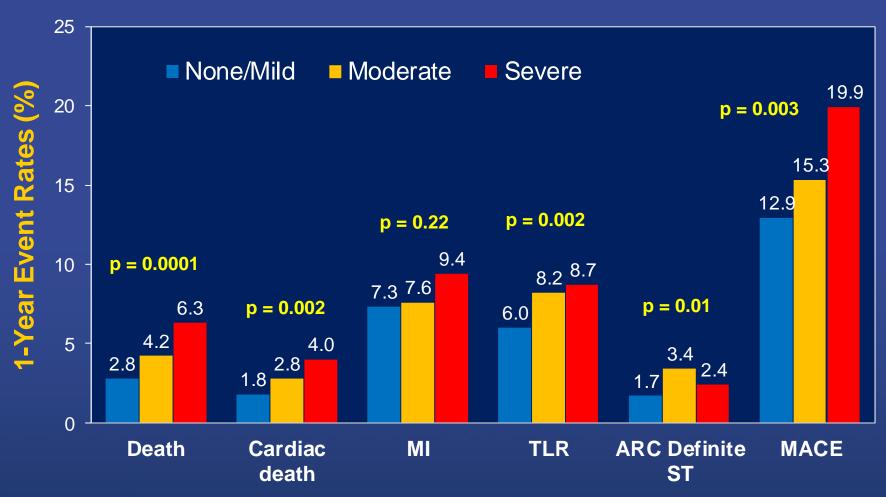
ACUITY/HORIZONS-AMI: Implications of Calcified Lesions on PCI in ACS

| Post-PCI | Moderate/Severe (n=2,958) | None/Mild (n=5,783) | P value |
|----------------|---------------------------|------------------------|---------|
| TIMI flow 0/1 | 2.6% | 1.6% | 0.001 |
| TIMI flow 2 | 6.8% | 5.2% | 0.004 |
| TIMI flow 3 | 90.6% | 93.1% | <0.0001 |
| No reflow | 0.4% | 0.1% | 0.02 |
| Perforation | 0.1% | 0.1% | 0.41 |
| Spasm | 1.1% | 0.6% | 0.02 |
| Dissection | 2.9% | 1.2% | <0.0001 |
| Abrupt closure | 0.5% | 0.1% | 0.001 |





ACUITY/HORIZONS-AMI: Implications of Calcified Lesions on PCI in ACS







ACUITY/HORIZONS-AMI: Implications of Calcified Lesions on PCI in ACS

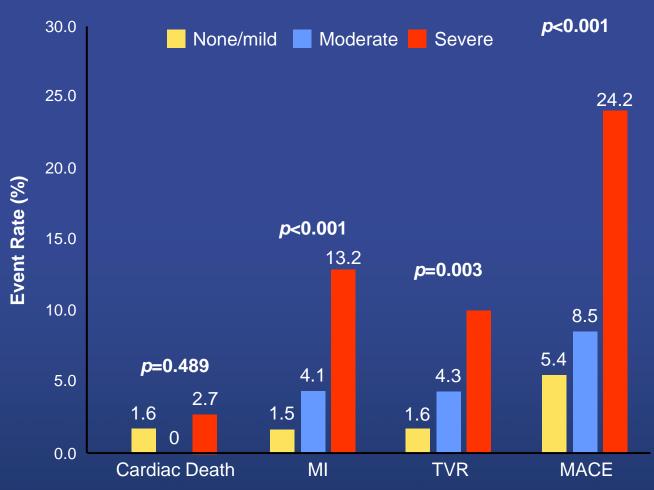
| | Adjusted Hazard Ratio [95% CI] | P Value |
|-----------------|--------------------------------|---------|
| Death | 1.10 [0.81,1.48] | 0.55 |
| MI | 1.06 [0.86,1.30] | 0.58 |
| Ischemic TLR | 1.44 [1.17,1.78] | 0.0007 |
| ARC definite ST | 1.62 [1.14,2.30] | 0.007 |





Impact of calcification on percutaneous coronary intervention:

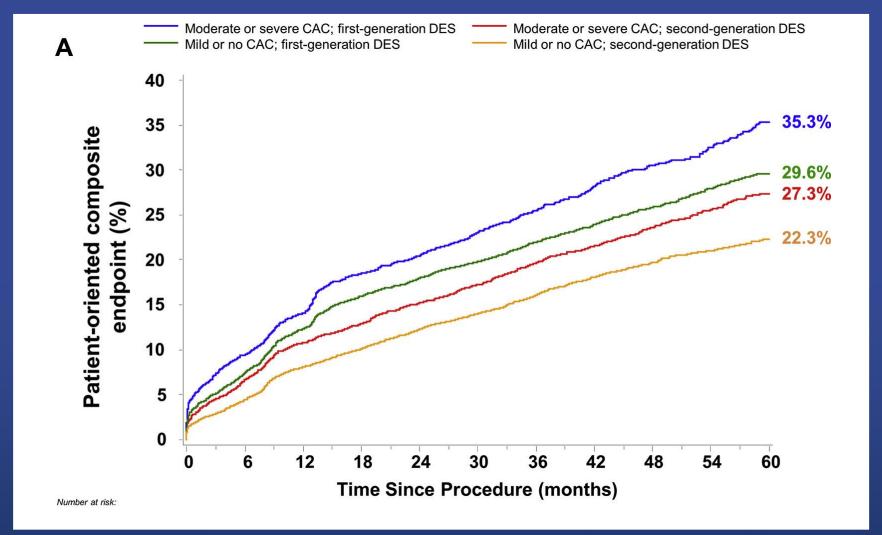
MACE-Trial 1-year results







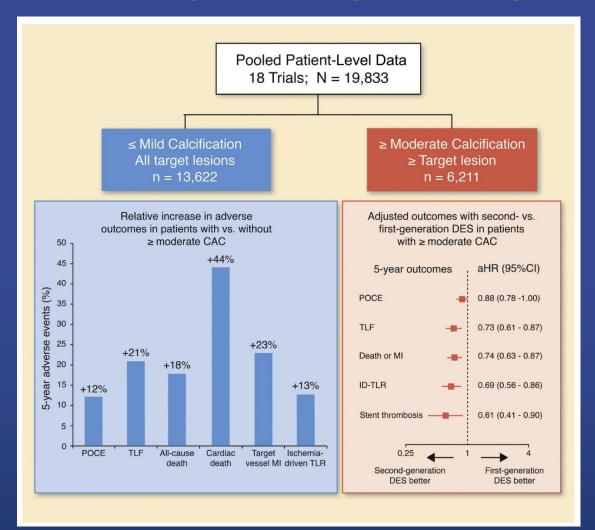
Coronary Calcification and Long-Term Outcomes According to Drug-Eluting Stent Generation







Coronary Calcification and Long-Term Outcomes According to Drug-Eluting Stent Generation



PCI of target lesion moderate or severe CAC was associated with adverse patient-oriented and deviceoriented adverse outcome at 5 years





Treatment of Calcified Lesions

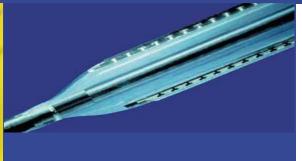
NC balloons

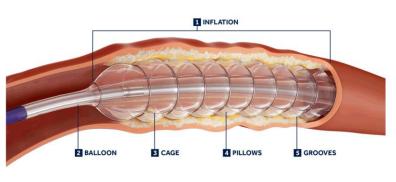
Cutting balloon

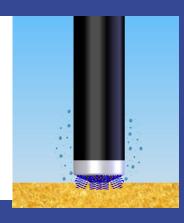
Chocolate PTCA balloon

Laser









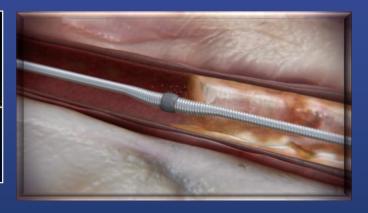
Angiosculpt



Rotational atherectomy



Orbital atherectomy





New Technics

Intravascular lithotripsy



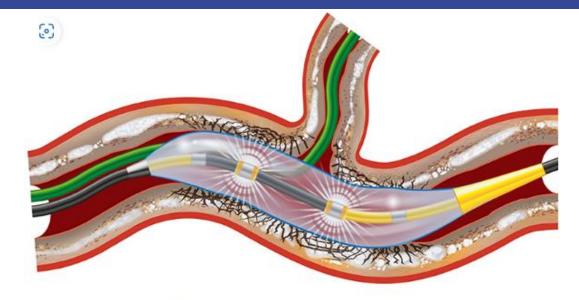
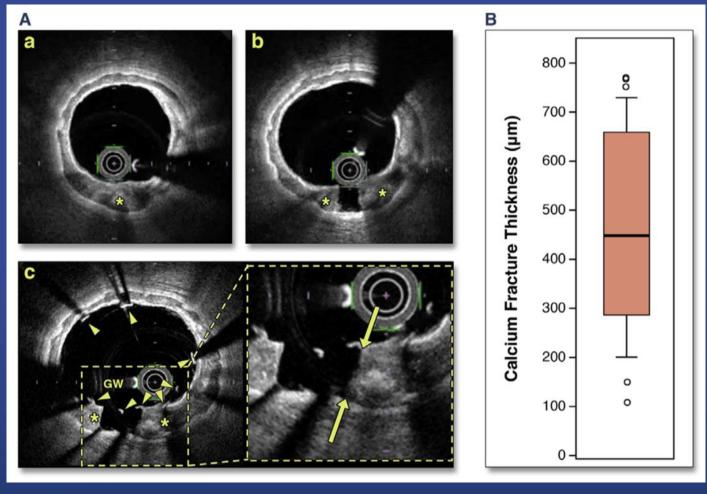


Photo Credit: Shockwave Medical

Calcium Fracture and Relation to Outcomes

61 pts with heavily calcified lesions studied serially with OCT Fracture was seen in 48% (more frequently with CB or atherectomy)



Fracture was associated with greater MSA and less restenosis/ID-TLR



Optical frequency-domain predictor good stent expansion after atherectomy

50 de novo heavily calcified lesions that underwent OFDI-guided RA)

| Variable | Univariate predictors | | Multivariate predictors | | |
|------------------------------|------------------------------|----------|------------------------------|--------------|----------|
| | Standardized coefficient (β) | P | Standardized coefficient (β) | t-statistics | P |
| Diabetes mellitus | 0.058 | 0.69 | | | |
| Hemodialysis | -0.073 | 0.61 | | | |
| Burr-to-artery ratio | 0.009 | 0.95 | | | |
| Arc of calcium | 0.075 | 0.60 | | | |
| Minimum thickness of calcium | -0.53 | < 0.001* | -0.45 | -3.78 | < 0.001* |
| Maximum thickness of calcium | 0.50 | 0.50 | | | |
| Length of calcium | -0.10 | 0.90 | | | |
| Dissection formation | 0.43 | 0.002* | 0.32 | 2.65 | 0.011* |

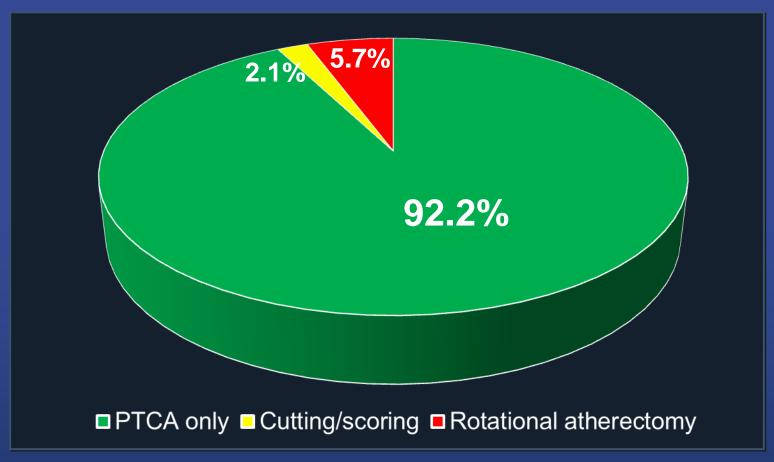
Minimum of thickness of calcification in the intima and dissection formation were positively associated with good stent expansion after RA.





