



VASCUPEDIA

MY TECHNICAL CHOICE IN THE TREATMENT OF SEVERELY CALCIFIED LESIONS OF THE SFA

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MY CRITERIA

- Age and comorbidity
- Functional status
- Prognosis
- Chronic kidney disease
- Length of the lesion
- Procedural challenges

PROCEDURAL CHALLENGES FOR SEVERELY CALCIFIED LESIONS

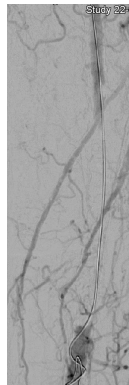
- Crossing
- Lumen gain
- Final treatment



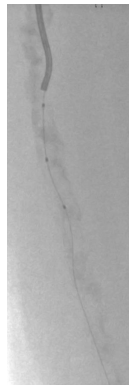
CROSSING THE LESION

MATERIALS

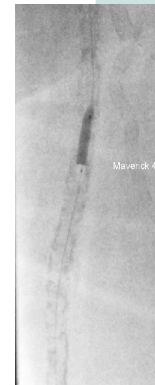
0.018/0.014
wire



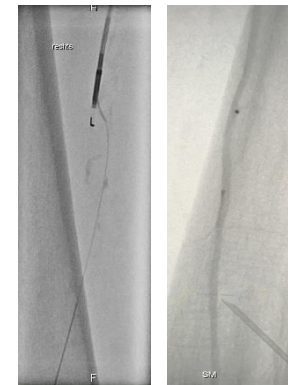
0.018/0.014
support
catheter



Coronary
balloon
catheters



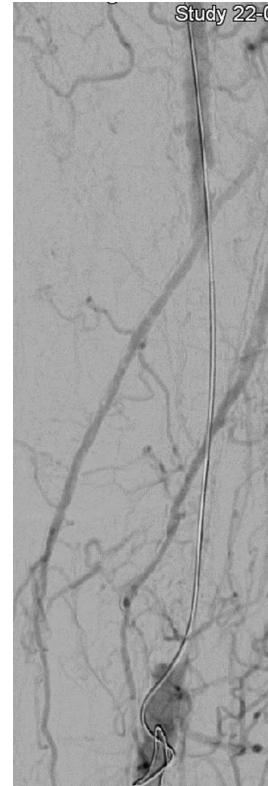
Outback or
retrograde
puncture



WIRES



- Kink resistance and shape retention capability (nitinol core)
- Crossability (hydrophilic coating)
- Steerability



My favorite one:

V18 (Boston Scientific) – 12g
Treasure 12 (Asahi) – 12g

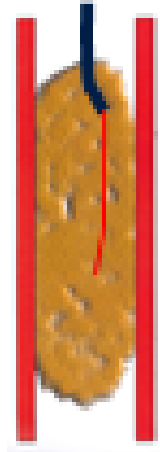


Command 0.014
(Abbott)

Astato 30 (Asahi) – 30g

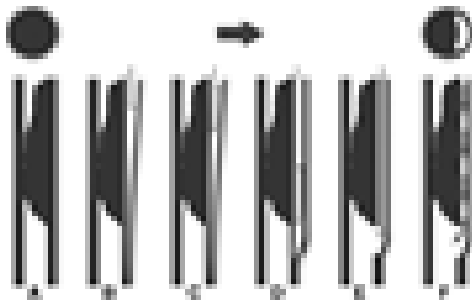
SUPPORT - RE-ENTRY CATHETERS – RETROGRADE APPROACH

Re-Direction

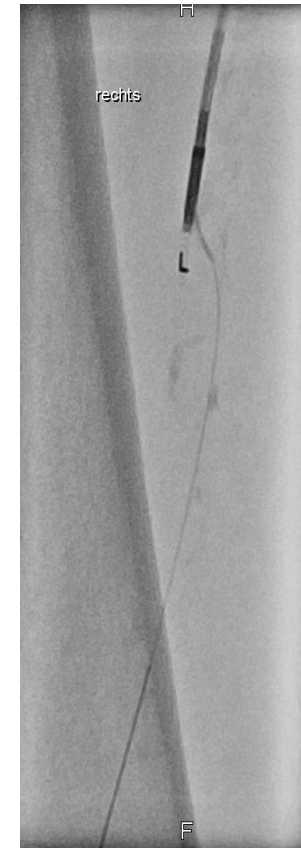


TrailBlazer (Medtronic)
Seeker (Bard)
QuickCross (Philips)

Re-Entry



Angled support catheters / Berenstein catheter



Outback (Cardinal Health)

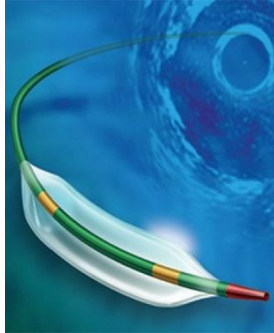


Retrograde tibial puncture

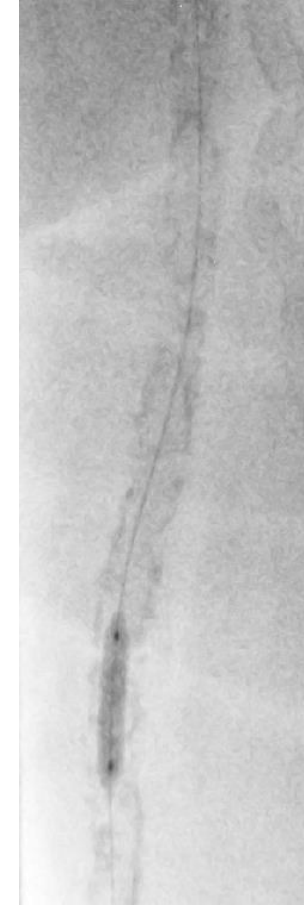
CORONARY BALLOONS



Maverick 4x2



Monorail PTCA
Katheter

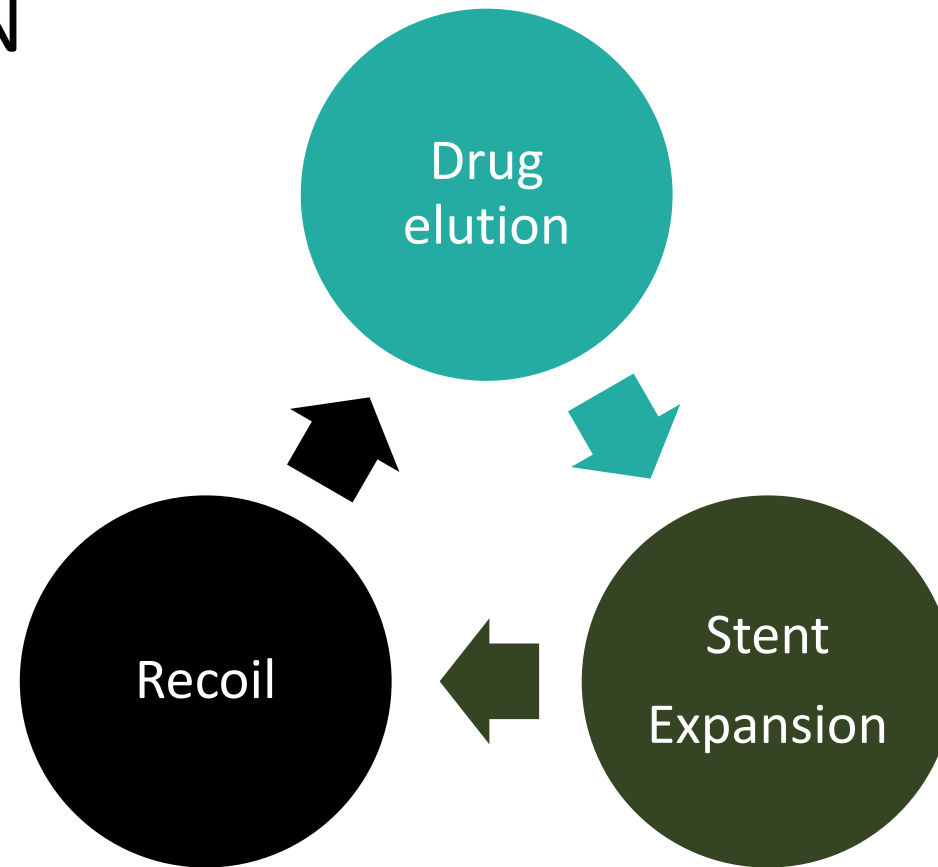


My favorite one:

