

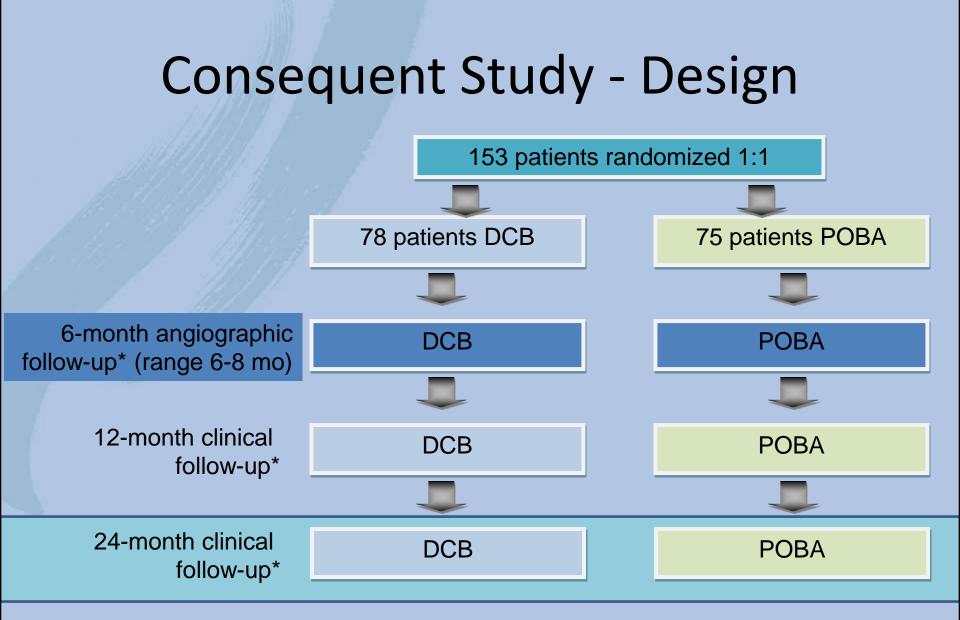
## MAKE COMBINATION THERAPY GREAT AGAIN DCBS AND AND SPOT STENTS IN FEMPOP LESIONS

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### Disclosure

Speaker name: Gunnar Tepe I have the following potential conflicts of interest to report: Study support and Advisory Board BBraun