ADAPT-DES (11 center all-comers registry): Calcified lesion preparation

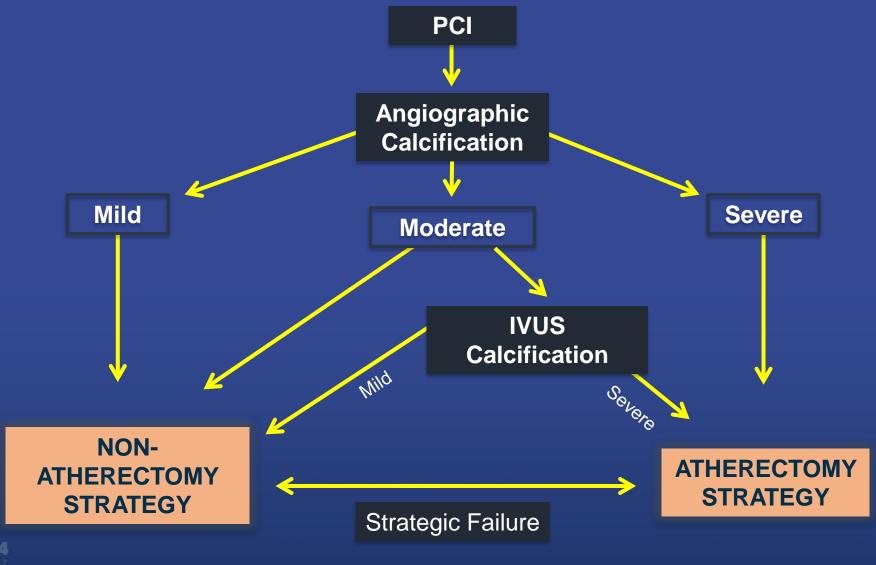
N = 2,644 patients







Potential Strategy for Calcified Lesions

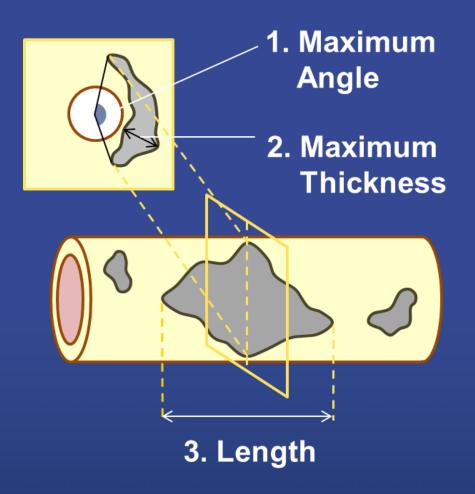




PCI Guideline recommendation

- In patients with fibrotic or heavily calcified lesions, plaque modification with rotational atherectomy can be useful to improve procedural success.
 (class 2a-B)
- In patients with fibrotic or heavily calcified lesions, plaque modification with orbital atherectomy, balloon atherotomy, laser angioplasty, or intracoronary lithotripsy may be considered to improved procedural success.
 (class 2b-B)
- Cutting or scoring balloon angioplasty or rotational atherectomy may be required in selected lesions—particularly those with heavy calcification—in order to adequately dilate lesions prior to stent implantation
- However, studies investigating the systematic use of these adjunctive technologies have failed to show clear clinical benefit.

Calcium Volume Index (CVI) Scoring System



| OCT-based CVI Score | | |
|----------------------------|------------------------------|--|
| Al | ≤ 180° ⇒ 0 point | |
| Angle | > 180° ⇒ 2 points | |
| Thick | ≤ 0.5 mm → 0 point | |
| ness | > 0.5 mm \Rightarrow 1 point | |
| Length | ≤ 5.0 mm → 0 point | |
| Length | > 5.0 mm → 1 point | |
| Total score: 0 to 4 points | | |

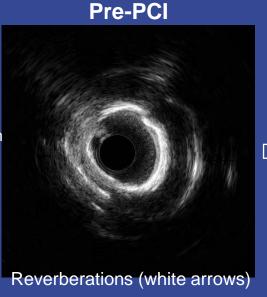




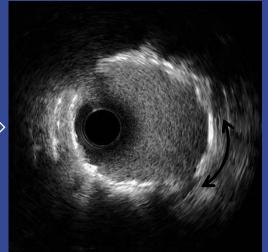
IVUS-Based Calcium Scoring System

Example: Calcium Score=0

- Length of Ca >270° = 4.1mm
- Calcified nodule (-)
- Vessel diameter = 4.4mm
- Reverberation arc >90°



Post-PCI



Excellent expansion despite severe Ca
Stent area= 9.7mm²

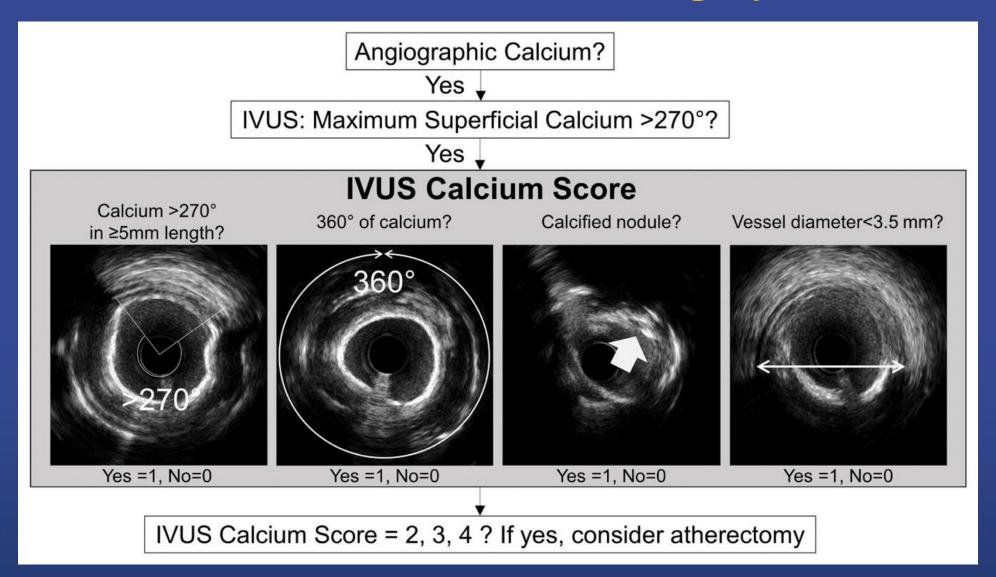
Note Ca fracture (newly visible perivascular tissue, doubleheaded white arrow)

| | Cut-off value | AUC | Score |
|--------------------------|---------------|------|---|
| Length of Calcium > 270° | 5.4 | 0.73 | ≤5mm → 0 point |
| (per 5mm) | 5.4 0.73 | | >5mm → 1 point |
| Vessel diameter | 3.4 | 0.74 | >3.5mm \rightarrow 0 point |
| (per 1mm) | 3.4 | 0.74 | ≤3.5mm → 1 point |
| Calcified nodule | NA | NA | Absent \rightarrow 0 point |
| Calcilled Hoddle | IVA | INA | Present → 1 point |
| Reverberation arc | 97° | 0.81 | $>90^{\circ} \rightarrow 0 \text{ point}$ |
| (per 90°) | 31 | 0.01 | ≤90° → 1 point |





IVUS-Based Calcium Scoring System





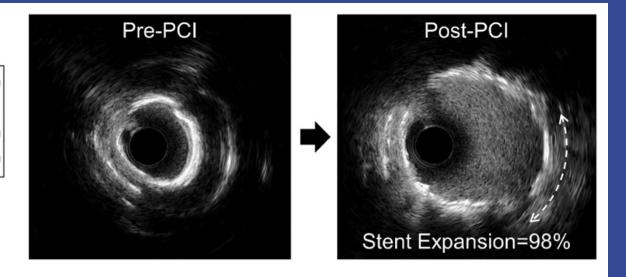


Calcium Scoring System (examples)

Case 1

- Length of Ca >270° = 4.1 mm = 0
- 360° of Calcium (+)
- Calcified nodule (-) = 0
- Vessel diameter = 4.4 mm = 0

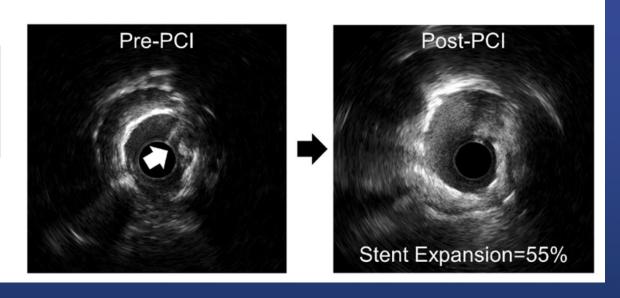




Case 2

- Length of Ca >270° = 8.9 mm = 1
- 360° of Calcium (-) = 0
- Calcified nodule (+) = 1
- Vessel diameter = 2.9 mm = 1









Angiosculpt Balloon

AngioSculpt is a scoring balloon catheter comprised of two main components:

1. Angioplasty balloon catheter

- semi-compliant nylon balloon
- coaxial, nylon shaft
- 2 marker bands

2. Scoring element

- - helical configuration







Scoring Mechanism of Action

AngioSculpt is the only device to offer 3 distinct benefits with one device:

- -Precision
- -Predictable Power
- -Safety







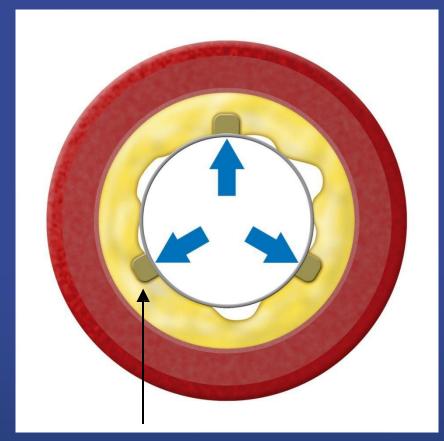
Precision – Minimal Slippage



- Rectangular edges "lock" the device into lesion
- No significant device slippage = less damage to healthy tissue