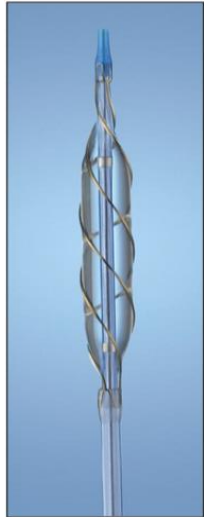
Maverick™ PTCA balloon catheter
Boston Scientific

CHALLENGES IN CALCIFIED LESIONS

LUMEN GAIN & DRUG ELUTION



LUMEN GAIN TOOLS



Scoring
balloon



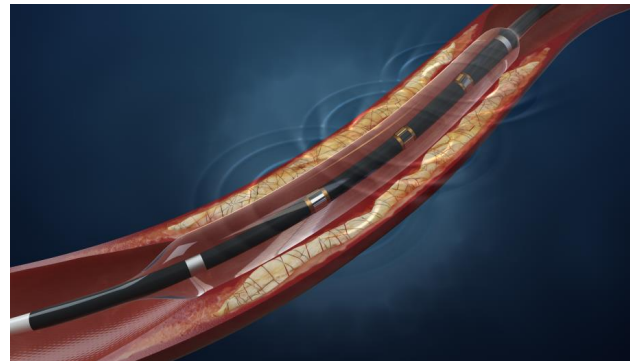
Cutting
balloon



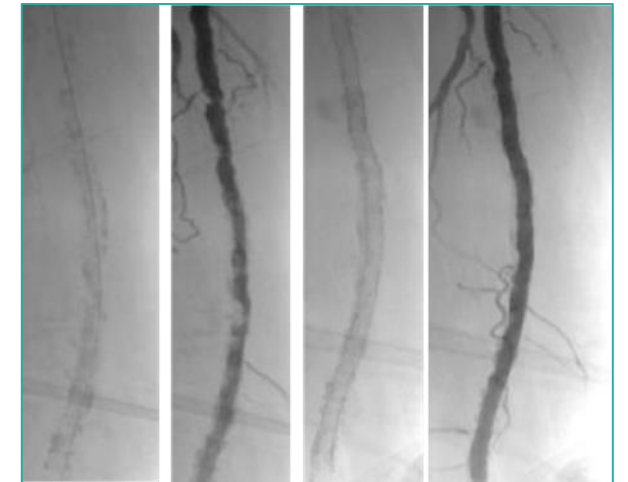
Chocolate
balloon



Atherectomy



Lithoplasty



Pave & Crack

SCORING BALLOON



Subgroups	Total	AS alone	AS + Stent	AS + DCB
	n=124	n=46	n=40	n=38
Lesion Length (mm)	7.4	6.1	10.1	5.9
Occlusions	16.1%	2.2%	31.6%	17.5%
12 mo. primary patency	81.2%	81.5%	77.8%	83.9%

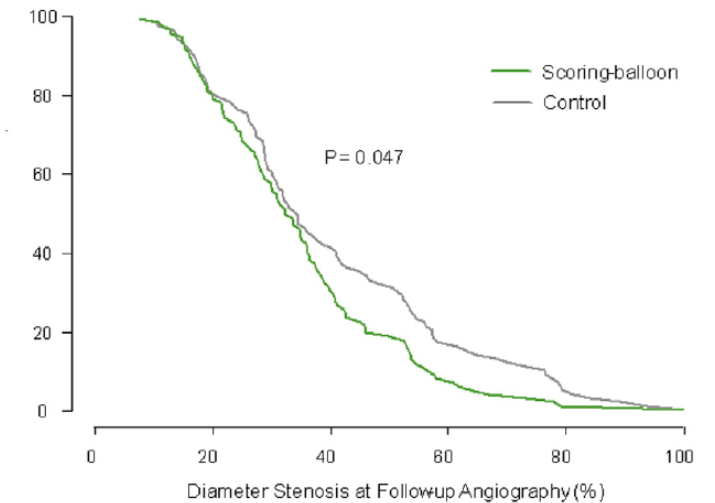
Calcification	Total	Mild	Moderate	Severe
		21.0%	34.7%	43.5%
12 mo. primary patency	81.2%	78.9%	81.3%	81.8%

AS= AngioSculpt®

- After vessel preparation with AngioSculpt®, calcium was not a predictor for loss of patency at 12 months.
- Lesion Preparation with AngioSculpt® in calcified lesions may improve DCB results

PANTHER Registry

FIGURE 1 Cumulative Frequency Distribution Curves for Primary Endpoint According to Treatment Group



Lugenbiel et al, VASA. 2017 Nov 8:1-7
Kufner et al, JACC Intv 2017;10:1332-40

DIRECTIONAL ATHERECTOMY



Medtronic



Case of a severely calcified lesion treated by directional atherectomy (HawkOne, Medtronic)

DIRECTIONAL ATHERECTOMY

DEFINITIVE Ca⁺⁺ TRIAL

Catheterization and Cardiovascular Interventions 84:236-244 (2014)

PERIPHERAL VASCULAR DISEASE

Original Studies

Effective Endovascular Treatment of Calcified Femoropopliteal Disease With Directional Atherectomy and Distal Embolic Protection: Final Results of the DEFINITIVE Ca⁺⁺ Trial

David Roberts,¹ MD, Khusrow Niazi,² MD, William Miller,³ MD, Prakash Krishnan,⁴ MD, Roger Gammon,⁵ MD, Theodore Schreiber,⁶ MD, Nicolas W. Shamma,⁷ MD, MS, and Daniel Clair,⁸ MD on behalf of the DEFINITIVE Ca⁺⁺ Investigators

Objectives: The purpose of the DEFINITIVE Ca⁺⁺ study was to evaluate the safety and effectiveness of directional atherectomy and distal embolic protection, used together to treat moderate to severely calcified femoropopliteal lesions. **Background:** Despite advances in endovascular treatment modalities, treatment of calcified lesions remains a challenge. **Methods:** A total of 133 subjects with 168 moderate to severely calcified lesions were enrolled. Lesions were treated with directional atherectomy devices, coupled with distal embolic protection. **Results:** The 30-day freedom from MAE rate was 93.1%. Per angiographic core laboratory assessment, the primary effectiveness endpoint (<50% residual diameter stenosis) was achieved in 92.0% (lower confidence bound of 87.6%) of lesions. By core lab analysis, these results did not achieve the success criteria (90%) for the primary effectiveness objective. Per site assessment, the objective was met with the endpoint being achieved in 97.0% (lower confidence bound 93.8%). A mean residual diameter stenosis of 33.3% was achieved with the directional atherectomy device. This was further decreased to 24.1% with the use of adjunctive therapy. The proportion of asymptomatic subjects [Rutherford Clinical Category (RCC)=0] increased from 0% at baseline to 52.3% at the 30-day follow-up visit. In total, 88.6% of subjects experienced an improvement of one or more Rutherford categories. **Conclusions:** The results of the DEFINITIVE Ca⁺⁺ study demonstrate that the SilverHawk™ and TurboHawk™ atherectomy devices are safe and effective in the endovascular treatment of moderate to severely calcified lesions in the superficial fem-