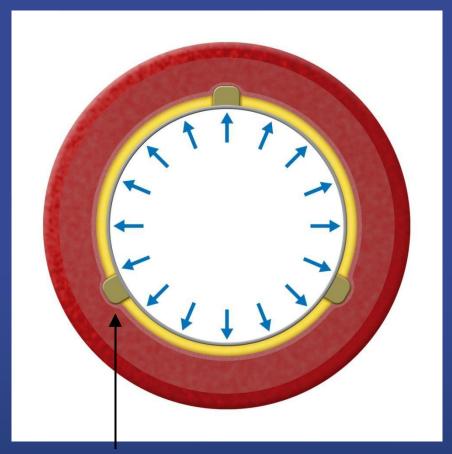
Power – More Dilatation Force



15-25X force of POBA*

- Leading edges drive outward force 15-25 times that of POBA
- Helical arrangement of scoring element creates uniform luminal enlargement

Safety – Low Dissection Rate



- Post-scoring, outward forces are designed to be equivalent to POBA
- Low dissection rate
- Low rate of adjunctive stenting

1X force post scoring*



Features & Benefits - Scoring Element

| Feature | Benefit |
|--------------------------------------|--|
| Nitinol material | Facilitates balloon deflation |
| Helical shape | Uniform, circumferential scoring |
| | Reduces balloon slippage |
| Electropolished | Provides safe scoring – |
| rectangular edges | minimize |
| | dissections |



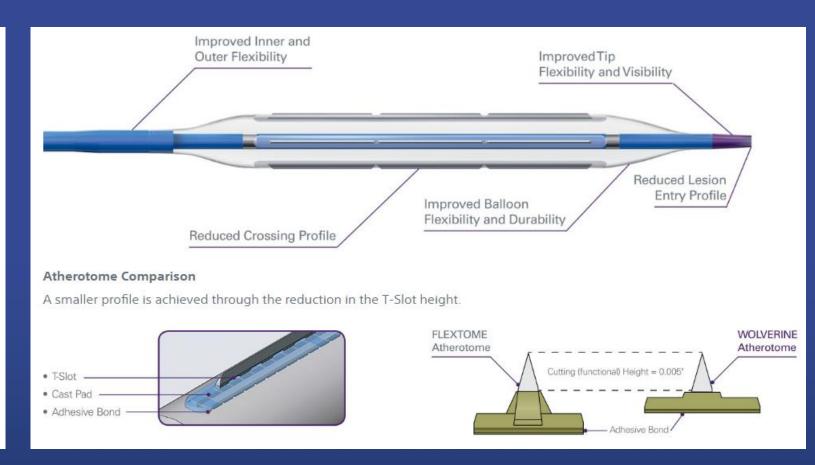
Element strut height .005" or .007"





Cutting Balloon







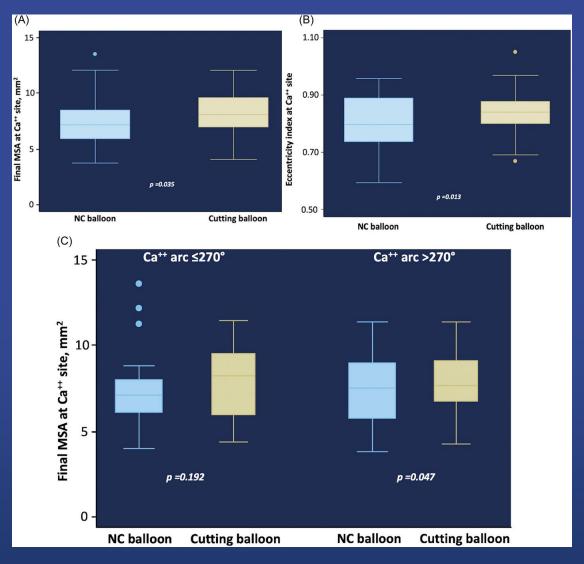


Cutting Balloon to Optimize Predilation for Stent Implantation: The COPS Randomized Trial

- 100 consecutive patients with calcified lesions
- Randomized to cutting balloon vs. non-compliant balloon
- Lesions excluded
 - In-stent restenosis
 - Graft restenosis
 - Thrombotic lesions
- Lesion characteristics
 - RVD 3.4 mm
 - Average calcium length: 12 mm
 - B2/C 71%



Cutting Balloon to Optimize Predilation for Stent Implantation: The COPS Randomized Trial



Israeli Registry -Baseline Characteristics

- 521 consecutive patients scheduled for PCI
- 521 patients and 745 lesions treated
- Lesions excluded
 - Without calcification
 - With untreated visible thrombus
- Lesion characteristics
 - RVD 2.48 mm
 - Average lesion length: 19.2 mm
 - Moderate/severe calcification: 75%
 - B2/C 53%
 - Bifurcations 18%
 - Angulated 43%





Israeli Registry – Results (Acute)

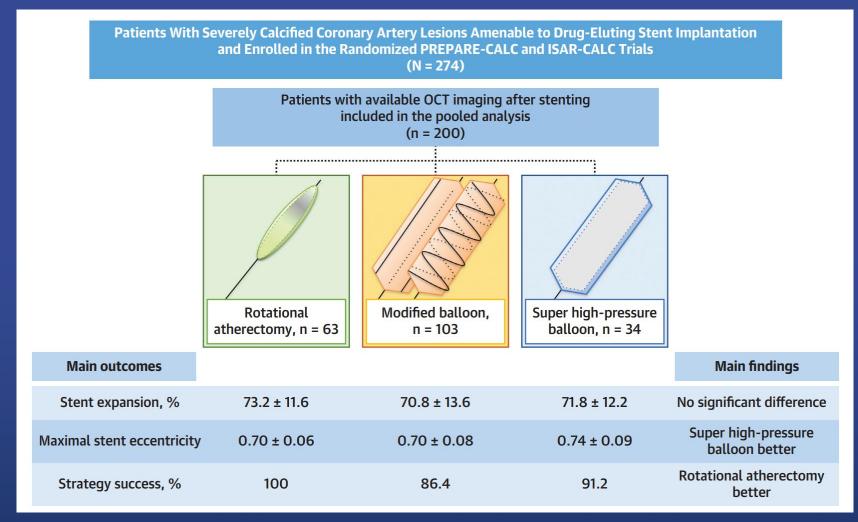
| | Pre-ASC | Post-ASC | Post-Stent |
|----------------|--------------------|---------------------|--------------------|
| MLD (QCA) mm | 0.22 <u>+</u> 0.17 | 2.04 <u>+</u> 0.57 | 2.49 <u>+</u> 0.69 |
| DS% | 84.8 <u>+</u> 13.9 | 21.7 <u>+</u> 12.7 | 5.7 <u>+</u> 2.4 |
| CSA (IVUS) mm2 | 2.49 <u>+</u> 0.39 | 3.72 <u>+</u> 1.12* | 5.30 + 2.05* |

*p<0.001

- Device slippage 1.2% lesions (9/745)
- Significant dissection (≥ type C) post ASC 1.5%
- No device-related perforations

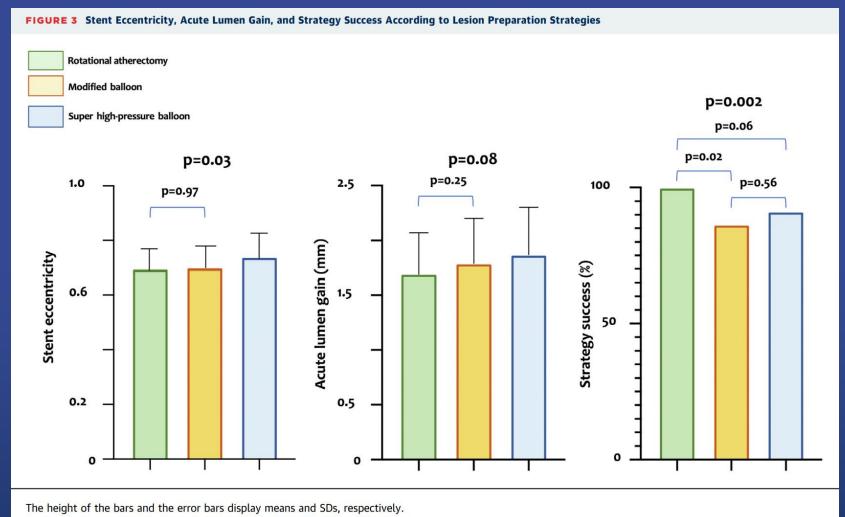


Rotational Atherectomy or Balloon-Based Techniques to Prepare Severely Calcified Coronary Lesions



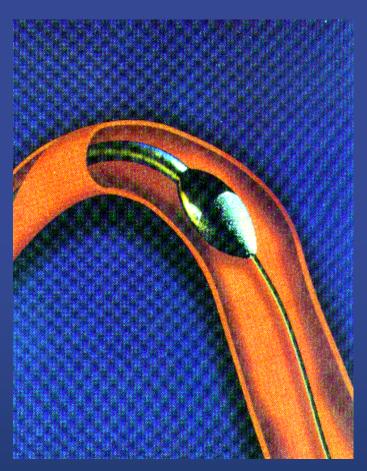


Rotational Atherectomy or Balloon-Based Techniques to Prepare Severely Calcified Coronary Lesions





Rotational Atherectomy (Rotablator)



- Burr : covered with 20-30 um diamond chips
- Guidewire : 0.009 inch with 0.014 inch tip









Rotablator Rotational Atherectomy System







Rotalink and Burr

RotaLink Plus System



RotaLink System Burrs



RotaLink System Advancer



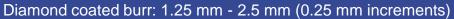
Pre-assembled

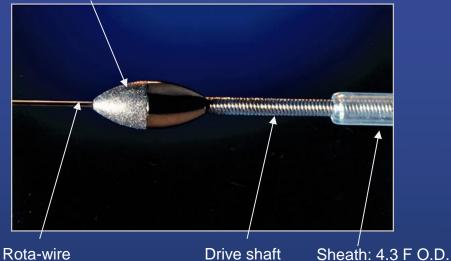
Separated



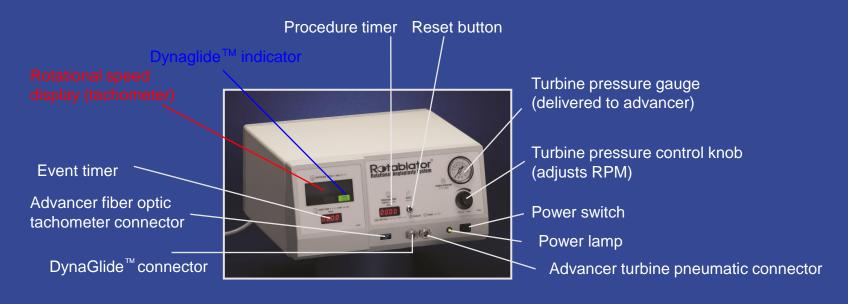


To avoid damage to the burr, remove distal gripper after connection to Advancer

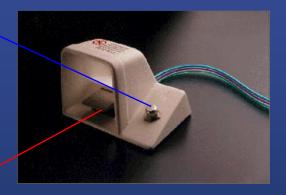




Console, Foot pedal, Gas, and Fluid



DynaGlide[™]
button (low/high
speed selector)



DynaGlide[™] Connectors (blue color)

Compressed Air or Nitrogen connector



Foot switch

Power Cord





Rotawire

RotaWire floppy guidewire



- Tip diameter= 0.014 inch, body diameter= 0.009 inch
- Spring tip length = 22 mm
- 'Long neck' segment: 130 mm, 0.005 0.0077 inch
- Total length 3300 mm

RotaWire extra support guidewire



- Tip diameter= 0.014 inch, body diameter= 0.009 inch
- Spring tip length = 28 mm
- 'Short neck' segment: 50 mm, 0.005 0.0077 inch
- Total length 3330 mm

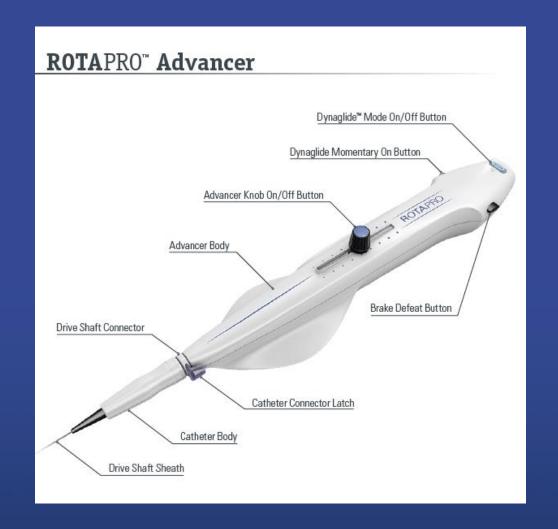
Cf) Rotalink length = 1350 mm





ROTAPRO™ Rotational Atherectomy System









Current Indications of Rotablator

Indication: lesion modification

- Undilatable lesion or severely calcified lesion
- Difficult to cross balloon or stent
- Stent ablation

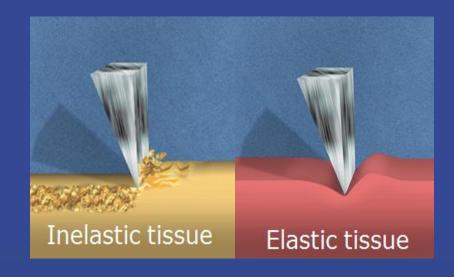
Relative contraindication

- Severe angulation
- Extremely eccentric lesion
- Vessel size is too small
- Pre-existing severe dissection or vasospasm
- High risk of no-reflow: thrombotic lesion, SVG





Principles of Rotational Atherectomy



Differential Cutting



Orthogonal displacement of friction



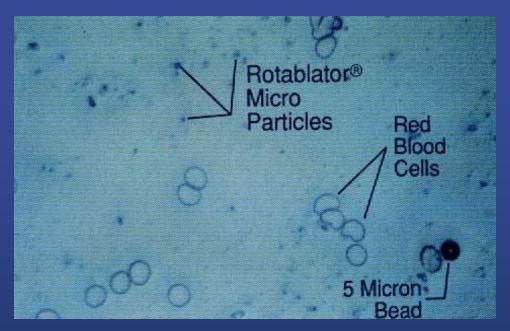


Microparticulate Debris

- Size: < 12 micron in 88%
- Increased size of debris when
 - Slow burr speed
 - Deceleration by pushing hard

< 75,000 rpm

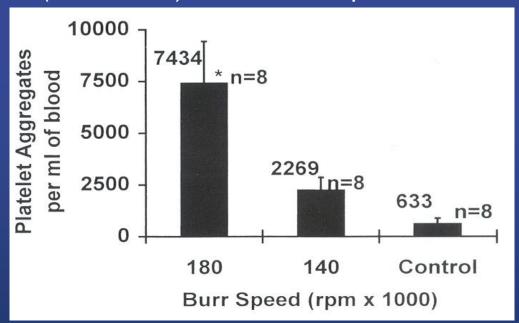
> 5000 rpm





Burr Selection

- Burr-to-artery ratio: upto 0.5
- One-burr vs. two-burr approach
- Burr speed
 - Large burr (≥ 2.0 mm) : 150,000 rpm
 - Small burr (≤ 1.75 mm) : 180,000 rpm





Burr Size and Guiding Catheter

| Rotablator Burr Size (mm) | Burr Diameter Inches/mm | Recommended Guide Catheter (Fr) | Minimum ID (Inches/mm) |
|------------------------------|----------------------------|------------------------------------|---------------------------|
| 1.25 | 0.049/1.245 | 5-6 | 0.053/1.346 |
| 1.50 | 0.059/1.499 | 6 | 0.063/1.600 |
| 1.75 | 0.069/1.753 | 7 | 0.073/1.854 |
| 2.00 | 0.079/2.007 | 8 | 0.083/2.108 |
| 2.15 | 0.085/2.159 | 8 | 0.089/2.261 |
| 2.25 | 0.089/2.261 | 9 | 0.093/2.362 |
| 2.38 | 0.094/2.388 | 9 | 0.098/2.489 |
| 2.50 | 0.098/2.489 | 10 | 0.102/2.591 |

^{*} Guiding catheters without abrupt primary or secondary curves are recommended (FR4, CLS, XB etc)





Cocktail solution

Infused into Rotalink advancer by pressure-bag (50~100 mmHg above the blood pressure)

• Infusion speed 6-8 ml/30 sec

Contents

Normal saline 500 ml

Nitroglycerin 2 mg

Heparin 2500 unit

Verapamil 5 mg

- Rotaflush study (Matsuo, AHJ 2007)
 - Nicorandil is better than verapamil in terms of ST resolution, and the risk of NQMI and QMI





Complications of Rotablation

- Slow or no-reflow
- Dissection
- Perforation
- Wire bias problems
- Lodged burr
- Spasm
- AV block







Slow Flow / No Flow

Overview

- Slow flow and no flow are observed in 5% of patients undergoing PTCRA
- Slow flow is a diminution of flow by 1-2 TIMI grades from the baseline antegrade flow
- No reflow is a cessation of flow into the distal coronary bed

Potential Course of Action

- Early recognition of flow disturbance is key
- Time
- IC Nitroprusside, verapamil or adenosine: careful of hypotension and bradycardia
- IABP if needed
- Intermittent injections of contrast media during ablation run for flow interrogation
- Appropriate burr run time for lesion and vessel complexity



Lodged Burr

Causes

- Oversized burr in diffuse calcium and too much pressure can jam
- Small burr in eccentric lesion and too much pressure can cause watermelon seeding thru lesion and with no diamonds on proximal side of burr, no way to get back

Potential Course of Action

- Do not attempt to start the burr spinning once it is stuck. Take an angiogram to determine burr position
- Nitro, cough and time
- DynaGlide[™]: Burp foot pedal while gently pulling catheter shaft. Brief spurt of energy and gentle pull back simultaneously
- Buddy wire with 1.5 mm ballooning if possible
- Pull the burr very hard, as the last resort!
- Surgery if required



AV Block

- Causes
 - No flow or slow flow for AV nodal branch
 - RCA > LCX > LAD
- Course of action
 - Cough CPR
 - Atropine
 - Temporary pacemaker



Incidence and Determinants of Complications in Rotational Atherectomy (J-PCI Registry)

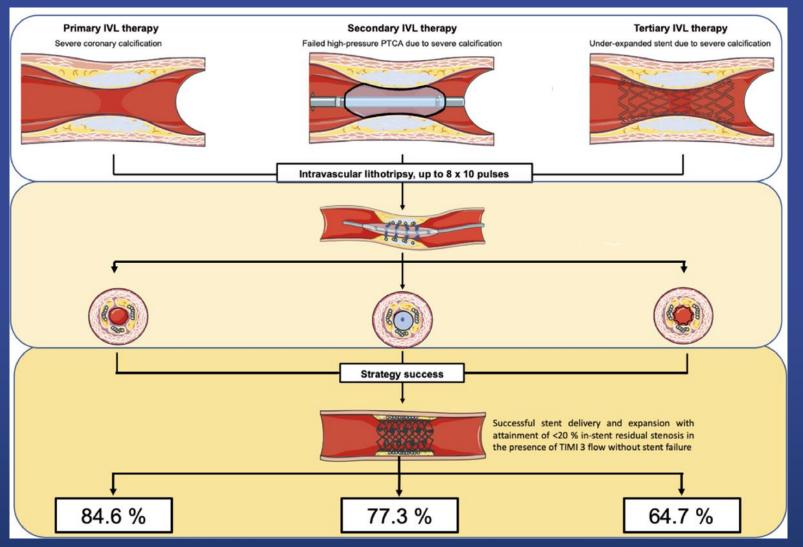
In hospital death, cardiac tamponade, emergent surgery

| | OR | 95% CI | P Value |
|---|------|-----------|---------|
| Age (1-y increase) | 1.03 | 1.02-1.05 | <0.001 |
| Impaired kidney function | 1.59 | 1.15–2.19 | 0.004 |
| History of previous myocardial infarction | 1.69 | 1.21-2.35 | 0.002 |
| Emergent PCI | 4.02 | 1.66-8.27 | <0.001 |
| Triple-vessel disease (vs single-vessel disease) | 2.17 | 1.43-3.28 | <0.001 |
| Left main disease (vs single-vessel disease) | 2.54 | 1.51–4.17 | <0.001 |
| High-volume institution (vs low-volume institution) | 0.56 | 0.36-0.89 | 0.011 |