TABLE IV. Procedure Characteristics

Procedure characteristic	Mean ± SD or % (n N ⁻¹)
Total procedure time (min)	74.2 ± 28.4 (133)
Total fluoroscopy time (min)	21.1 ± 9.7 (132)
Total contrast administered (cm ³)	182.3 ± 72.4 (132)
Visible debris in filter device	88.4% (122/138)
Adjunctive therapy	53.8% (91/169)
Adjunctive (bail-out) stenting	4.1% (7/169)
Preservation of run-off (per angiographic core lab)	98.3% (113/115)

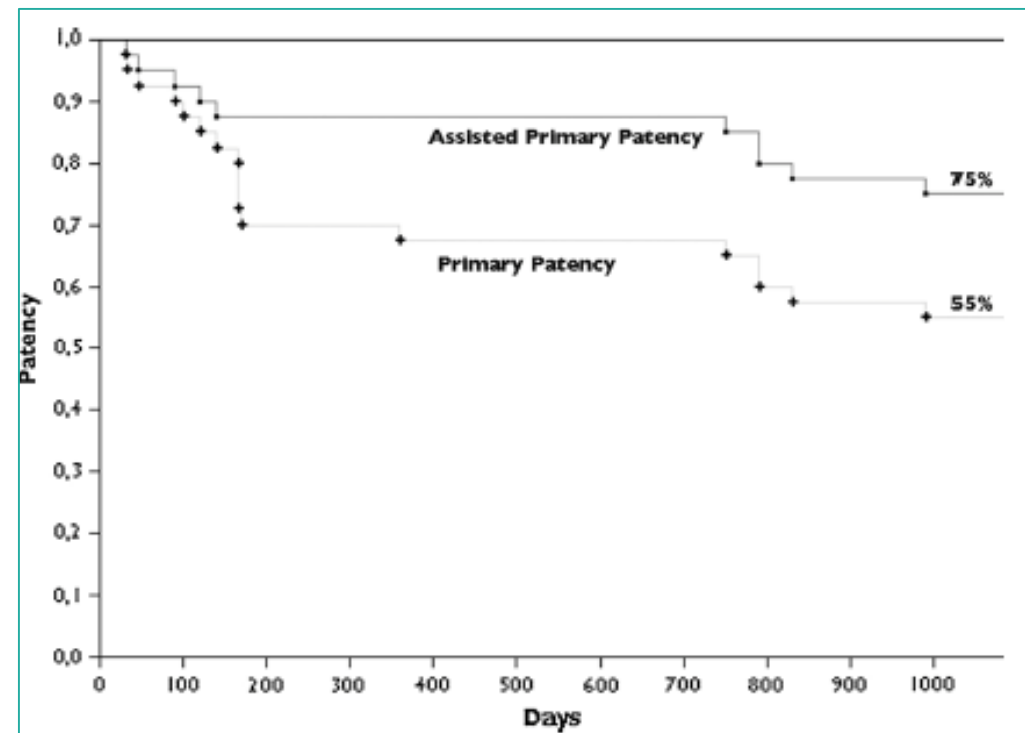
TABLE V. Summary of Major Adverse Events

Major adverse event	% of Subjects (no. of events)
Total	6.9% (9/131) [9]
Death	0.0% (0/131) [0]
Acute myocardial infarction	0.8% (1/131) [1]
Dissection, target vessel (C)	0.0% (0/131) [0]
Dissection, target vessel (grade D or greater)	0.8% (1/131) [1]
Vessel clinical perforation, target vessel	2.3% (3/131) [3]
Pseudoaneurysm, target vessel	0.0% (0/131) [0]
Thrombosis, target vessel	0.8% (1/131) [1]
Distal embolism	2.3% (3/131) [3]
Amputation, above metatarsal line	0.0% (0/131) [0]
Clinically driven TVR	0.0% (0/131) [0]