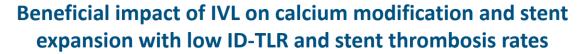


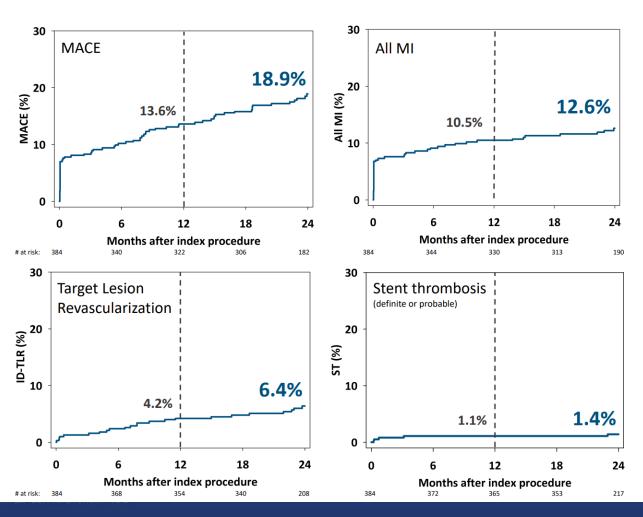


Intravascular coronary lithotripsy



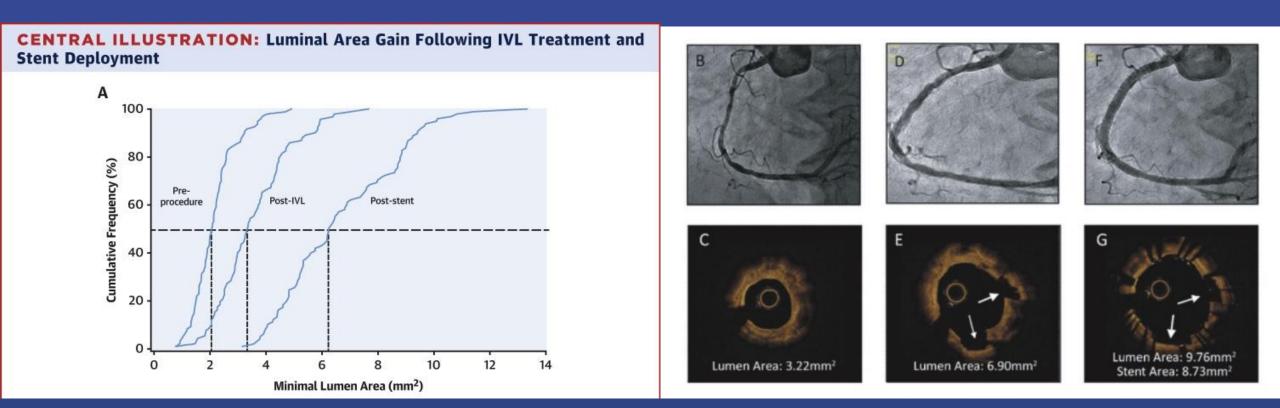
Intravascular coronary lithotripsy





Beneficial impact of IVL on calcium modification and stent expansion with low ID-TLR and stent thrombosis rates - Final 2-year result from the Disrupt CAD III study

Intravascular coronary lithotripsy



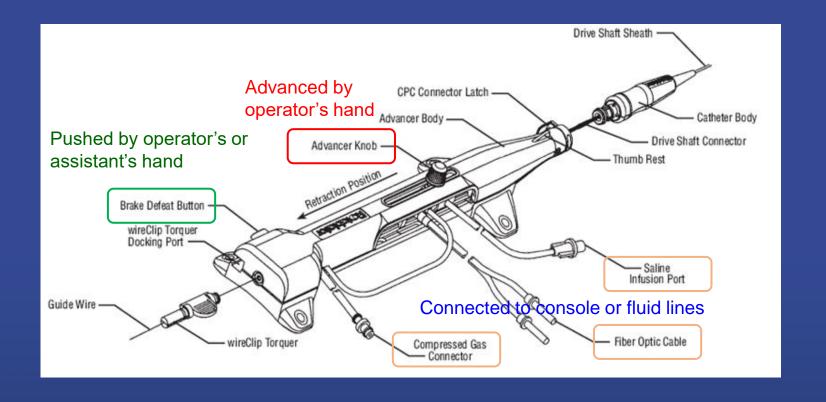
Coronary IVL safely and effectively facilitated stent implantation in severely calcified lesions



Technical Issues



Rotalink advancer







Basic procedural steps (1)

- 1. Place the rota-wire beyond lesion
 - Rota-wire is very delicate. No severe bends
 - Rota-wire has poor torque conduction. Use microcatheter or over-the-wire balloon to exchange with conventional guidewire.
- 2. Select burr size: Burr-to-artery ration < 0.5 0.7
- 3. Backload and advance assembled burr + advancer unit over rota-wire. Place wire clip at the end of rota-wire.



- 4. <u>Lock advancer knob 2 to 3cm forward</u> before advancing burr into guiding catheter. Turn on the flush solution and do <u>brief RPM check</u> while holding the Y-connector firmly.
 - It removes tension/inertia on the burr (sudden burr advancement or jump)
 - (Cover the burr with wet gauze to prevent damage)







Basic procedural steps (2)

5. Press Dynaglide button to activate Dynaglide mode (60,000 – 80,000 rpm).



- 6. Advance the burr to the '<u>landing zone</u>' (non-stenotic site proximal to the lesion) in the proximal coronary artery
 - Avoid tightening of Y-connector. The hemostasis valve should be closed just tight enough to prevent blood loss, but still allow the RotaLink Sheath to slide through the valve.
- 7. Remove residual tension/inertia of burr at landing zone
 - Move advancer knob back and forth to remove tension between drive shaft and Teflon sleeve
 - Release Y-connector and move burr back and forth to remove tension between guidewire and rota burr
 - Brief Dynaglide run under fluoroscopic guidance. If there is residual tension/inertia, sudden burr advancement or jump occurs.





Basic procedural steps (3)

8. Basics of rotablation

- 1. Burr motion: To-and-fro pecking motion > slow advancement
- 2. Burr run time: the shorter is the better, 15–20 sec
- 3. Burr speed: the higher is the better, > 180,000 rpm*
- 4. Advance burr no more than 3 cm back and forth. Moving forward only when there is <u>light resistance</u>.
- 5. Avoid running the burr in static position. Always keep the burr advancing or retracting while it is rotating.
- 6. Avoid significant drop in rpm (> 5000 RPM for > 5 sec)
- 7. Aggressively keep blood pressure and heart rate.
- 8. Do final 'polish run' (no rpm drop, no resistance) after completion of rotablation.
- 9. Long lesions were divided into segments and each segment was separately ablated.





Basic procedural steps (4)

- 9. Get feedback of rotablation
 - Tactile: advancer knob resistance or driveshaft vibration
 - excessive load on burr
 - too rapid advancement
 - a kink in the drive shaft coil
 - too large burr
 - Visual: smooth advancement under fluoroscopy
 - Auditory: Pitch changes relative to resistance encountered by the burr

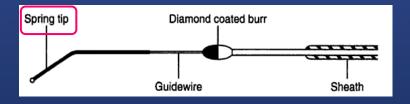


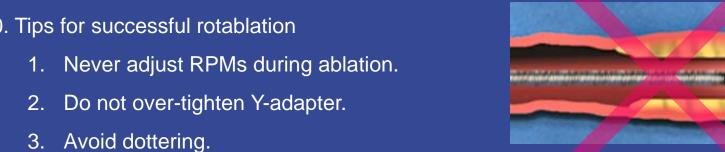


Basic procedural steps (5)

10. Tips for successful rotablation

- Avoid burring in the guide catheter (except Dynaglide mode).
- Gently advance or retract the burr while it is at high speed rotary motion.
- Never stop burr in lesion or distal to lesion. Burr should be located at the proximal 'landing zone' or within guiding catheter when not running.
- Do not allow the burr to remain in any location while rotating at high speeds. Always keep the burr advancing or retracting while it is rotating.
- Never advance rotating burr to point of contact with the guidewire spring tip. The guidewire can be destructed easily.









Procedure

- Place the rota-wire beyond lesion
 Easy to bend, poor torque control
 Use microcatheter or OTW balloon for wire exchange
- 2. Select burr size: Burr-to-artery ration ≈ 0.5
- 3. Backload and RPM check
 150K RPM for 1.75 or larger burr, 180K for smaller burr
- 4. Advance the burr upto landing zone (with or without dynaglide)
- 5. Tension release and dye injection
- 6. Start ablation
 - 1) Burr motion: To-and-fro pecking motion for 15~20 sec
 - 2) Never stop burr in lesion or distal to lesion
 - 3) Get feedback: visual, auditory for drop in rpm > 5000 RPM for > 5 sec
 - 4) Intermittent dye injection for slow flow
 - 5) Polish run after cross
- 7. Remove burr using dynaglide



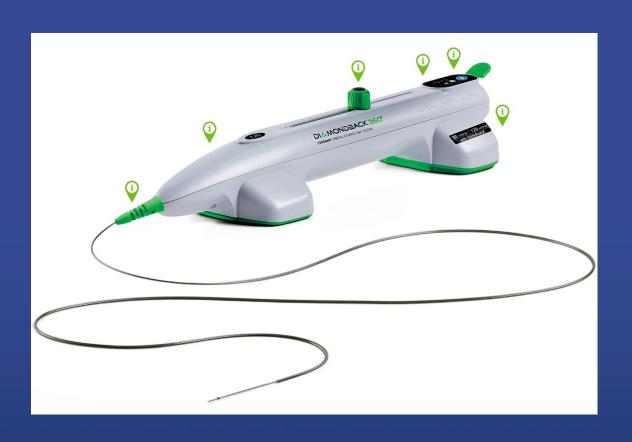


DIAMONDBACK 360: Coronary Orbital Atherectomy System





Orbital atherectomy



A DEEPER LOOK

Differential Sanding⁴

The diamond-coated crown sands intimal calcium into particulate with an average size of approximately 2 μ m – which is smaller than a capillary vessel.



Pulsatile Forces¹⁻⁴

The pulsatile impact of the crown may facilitate fracture of deep calcium.

Procedural Safety⁵

With the Diamondback 360 Oronary Orbital
Atherectomy System, healthy tissue safely flexes away
from the crown during operation, reducing impact to the
medial layer. The orbital movement of the crown allows
blood and saline to flow continuously during procedures,
minimizing risk of thermal injury and slow flow/no reflow
events.

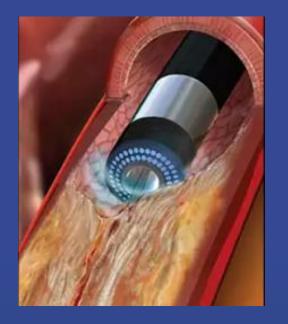


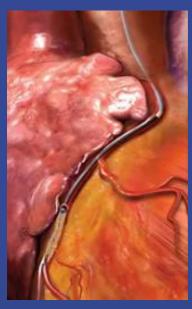




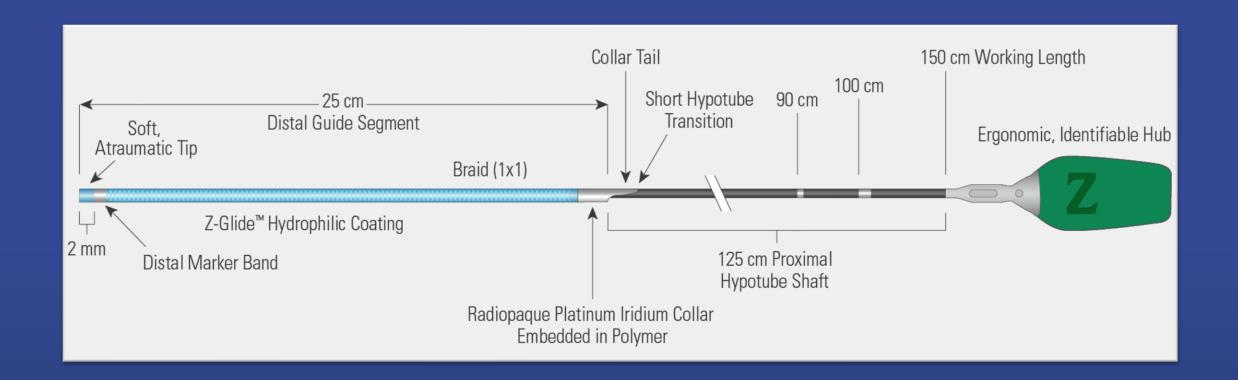
Laser atherectomy







Guidezilla II







Guidezilla II

Powerful Reach. Predictable Performance.