Primary
effectiveness
92%

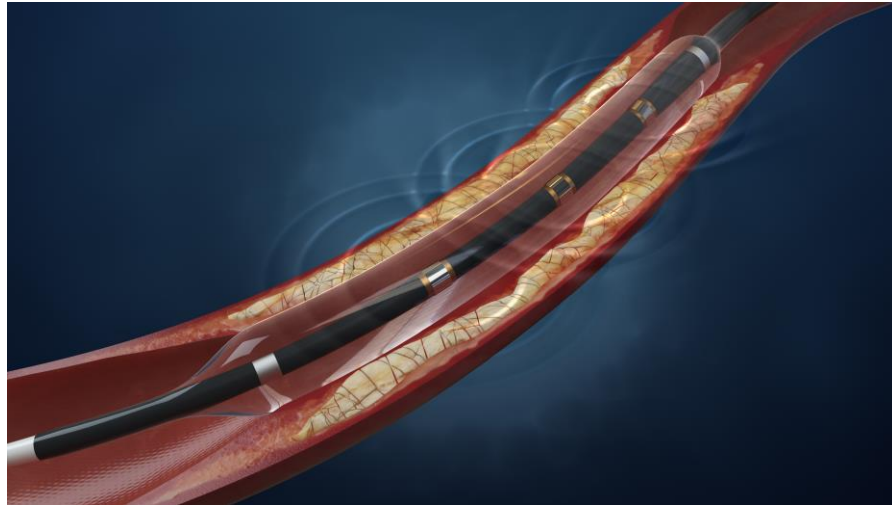
DIRECTIONAL ATHERECTOMY

ADDITIONAL EVIDENCE



Minko et al, Cardiovasc Intervent Radiol (2014) 37:1165–1170

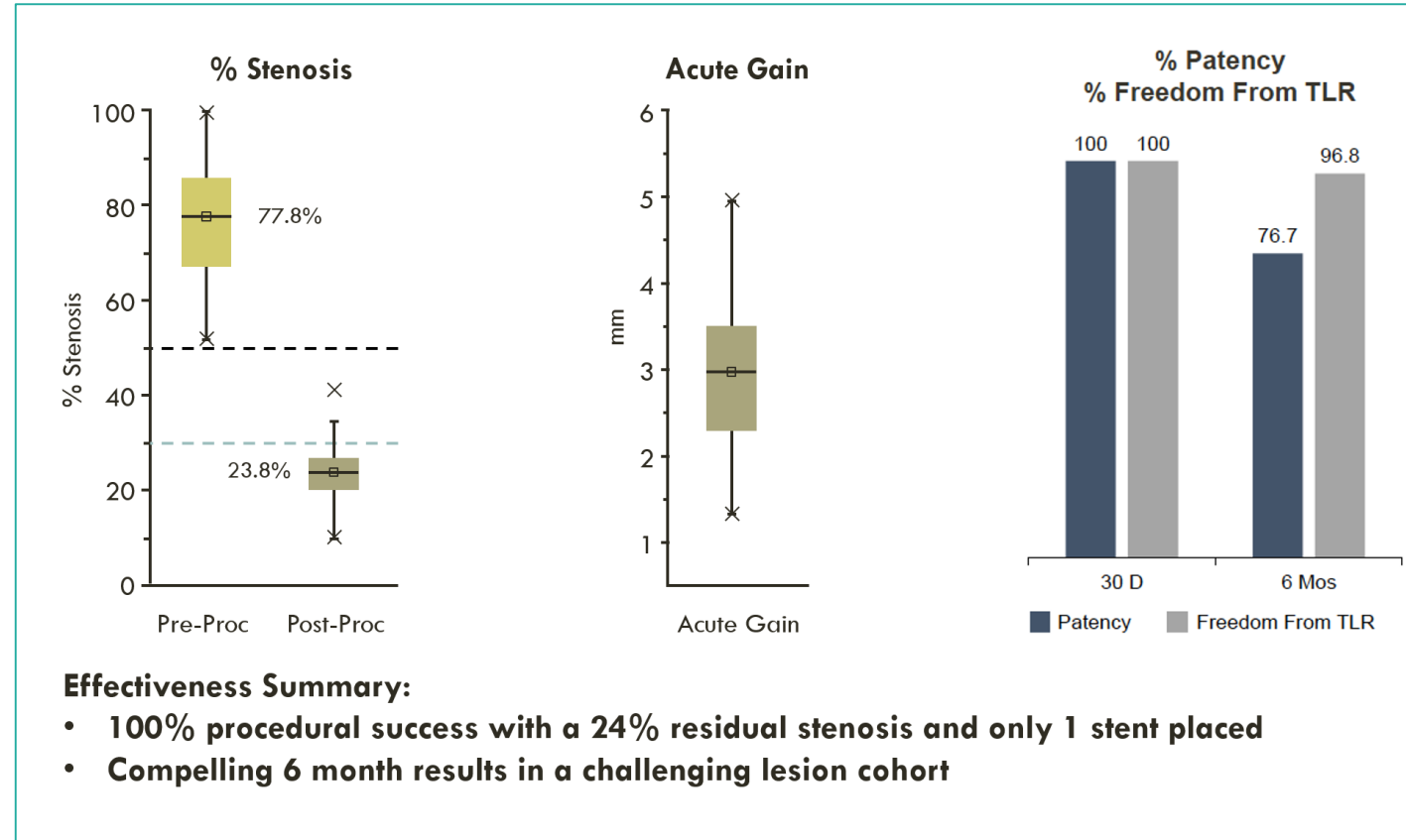
LITHOPLASTY



Schockwave IVL Catheter, Schockwave Medical, Inc



LITHOPLASTY EVIDENCE



DISRUPT PAD trials I & II

PREVENTION OF RESTENOSIS – FINAL TREATMENT

TOOLS



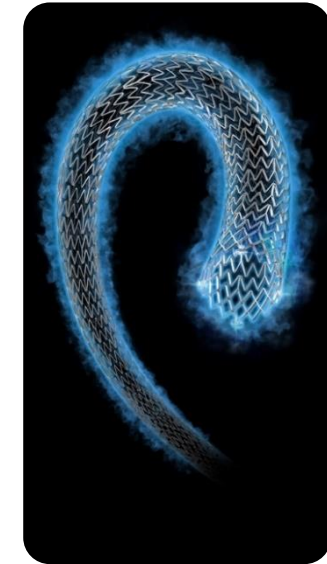
Bare-metal
stent



Drug-coated
balloon



Covered stent



Drug-eluting/coated
stent

BARE METAL STENTS

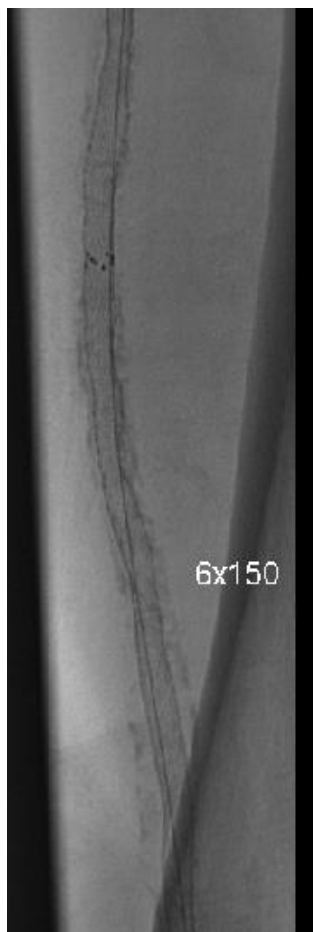


TABLE II. Effects of Patient Characteristics on Outcomes—Univariable Models

Patient characteristic	primary patency			TLR		
	Parameter estimate	Standard error	<i>P</i> -value	Parameter estimate	Standard error	<i>P</i> -value
Age (per 10 year)	0.194	0.072	0.007	-0.167	0.09	0.063
Male	0.265	0.153	0.085	-0.408	0.192	0.034
Diabetes	-0.155	0.148	0.295	0.035	0.19	0.856
Hypertension	0.630	0.211	0.003	-0.150	0.269	0.577
Hyperlipidemia	0.335	0.199	0.092	-0.270	0.235	0.251
Ever Smoker	-0.320	0.173	0.065	0.268	0.222	0.227
Current Smoker	-0.383	0.206	0.063	0.670	0.266	0.012
Rutherford	0.019	0.124	0.876	0.036	0.155	0.812
<i>ABI</i>	1.499	0.423	<.001	-1.490	0.529	0.005
<i>Lesion length (per 10mm)</i>	-0.094	0.016	<.001	0.075	0.023	0.001
Stenosis (per 10%)	-0.023	0.040	0.565	-0.021	0.044	0.640
Calcification	0.107	0.137	0.436	-0.141	0.172	0.412

Rocha-Singh et al. Catheterization and Cardiovascular Interventions 89:1250–1256 (2017)

Calcification is not a risk factor for patency loss or TLR

BARE METAL STENTS

Results of primary stent therapy for femoropopliteal peripheral arterial disease at 7 years



Konstantinos Stavroulakis, MD, Giovanni Torsello, MD, PhD, Ayad Manal, MD, Arne Schwandt, MD, Christiane Hericks, MA, Arne Stachmann, MA, Eva Schönefeld, MD, and Theodosios Bisdas, MD, PhD, Münster, Germany

ABSTRACT

Objective: Primary stenting is a well-established treatment option for femoropopliteal arterial disease. However, there is a paucity of data concerning the performance of this modality at ≥5 years. This study evaluated the long-term clinical and radiologic outcomes of primary stent therapy in patients with femoropopliteal arterial disease.

Methods: A prospective data collection and analysis was conducted in patients undergoing primary stent placement in femoropopliteal lesions between September 2006 and September 2007. The EverFlex (Medtronic/Covidien, Plymouth, Minn) bare-metal stent was used. The primary outcome of this study was the primary patency rate. Secondary outcomes were secondary patency rate, amputation-free survival (AFS), and freedom from target lesion revascularization (TLR). A Cox regression analysis identified risk factors for the primary and the secondary measure outcomes.

Results: Included were 89 patients (102 stents). The prevalence of critical limb ischemia was 34% (n = 30). The initial angiography revealed a TransAtlantic Inter-Society Consensus for the Management of Peripheral Arterial Disease C/D lesion in 31 patients (35%). Occlusions were present in 49 patients (55%) and the mean lesion length was 116 ± 33 mm. Popliteal artery disease was present in 39 treated limbs (35%). The primary patency rate at 1, 3, 5, and 7 years was 73%, 64%, 47%, and 33% respectively. At 7 years, secondary patency rate was 67%, freedom from TLR was 47%, and the AFS was 73%. Cox regression analysis revealed a decreased AFS among diabetic patients (hazard ratio [HR], 2.6; 95% confidence interval [CI], 1.08-6.28; P = .03), whereas secondary endovascular interventions showed a protective effect for AFS (HR, 0.14; 95% CI, 0.03-0.65; P = .01). Popliteal artery disease was identified as independent risk factor for secondary interventions (HR, 2.07; 95% CI, 1.05-4.06; P = .04) and TLR (HR, 1.99; 95% CI, 1.03-3.83; P = .04). Critical limb ischemia was associated with an increased incidence of surgical conversion owing to endovascular treatment failure (HR, 5.46; 95% CI, 2.44-12.17; P < .001).

Conclusions: This study found primary stenting was associated with acceptable clinical and radiologic long-term outcomes. Diabetes was associated with poor AFS, and popliteal artery involvement correlated with an increased need for reinterventions. AFS was better among patients undergoing secondary procedures. (J Vasc Surg 2016;64:696-702.)

Calcification did not affect primary patency rate

Table II. Preoperative angiographic findings with respect to Rutherford class

Angiographic findings ^a	Rutherford class			P value
	3 (IC)	4 (CLI)	5 (CLI)	
Total No.	58 (66)	14 (16)	16 (18)	
TASC II A/B lesions	42 (72)	9 (64)	7 (44)	.036
TASC II C/D lesions	16 (28)	5 (36)	9 (56)	
Occlusion	24 (41)	10 (71)	13 (81)	.002
Popliteal involvement	22 (38)	11 (79)	6 (38)	.019
P2/P3 involvement ^b	9 (16)	7 (50)	4 (25)	.021
Lesion length, mm	120 (100-140)	118 (100-140)	123 (80-220)	.965
Calcification	40 (69)	9 (64)	11 (69)	.943
Mild Ca ²⁺	13 (22)	3 (21)	4 (25)	.969
Moderate Ca ²⁺	18 (31)	4 (29)	4 (25)	.893
Severe Ca ²⁺	9 (16)	2 (15)	3 (19)	.950
Runoff vessels				
One	22 (38)	7 (50)	8 (50)	.554
Two	25 (43)	3 (21)	6 (38)	.325
Three	11 (19)	4 (29)	2 (12)	.535

CLI, Critical limb ischemia; IC, intermittent claudication; TASC II, TransAtlantic Inter-Society Consensus for the Management of Peripheral Arterial Disease.
^aCategorical data are presented as number (%) and continuous data as median (interquartile range).
^bP2 is the popliteal artery segment between the proximal part of patella and the center of knee joint space, and P3 is the segment between the knee joint space and the origin of the anterior tibial artery.

0.90-3.5; P = .09) affected PPR. The presence of distal popliteal artery disease (HR, 0.79; 95% CI, 0.26-2.41; P = .68), moderate or severe calcification (HR, 1.2; 95% CI, 0.6-2.45; P = .49), and lesion length >120 mm (HR, 1.4; 95% CI, 0.75-2.80; P = .26) did not affect PPR. The number of runoff

INTERWOVEN NITINOL STENTS

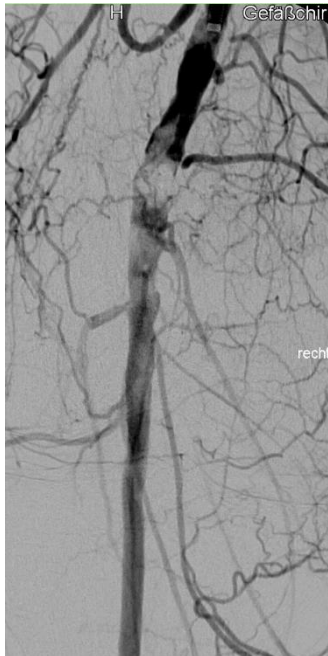


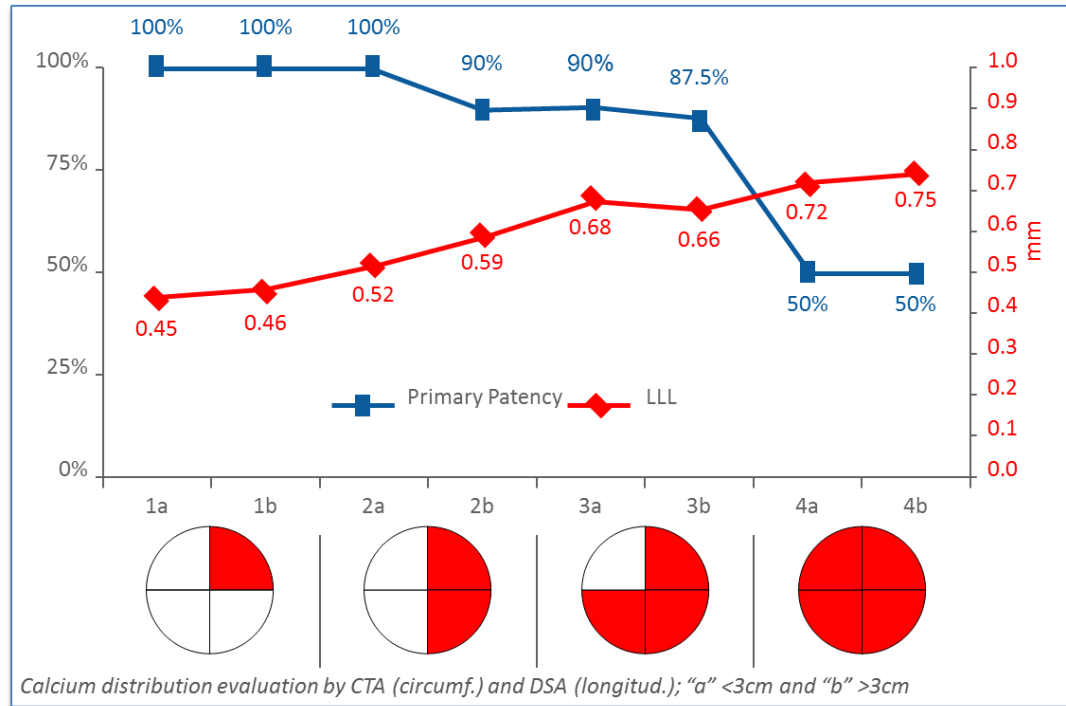
TABLE IIIa. Baseline Angiographic Findings—ITT, Single Stented Subjects

Lesion Characteristics	Nominal (-10% to 10%) (N of patients = 84, N of segments = 84)	Compressed (< -20%) (N of patients = 10, N of segments = 10)	Minimal compression (-20% to -10%) (N of patients = 23, N of segments = 23)	Minimal elongation (10-20%) (N of patients = 44, N of segments = 44)	Moderate elongation (20-40%) (N of patients = 44, N of segments = 44)	Severe Elongation (>40%) (N of patients = 31, N of segments = 31)
Lesion Length, mm Mean±SD (N)	76.62 ± 35.82 (84)	60.94 ± 27.12 (10)	73.98 ± 31.89 (23)	76.04 ± 36.56 (43)	69.41 ± 39.83 (44)	81.92 ± 53.13 (31)
Calcification, Any†						
Mild	21.4% (18/84)	20.0% (2/10)	17.4% (4/23)	25.0% (11/44)	34.1% (15/44)	35.5% (11/31)
Moderate	27.4% (23/84)	40.0% (4/10)	34.8% (8/23)	38.6% (17/44)	22.7% (10/44)	32.3% (10/31)
Severe	51.2% (43/84)	40.0% (4/10)	47.8% (11/23)	36.4% (16/44)	43.2% (19/44)	32.3% (10/31)
Total Occlusion (Per Patient)	29.8% (25/84)	30.0% (3/10)	17.4% (4/23)	25.0% (11/44)	20.5% (9/44)	19.4% (6/31)

Patients with calcification defined via angiographic core lab as severe with at least 1 cm of calcification noted on two sides of the artery (72.4% of the cohort) did not have a decrease in CD-TLR rates. At 3 years, freedom from CD-TLR was 87.6%.