Short Hypotube Transition for reduced device interaction

Radiopaque Helical Collar
Designed for improved
strength and visibility

Z-Glide[™] Coating For improved deliverability

Green Ergonomic Hub
Unique and easily
identifiable



Expanded Size Matrix
6, 7 & 8F 25 cm;
6F 40cm
(Rapid Exchange
Length Noted)



Design Changes (Guidezilla to GUIDEZILLA™ II)

| Features | Guidezilla | GUIDEZILLA II | Design Goal | |
|------------------------|----------------------------------|---|---|--|
| Sizes | 6F | 6F, 7F, 8F, and 6F Long | Expanded Size Matrix | |
| Guide Segment | 25 cm | 25 cm on 6F,7F,8F (40 cm on 6F Long) | 40cm 6F Long Designed for TRI | |
| Working Length | 145cm | 150cm | Extra 5 cm Proximal Hypotube Shaft | |
| Collar | Stainless Steel | Helical Platinum Iridium | Visibility, Strength, and Smooth Device Interaction | |
| Coating | Bioslide™ | Z-Glide™ | Deliverability | |
| Radiopaque | Distal Marker Proximal Marker | Distal Marker band Radiopaque Collar | True Device Positioning with Added Visibility | |
| Hypotube Transition | 19mm | 6mm | Optimized to Reduce Device Interaction | |
| Hub Design | | | | |

Guidezilla



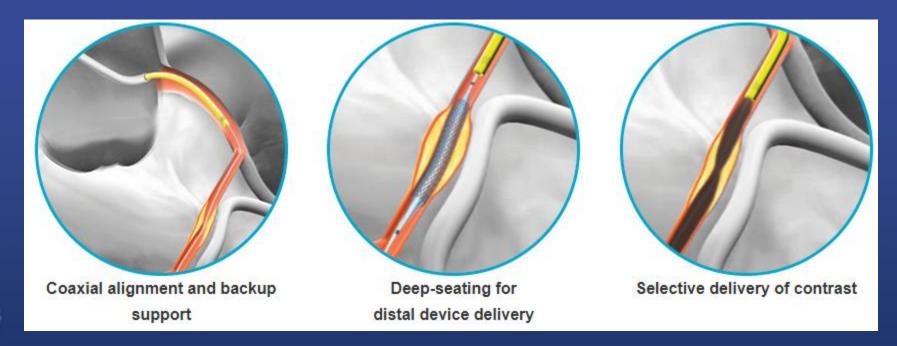


GUIDEZILLA II



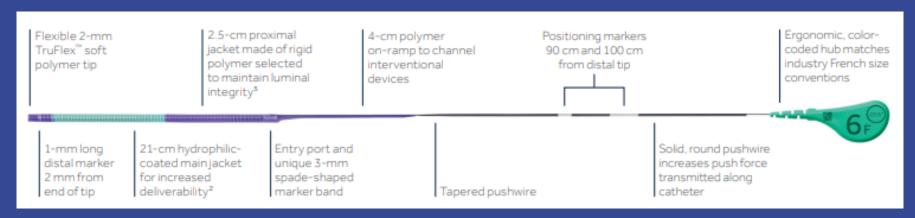
GuideLiner

GuideLiner®
V3 Catheter
Beyond Tried. True.





Telescope



TECHNICAL FEATURES

| Technical Features | Telescope™ GEC |
|-----------------------------------|--|
| Catheter length | 150 cm |
| Distal extension length | 25 cm |
| Marker band material | Platinum iridium |
| Marker band lengths and locations | 1 mm long, 2 mm from distal tip 3 mm long, spade-shaped at entry port |
| Coating | Hydrophilic, outer layer of distal 21 cm |
| Pushwire length | 125 cm |
| Tapered pushwire portion | 10 cm |
| On-ramp length | 4 cm |
| On-ramp material | Nylon-based polymer |
| TruFlex™ tip | 2 mm |
| Shelf life | 2 years |

DIMENSIONAL COMPARISON

| French Size (F) | GEC Name | I.D. (in) | O.D. (in) | Required GC I.D. (in) |
|--------------------|----------------------------------|-----------|-----------|--------------------------|
| 5.5 | GuideLiner™* V3 GEC ⁶ | 0.051 | 0.063 | 6 F ≥ 0.066 |
| | Telescope™ GEC | 0.056 | 0.067 | 6 F ≥ 0.070 |
| 6 | GuideLiner™* V3 GEC ⁶ | 0.056 | 0.067 | 6 F ≥ 0.070 |
| 6 | Guidezilla™* II GEC ⁷ | 0.057 | 0.067 | 6 F ≥ 0.070 |
| | Telescope™ GEC | 0.062 | 0.075 | 7 F ≥ 0.078 |
| 7 | GuideLiner™* V3 GEC ⁶ | 0.062 | 0.075 | 7 F ≥ 0.078 |
| 7 | Guidezilla™* II GEC ⁷ | 0.063 | 0.073 | 7 F ≥ 0.078 |





Heartrail

Large I.D. & Superb Back up force







ARTIST trial

Balloon angioplasty (PTCA) vs. Rotablation in ISR (PTCR)

| TABLE 4 Angiographic Outcome | | | |
|---|-------------------|-------------------|---------|
| | PTCA (n = 138) | PTCR (n = 139) | p Value |
| MLD after rotational ablation (mm) | _ | 1.33 ± 0.39 | |
| Mean diameter after rotational ablation (mm) | _ | 1.7 ± 0.28 | |
| Diameter stenosis after rotational ablation (%) | _ | 35 ± 15 | |
| Final MLD (mm) | 1.9 ± 0.3 | 1.9 ± 0.4 | 0.57 |
| Final mean diameter (mm) | 2.2 ± 0.35 | 2.2 ± 0.37 | 0.2 |
| Final diameter stenosois (%) | 29 ± 10 | 28 ± 12 | 0.38 |
| Acute gain (mm) | 1.3 ± 0.4 | 1.4 ± 0.4 | 0.45 |
| Acute gain index | 50 ± 16 | 52 ± 16 | 0.43 |
| Final plaque area (mm²) | 6.4 ± 5.2 | 6.8 ± 5.4 | 0.55 |
| Plaque area reduction (%) | 69 ± 17 | 68 ± 17 | 0.68 |
| Angiographic success | 139/146 (95%) | 144/152 (94%) | 1.0 |
| Diameter stenosis ≤30% | 78/137 (57%) | 87/143 (61%) | 0.54 |

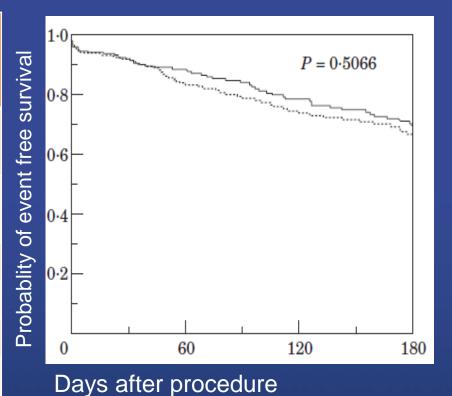
| TABLE 5 Angiographic Outcome After Six Months | | | | |
|---|-------------------|-------------------|---------|--|
| | PTCA (n = 123) | PTCR (n = 131) | p Value | |
| Diameter stenosis (%) | 56 ± 20 | 64 ± 22 | 0.005 | |
| MLD (mm) | 1.2 ± 0.6 | 1.0 ± 0.6 | 0.008 | |
| Mean stenosis diameter (mm) | 1.83 ± 0.74 | 1.7 ± 0.45 | 0.03 | |
| Late loss (mm) | 0.68 ± 0.5 | 0.92 ± 0.6 | 0.0015 | |
| Loss index | 50 ± 46 | 69 ± 42 | 0.0007 | |
| Net gain (mm) | 0.67 ± 0.5 | 0.45 ± 0.6 | 0.0019 | |
| Net gain index | 24.5 ± 20 | 16.8 ± 22 | 0.005 | |
| Neo-plaque area (mm²) | 5.1 ± 5.8 | 6.1 ± 6.3 | 0.25 | |
| Net plague reduction (mm ²) | 11.6 ± 14 | 7.9 ± 12 | 0.04 | |
| Restenosis rate (%) | 51.2 | 64.9 | 0.027 | |



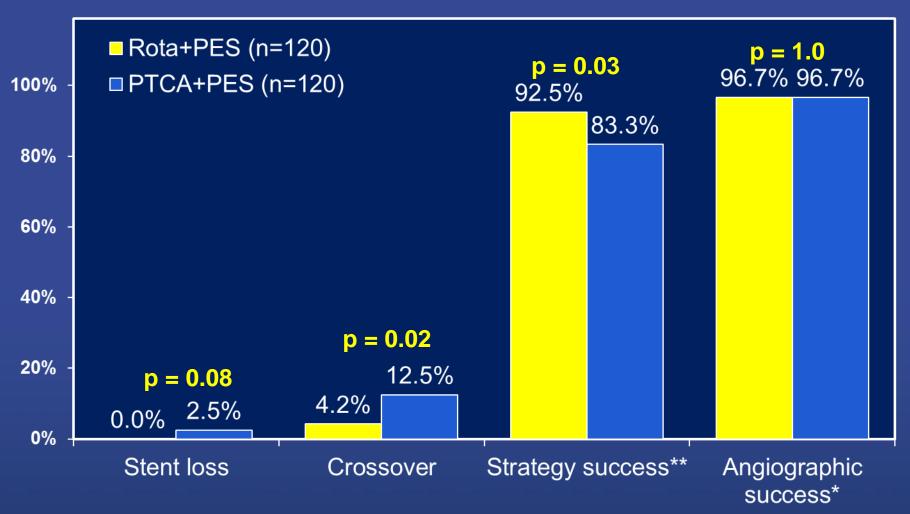


COBRA study
A randomized comparison of balloon angioplasty versus rotational atherectomy in complex coronary **lesions**

| | RA (n=25 2) | PTCA (n=250) | P value |
|--------------------|-------------------|-----------------|---------|
| Procedural success | 85% | 78% | 0.038 |
| 6 months success | 48.9% | 51.1% | 0.333 |
| Major cardiac | events d | uring follow | up |
| Q wave MI | 0.5% | 0% | |
| CABG | 4.2% | 6.5% | |
| Death | 0% | 0% | |