Garcia et al, Catheterization and Cardiovascular Interventions 89:1259–1267 (2017)

DRUG-COATED BALLOONS IN CALCIFIED LESIONS



Fanelli et al. Cardiovasc Intervent Radiol. 2014 Aug;37(4):898-907

Clinical Investigation

JOURNAL OF
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THERAPY.

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www.jevt.org
SAGE

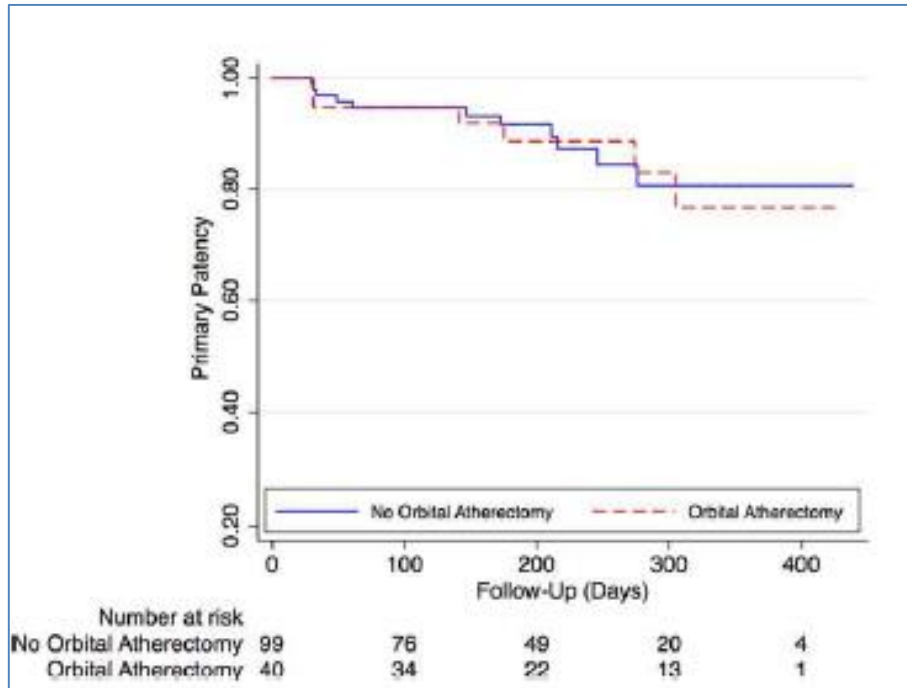
Drug-Eluting Balloon Therapy for Femoropopliteal Occlusive Disease: Predictors of Outcome With a Special Emphasis on Calcium

Gunnar Tepe, MD¹, Ulrich Beschorner, MD², Charlotte Ruether, MD¹, Imma Fischer, PhD³, Peter Pfaffinger, MD¹, Elias Noory, MD², and Thomas Zeller, MD²

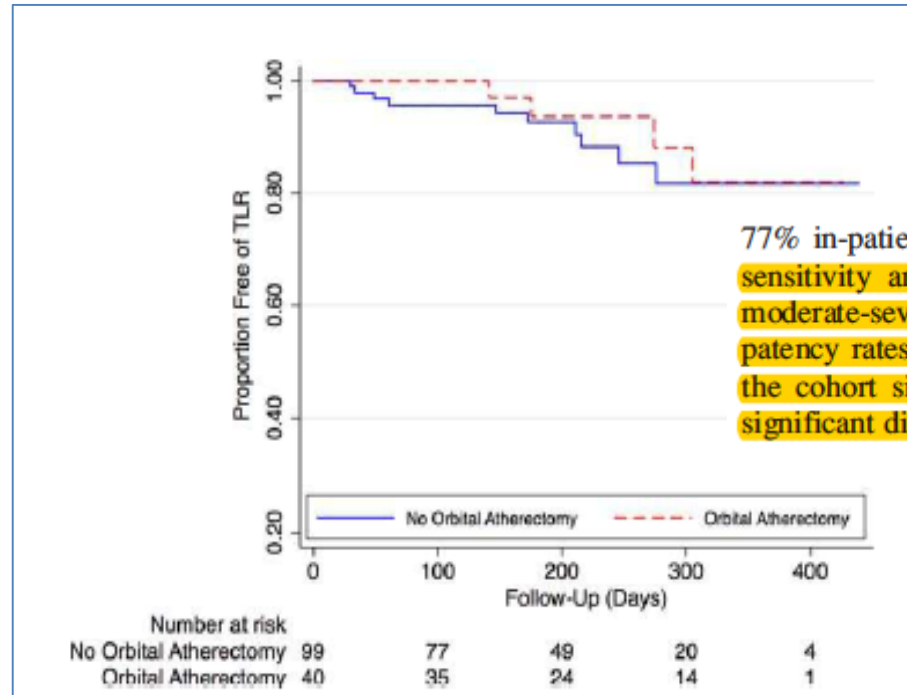
- Retrospective analysis of 91 patients²
- Analysed at 6M post DEB
- Lesion calcification analysed by core labs (PACSS score + angiographic calcium score)
- Severity of lesion calcification is associated with LLL after treatment with DCB.
- **Author conclusion: "One possible approach to overcome this limitation might be plaque modification or removal prior to DEB usage"**

ORBITAL ATHERECTOMY + DCB VS DCB

Freedom from TLR



12 M Primary Patency



Foley et al. Catheterization and Cardiovascular Interventions 89:1078–1085 (2017)

DIRECTIONAL ATHERECTOMY + DCB (DAART)

Severely calcified lesions

Mean lesion length: 11.5 cm

CTOs: 13 %

Bailout Stent: 6.7 %

12 M PPR: 90 %

12 M TLR: 10 %

Amputation rate: 0 %

Cardiovascular Revascularization Medicine 13 (2012) 219–223

Contents lists available at ScienceDirect

ELSEVIER Cardiovascular Revascularization Medicine

Combined treatment of heavy calcified femoro-popliteal lesions using directional atherectomy and a paclitaxel coated balloon: One-year single centre clinical results[☆]

Angelo Cioppa^{*}, Eugenio Stabile, Grigore Popusoi, Luigi Salemme, Linda Cota, Armando Pucciarelli, Vittorio Ambrosini, Giovanni Sorropago, Tullio Tesorio, Alessia Agresta, Giancarlo Biamino, Paolo Rubino

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Atherectomy
Drug coated balloons

ABSTRACT

Background: The use of Directional Atherectomy (DA) for the treatment of calcified femoro-popliteal lesions seems to improve the acute procedural success, however without reducing the long term restenosis rate. Drug coated balloons (DCB) reduced restenosis rate in non heavy calcified lesions. Aim of this study was to demonstrate safety and efficacy of a combined endovascular approach using DA and DCB for the treatment of heavy calcified lesions of the femoro-popliteal tract.

Methods: From January 2010 to November 2010, 240 patients underwent PTA of the femoro-popliteal tract in our institution. Within this cohort a total of 30 patients had Life Limiting Claudication (LLC) (n=18) and 12 a Critical Limb Ischemia (CLI) with baseline Rutherford class 4.2 ± 1.2 underwent PTA of heavy calcified lesions with intravascular ultrasound guided DA and DCB. All procedures have been performed using a distal protection device. Stent implantation was allowed only in case of flow limiting dissections or suboptimal result (residual stenosis >50%) by visual estimation. After the intervention patients were followed up to 12 months.

Results: Procedural and clinical success, was achieved in all cases. Bail-out stenting was necessary in only two (6.7%). At twelve month follow up median Rutherford class was 2.2 ± 1.2, ABI was 0.8 ± 0.1 and limb salvage rate was 100%. Two minor foot finger or forefoot amputations, were performed to reach complete wound healing and/or preserve deambulation. Duplex control was performed in all the cases (n=30). In three cases duplex scan showed a significant target lesion restenosis requiring a reintervention (TLR=10%) leading a total one-year secondary patency rate of 100%. All the three restenosed patients were insulin dependent diabetics and none of them were started during the procedure.

Conclusion: The data suggest that combined use of DA and DCB may represent a potential alternative strategy for the treatment of femoro-popliteal severely calcified lesions. These very promising data and the considered hypothesis have to be confirmed in a multicentre randomized trial.

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1. Introduction

Restenosis rate after percutaneous transluminal angioplasty (PTA) of calcified superficial femoral and popliteal artery is still the "Achilles' Hill" of this endovascular procedure [1,2].

One of the most important limiting factors in terms of acute procedural success rate and need of stent implantation, as well as long term patency seems to be the presence of diffuse and severe calcification of the target lesion.

Moreover the majority of randomised controlled studies considered severe calcification of the lesion as an exclusion criterion. The presence of severe calcification of the atherosclerotic lesion is responsible for a poor response to balloon dilation, due to significant acute vessel recoil and frequent flow limiting dissections. Directional atherectomy (DA) improves acute success by debulking fibrocalcific portion of the atherosclerotic plaque [3], but does not provide any benefits in terms of patency rate [4–9].

The use of drug coated balloons (DCB) for the endovascular treatment of the femoro-popliteal tract has been demonstrated to reduce the occurrence of restenosis [10–13]. There are no data available, at the moment, regarding the use of DCB in heavy calcified lesions.

In this registry we applied a new endovascular approach for the treatment of severely calcified lesions in the femoro-popliteal tract, based on the combination of DA followed by DCB, to prevent long-term restenosis and to reduce the need of stenting.

2. Methods

2.1. Study design

This is a single centre, single arm, prospective non randomised study.

[☆] Conflict of Interest: None.
^{*} Corresponding author. Tel.: +39 0825 680022; fax: +39 0825 680022.
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DEFINITIVE AR TRIAL

Baseline Lesion Characteristics

Per Core Lab

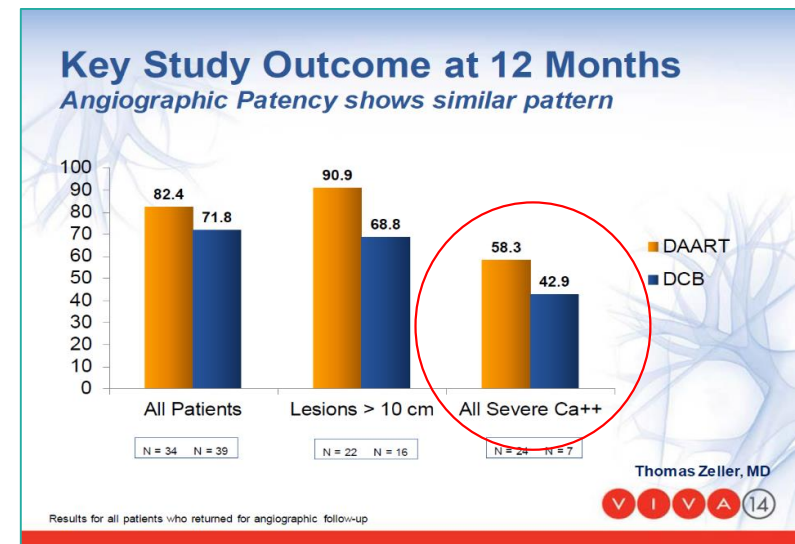
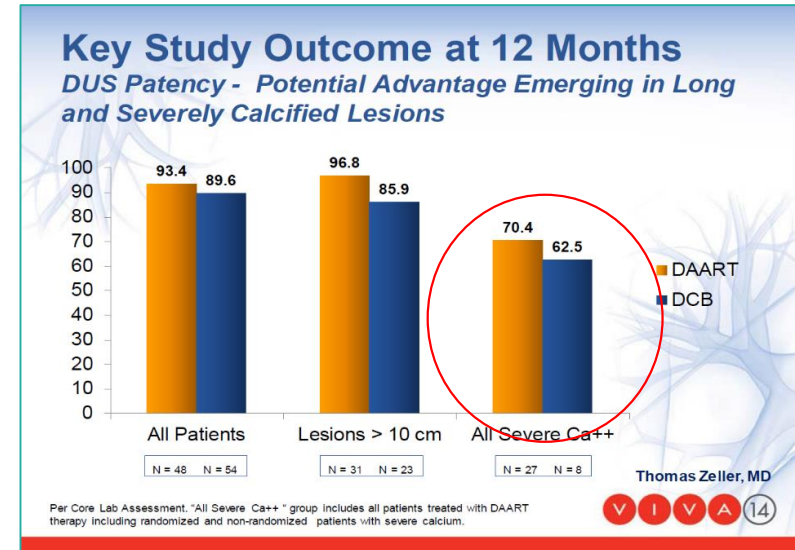
Baseline Characteristics	DAART (N= 48)	DCB (N = 54)	p-Value*	DAART Severe Ca++ Arm (N=19)
Lesion Length (cm)	11.2	9.7	0.05	11.9
Diameter Stenosis	82%	85%	0.35	88%
Reference vessel diameter (mm)	4.9	4.9	0.48	5.1
Minimum lumen diameter (mm)	1.0	0.8	0.34	0.7
Calcification	70.8%	74.1%	0.82	94.7%
Severe calcification	25.0%	18.5%	0.48	89.5%