ROTAXUS; Procedural Outcomes



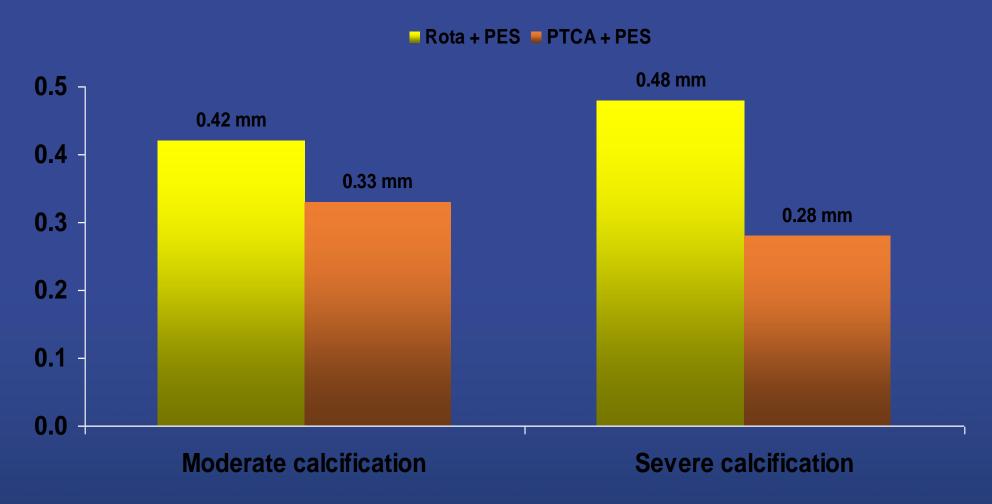
^{*} Defined as <20% residual stenosis + TIMI 3 flow





^{**} Defined as angiographic success with no crossover or stent loss

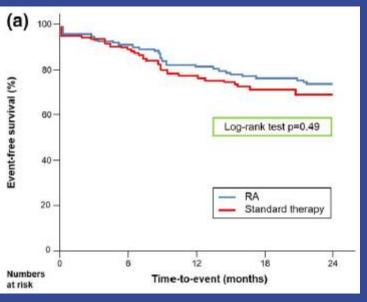
ROTAXUS Strategy Success according to calcification

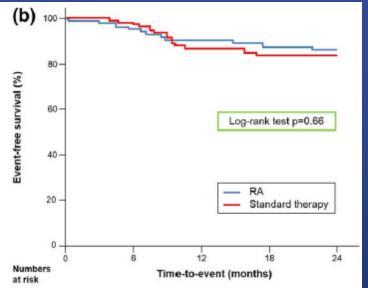






Rotaxus; 2 year clinical outcome





| | RA+DES (n=109) | DES (n=108) | P value |
|-------------------|-------------------|----------------|---------|
| Procedure success | 92.5% | 83.3 | 0.03 |
| MACE | 29.4% | 34.3% | 0.47 |
| Death | 8.3% | 7.4% | 1.00 |
| MI | 8.3% | 6.5% | 0.8 |
| TLR | 13.8% | 16.7% | 0.58 |
| TVR | 19.3% | 22.2% | 0.62 |

Increse procedure success
But does not increase clinical outcome

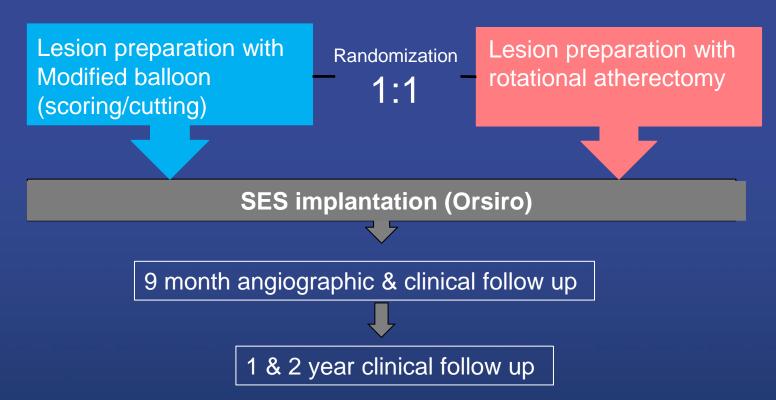


PREPARE-CALC Trial

Study design

Prospective, randomized, active controlled clinical trial in 2 German centers

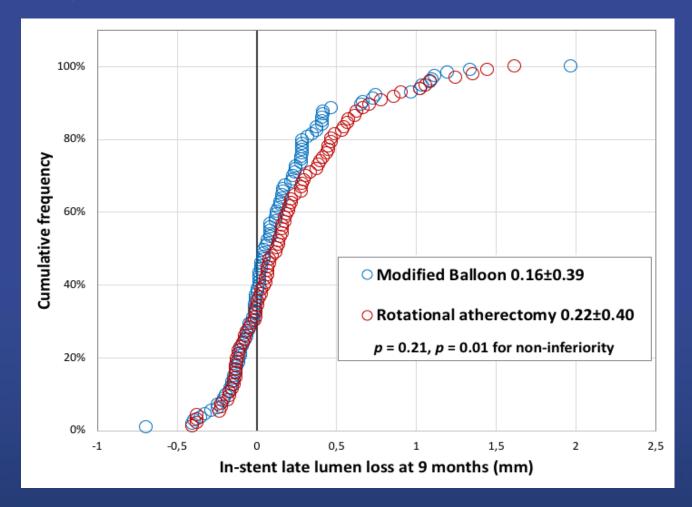
PCI in 200 patients with severely calcified lesions







PREPARE-CALC Trial Co-Primary Endpoint – In stent LLL at 9 Month







PREPARE-CALC Trial QCA 9 months

| | Modified balloon (n = 112 lesions) | Rotational atherectomy (n = 97 lesions) | p-value |
|-----------------------------|---------------------------------------|---|---------|
| Minimal lumen diameter (mm) | | | |
| In-stent | 2.68±0.59 | 2.64±0.51 | 0.59 |
| In-segment | 2.50±0.54 | 2.50±0.55 | 0.96 |
| Diameter stenosis (%) | | | |
| In-stent | 18.83±13.42 | 19.75±11.54 | 0.49 |
| In-segment | 22.40±11.36 | 23.30±11.43 | 0.52 |
| Late lumen loss (mm) | | | |
| In-stent | 0.16±0.40 | 0.22±0.41 | 0.21 |
| In-segment | 0.07±0.52 | 0.18±0.74 | 0.25 |
| Binary restenosis (%) | | | |
| In-stent | 6 (5.3%) | 2 (2.1%) | 0.30 |
| In-segment | 5 (4.5%) | 2 (2.1%) | 0.32 |





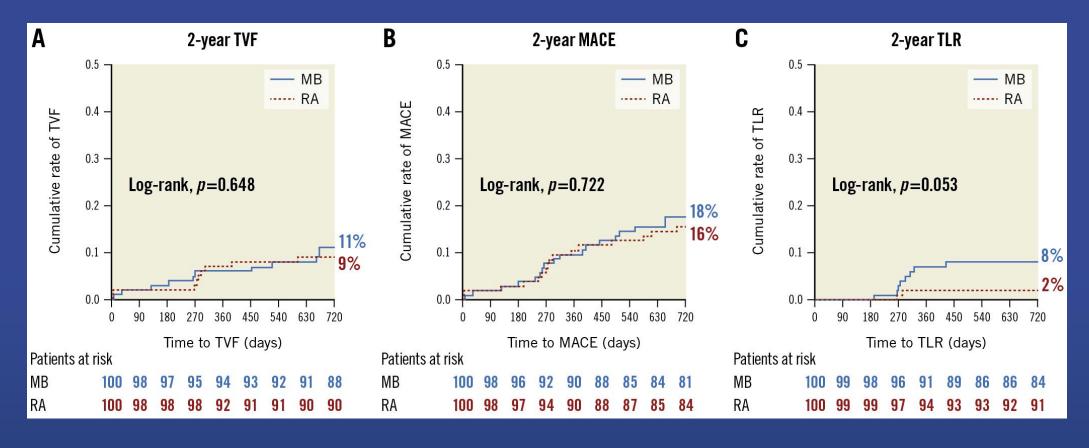
PREPARE-CALC Trial Clinical outcome 9 months

| | Modified balloon (n = 100 pts.) | Rotational atherectomy (n = 100 pts.) | p-value |
|-------------------------------|------------------------------------|---------------------------------------|---------|
| Death | 2 (2%) | 2 (2%) | 1.00 |
| Cardiac death | 1 (1%) | 1 (1%) | 1.00 |
| Non-cardiac death | 1 (1%) | 1 (1%) | 1.00 |
| Myocardial infarction | 3 (3%) | 2 (2%) | 1.00 |
| Target vessel MI | 1 (1%) | 2 (2%) | 1.00 |
| Periprocedural MI | 1 (1%) | 2 (2%) | 1.00 |
| Spontaneous MI | 2 (2%) | 0 (0%) | 0.50 |
| Stent thrombosis (def./prob.) | 0 (0%) | 0 (0%) | 1.00 |
| TVR | 8 (8%) | 3 (3%) | 0.21 |
| Target vessel failure | 8 (8%) | 6 (6%) | 0.78 |





PREPARE-CALC Trial Clinical outcome 2-year







PREPARE-CALC-COMBO Study

Combined rotational atherectomy and cutting balloon angioplasty prior to drug-eluting stent implantation in severely calcified coronary lesions

 To assess whether the Rota-Cut combination improves stent performance in severely calcified coronary lesions

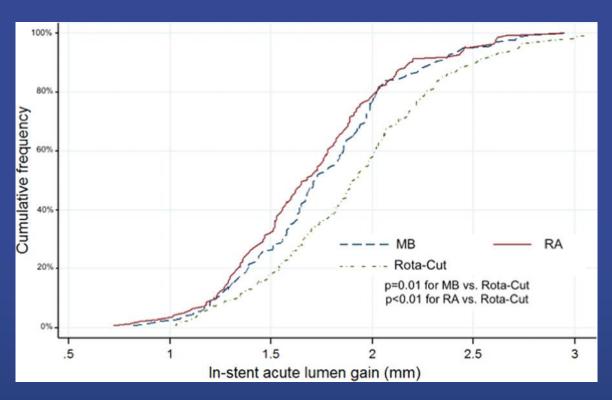
Prospective, single-arm, single center study

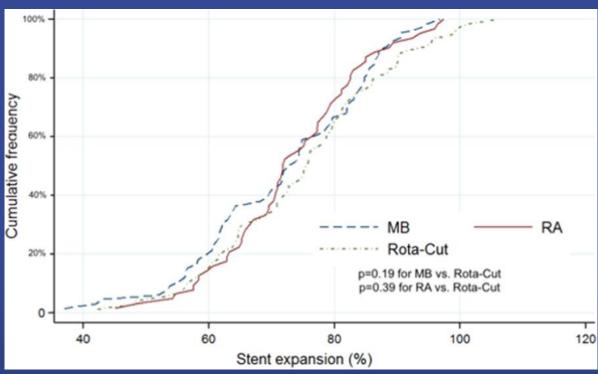
Primary endpoint : in-stent acute lumen gain(ALG), stent expansion(SE)