Thomas Zeller, MD

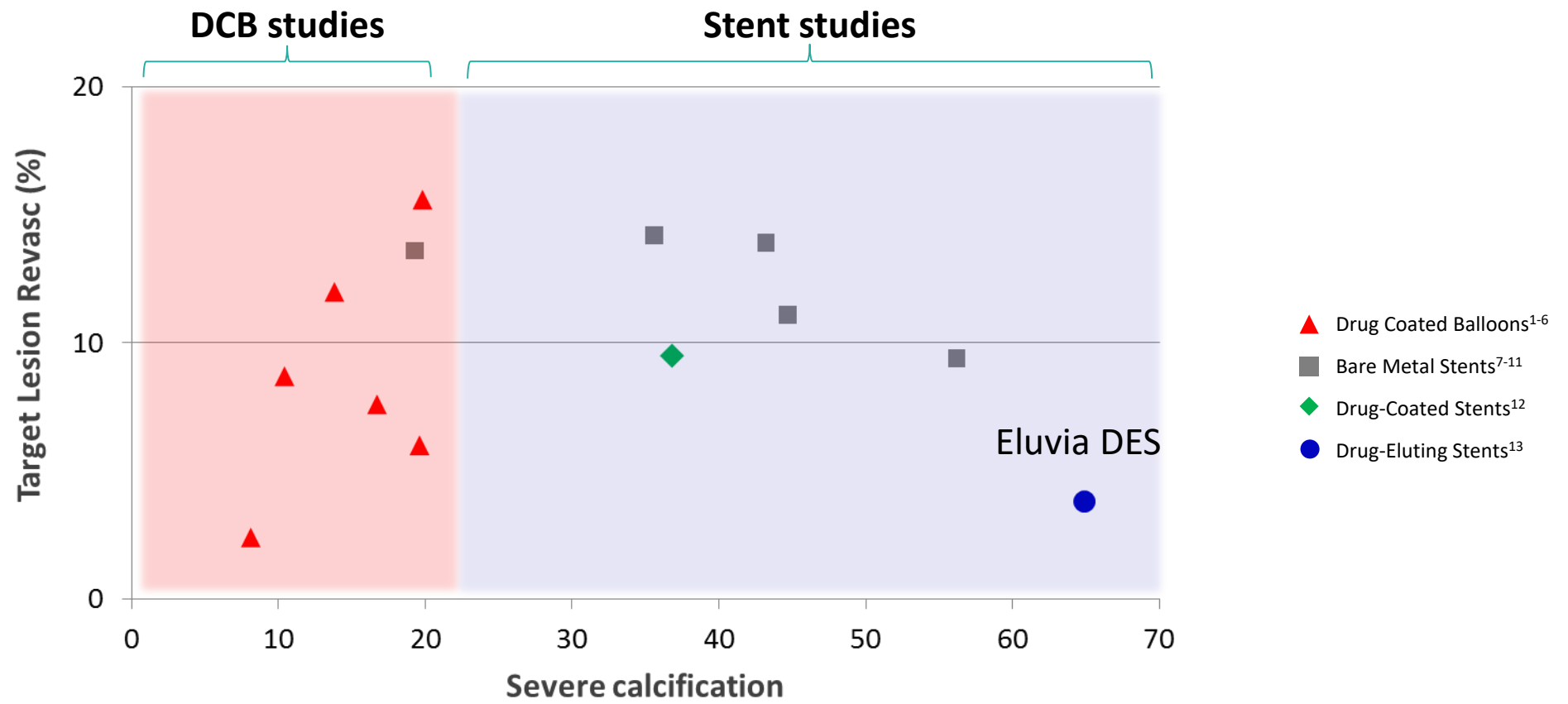
* p-value for DAART and DCB groups

V I V A 14

Zeller et al. Circ Cardiovasc Interv 2017 Sep;10(9). pii: e004848



DRUG-ELUTING STENTS



¹Micari A Et al. J Am Coll Cardiol Interv 2012; ²Tepe G et al. Circulation 2015; ³Zeller T et al. J Endovasc Therapy 2014; ⁴Schroeder H et al. Catheter Cardiovasc Interv 2015; ⁵Laird J. Endovascular Today Feb 2015; ⁶Ansel G. TCT 2015; ⁷Matsumura et al. J of Vasc Surg. Jul 2013; ⁸⁻⁹www.accessdata.fda.gov; ¹⁰www.endovascularmagazine.eu 2013; ¹¹Powell, R. Charing Cross 2015; ¹²Dake MD et al. Circ Cardiovasc Interv 2011; ¹³Müller-Hülsbeck, S. VIVA 2015.

‘PAVE AND CRACK’ TECHNIQUE

Clinical Investigation

Endovascular Treatment of Severely Calcified Femoropopliteal Lesions Using the “Pave-and-Crack” Technique: Technical Description and 12-Month Results

Journal of Endovascular Therapy
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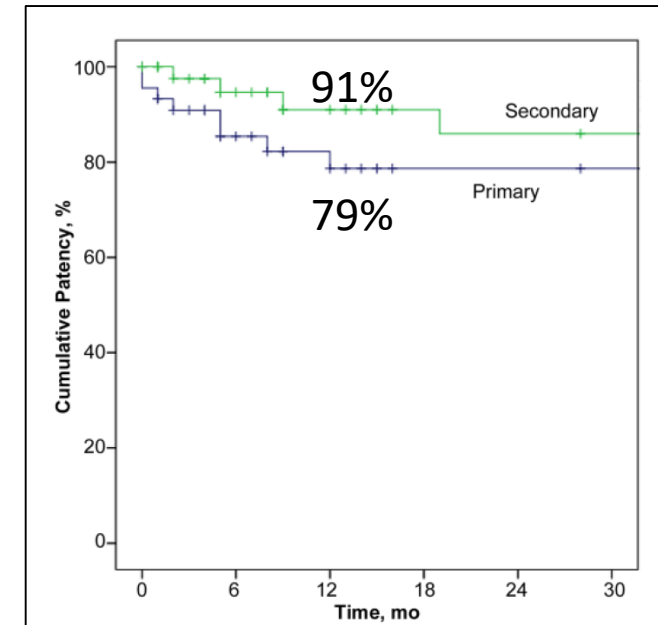
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Calcification (PACSS)	
1	3/64 (5)
2	2/64 (3)
3	19/64 (30)
4	40/64 (62)
Severe calcification ^b	40/66 (61)

Use of high pressure or cutting balloon	23/65 (35)
Perforation	39/66 (59)
Viabahn cumulative length, cm	16.5±8.7 (n=51)
Viabahn maximum diameter, mm	
5	2/67 (3)
6	34/67 (51)
7	28/67 (42)
8	3/67 (5)
SUPERA cumulative length, cm	22.9±12.0 (n=51)
SUPERA maximum diameter, mm	
5	32/67 (48)
6	34/67 (51)
7	1/67 (2)

Technique:

1. Local anesthesia in the surrounding issues of the SFA lesion
2. PTA
3. Angiographic control and confirmation of the residual stenosis
4. Deployment of a Viabahn endoprosthesis and aggressive dilatation with a high-pressure balloon catheter
5. Angiographic control and relining with Supera stents



MY TECHNICAL CHOICE IN SEVERE CALCIUM IN THE SFA

In fit patients:

LEAVE NOTHING BEHIND - STRATEGY

- **DAART + DCB**
 - LONG LESIONS
 - NORMAL RENAL FUNCTION
 - CLAUDICANTS
- **LITHOPLASTY + DCB**
 - SHORT LESIONS
 - CHRONIC KIDNEY DISEASE
 - CLI + CLAUDICANTS
- **IF STENT NECESSARY, THEN:**
 - DRUG-ELUTING STENT

In multimorbid patients:

LEAVE DIRECTLY A STENT BEHIND

- **PTA ± SCORING BALLOON**
 - **RECOIL < 30% ?**
BARE METAL or DRUG-ELUTING STENTS
 - **RECOIL < 30% in the popliteal artery**
SUPERA STENT
 - **RECOIL > 30%**
VIABAHN + HIGH-PRESSURE BALLOON CATHETER ± SUPERA

QUESTIONS TO VASCUPEDIANS

- What is your technical choice in severely calcified lesions?
- Which of the new technologies are available in your cath lab?
- Do you use DCBs in calcified lesions?
- Which kind of study would be helpful in your daily practice regarding calcified lesions in the SFA?