PREPARE-CALC-COMBO Study

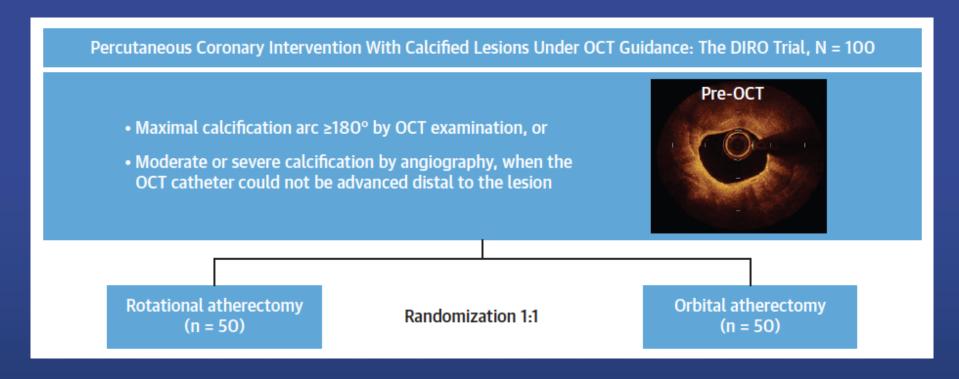
Combined rotational atherectomy and cutting balloon angioplasty prior to drug-eluting stent implantation in severely calcified coronary lesions





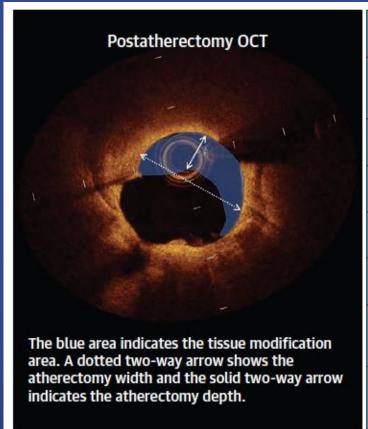
DIRO Study

 Direct Comparison of Rotational vs Orbital Atherectomy for Calcified Lesions Guided by Optical Coherence Tomography





DIRO Study



| | RA | OA | P Value |
|---|-------------------|-------------------|---------|
| Maximum tissue modification area, mm² | 1.24 (0.84-1.74) | 0.89 (0.59-1.11) | <0.01 |
| Atherectomy width, mm | 1.50 (1.32-1.89) | 1.22 (1.12-1.40) | <0.01 |
| Atherectomy depth, mm | 0.54 (0.39-0.83) | 0.55 (0.31-0.73) | 0.62 |
| Percentage of lumen area increase, % | 72.2 (49.0-98.3) | 39.2 (17.0-48.1) | <0.01 |
| Ratio of atherectomy width to burr size | 0.94 (0.79-0.98) | 0.98 (0.89-1.12) | 0.03 |
| Stent expansion assessed by distal reference, % | 99.5 (89.3-107.3) | 90.6 (80.0-102.3) | 0.02 |
| Stent expansion assessed by mean reference, % | 72.2 (60.6-86.3) | 64.1 (54.0-77.7) | 0.05 |

- Procedural outcomes including periprocedural MI were comparable
- · Clinical outcomes at 8 months were similar



ECLIPSE

Evaluation of Treatment Strategies for Severe CaLciflc Coronary Arteries: Orbital Atherectomy vs. Conventional Angioplasty Prior to Implantation of Drug Eluting StEnts

~2000 pts with severely calcified lesions; ~150 US sites

Randomize

Orbital Atherectomy Strategy

(1.25 mm Classic Crown followed by balloon pre-dilation)

2nd generation DES implantation and optimization

1:1

Conventional Angioplasty Strategy

(Conventional and/or specialty balloons per operator discretion)

2nd generation DES implantation and optimization

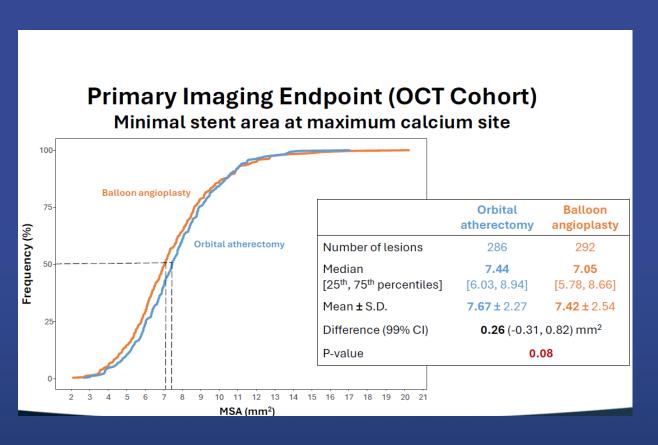
- 1° endpoints: 1) Post-PCI in-stent MSA by OCT (N~500 in imaging sub-study)
 - 2) 1-year TVF (all subjects)
- 2° endpoints: 1) Procedural Success (Stent deployed w/RS<20% & no maj complications)
 - 2) Strategy Success (Procedural success w/out crossover)

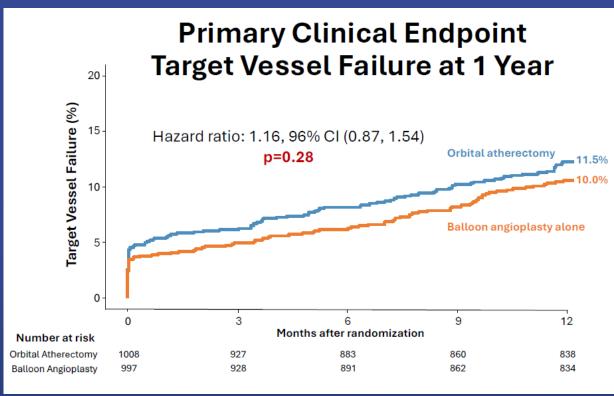




ECLIPSE

Evaluation of Treatment Strategies for Severe CaLciflc Coronary Arteries: Orbital Atherectomy vs. Conventional Angioplasty Prior to Implantation of Drug Eluting StEnts







Disrupt CAD III

Intravascular Lithotripsy for Treatment of Severely Calcified Coronary Artery Disease

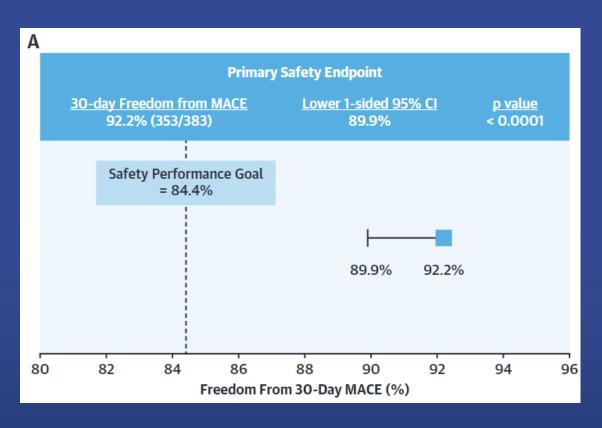
 To assess safety and effectiveness of IVL in severely calcified de novo coronary lesions

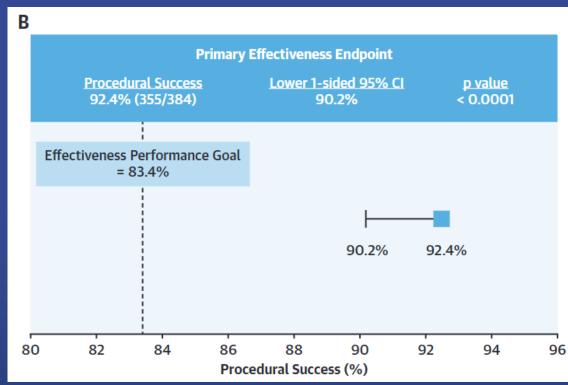
- Prospective, single-arm multicenter study
- Primary safety endpoint: freedom from major adverse cardiovascular events (cardiac death. MI, or target vessel revascularization) at 30 days
- Primary effectiveness endpoint : procedural success



Disrupt CAD III

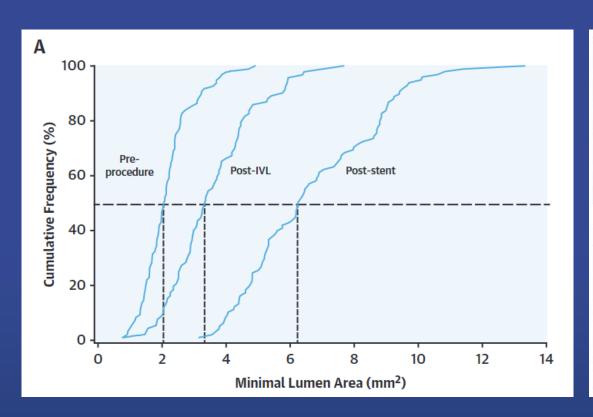
Intravascular Lithotripsy for Treatment of Severely Calcified Coronary Artery Disease

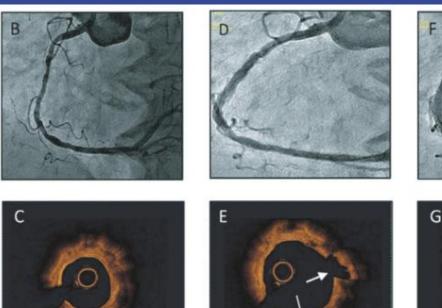




Disrupt CAD III

Intravascular Lithotripsy for Treatment of Severely Calcified Coronary Artery Disease





Lumen Area: 3.22mm²

Lumen Area: 6.90mm³

Lumen Area: 9.76mm²

Stent Area: 8.73mm²

DISRUPT-CAD Studies

| | DISRUPT-CAD | DISRUPT-CAD | DISRUPT-CAD | DISRUPT-CAD IV | Pooled result |
|--------------------|-------------|-------------|-------------|-------------------|---------------|
| Patients | 60 | 120 | 384 | 64 | 628 |
| Procedural success | 95% | 94% | 92.4% | 93.8% | 92.4% |
| Stent delivery | 100% | 100% | 99.2% | 100% | 99.5% |
| Severe dissection | 0% | 0% | 0.3% | 0% | 0.2% |
| Perforation | 0% | 0% | 0.3% | 0% | 0.2% |
| Abrupt closure | 0% | 0% | 0.3% | 0% | 0.2% |
| Slow/no flow | 0% | 0% | 0% | 0% | 0% |

Kereiakes, D.J. et al. J Am Coll Cardiol Intv. 2021; 14 (12) 1337–1348.





ISAR-CALC 2 trial

Randomized Comparison of Strategies to Prepare Severely Calcified Coronary Lesions 2

- To compare a lesion preparation strategy with either super high-pressure balloon or intravascular lithotripsy in severely calcified undilatable coronary lesion
- Prospective, randomized, multicenter, assessors-blind, open-lable study
- Primary end point : final angiographic minimal lumen diameter after stent implantation



ISAR-CALC 2 trial

Randomized Comparison of Strategies to Prepare Severely Calcified Coronary Lesions 2