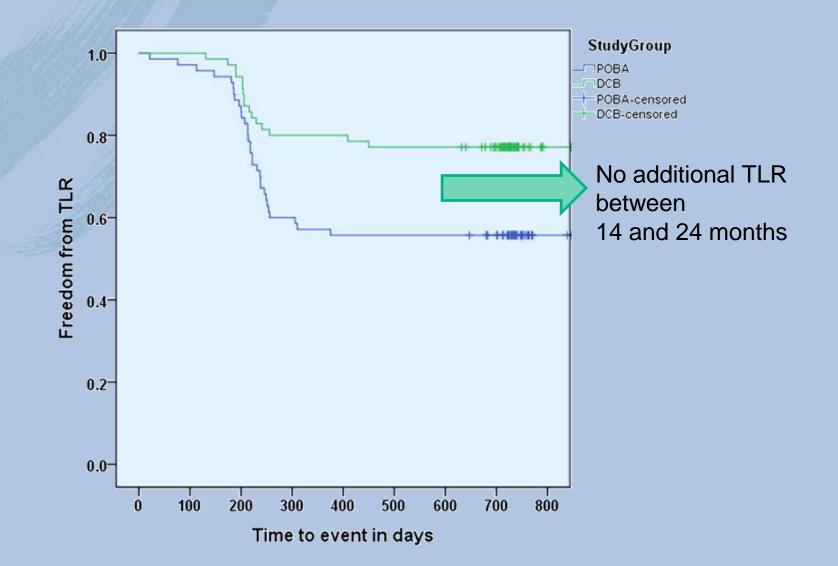


including walking test, ABI and Duplex

## Lesion details – target lesions

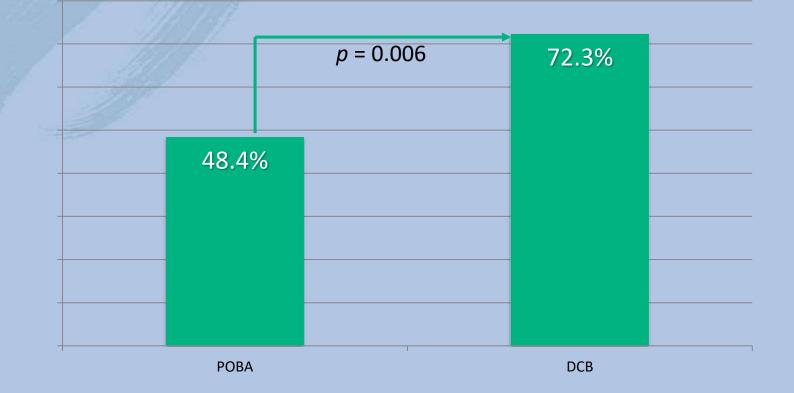
	All patients	Drug Coated Balloon	Uncoated Balloon	p-value
Target lesions	153	78	75	-
Location SFA P1/P2 SFA + P1/P2	122 (79.7%) 9 (5.9%) 22 (14.4%)	63 (80.8%) 4 (5.1%) 11 (14.1%)	59 (78.7%) 5 (6.7%) 11 (14.7%)	0.912
TASC A TASC B TASC C TASC D	54 (35.3%) 63 (41.2%) 26 (17.0%) 10 (6.5%)	28 (35.9%) 31 (39.7%) 13 (16.7%) 6 (7.7%)	26 (34.7%) 32 (42.7%) 13 (17.3%) 4 (5.3%)	0.934
Diameter stenosis, %	76.6 ± 18.1	76.0 ± 17.7	77.1 ± 18.5	0.703
Total occlusions	40 (26.1%)	18 (23.1%)	22 (29.3%)	0.462
Lesion length, cm	13.2 ± 10.4	13.7 ± 12.2	12.6 ± 8.2	0.540
Reference diameter, mm	5.22 ± 0.87	5.06 ± 0.77	5.38 ± 0.94	0.050
2 <sup>nd</sup> non-target lesion	18 (11.8%)	9 (11.5%)	9 (12.0%)	0.929

## 24-month Kaplan-Meier Curve



## 24-month patency

CONSEQUENT trial: 24 month Patency

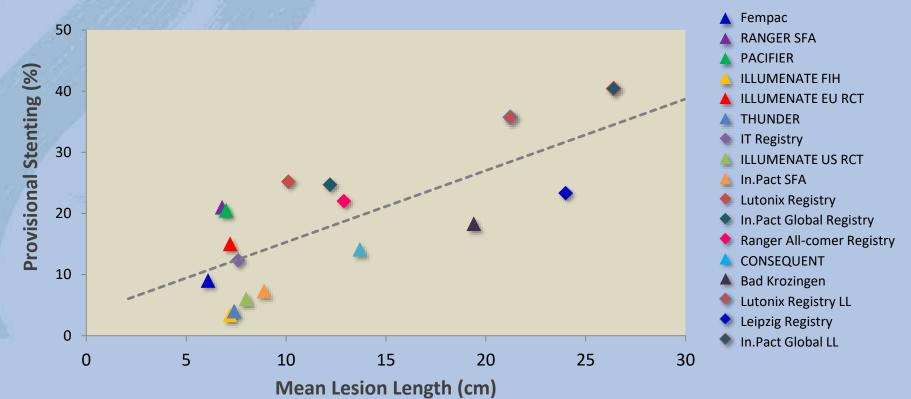


Patency defined as binary restenosis with diameter stenosis >50% (angiographic) or PSVR>2.4 (sonographic), definition by Diehm et al. [8]

#### Stents + DCBs



• Longer mean lesion length correlates with higher provisional stenting rate



Provisional Stenting in Randomized Controlled Trials may not be representative of actual stenting in studies due to study design

FEMPAC- Werk M et al. Circulation 2008 RANGER SFA-Bausback et al. J Endovasc Ther 2017 PACIFIER- Werk et al. Circ Cardiovasc Interv 2012 THUNDER- Tepe G et al. N Engl J Med 2008 IT Registry- Micari A Et al. J Am Coll Cardiol Intv 2012 IN.PACT SFA- Tepe et al. Circulation 2015

Lutonix Registry- Thieme M, et al. JACC Cardiovasc Interv. 2017 Interv 2015 CONSEQUENT- Tepe et al. Cardiovasc Intervent Radiol 2017 Bad Krozingen- Zeller T et al. J Endovasc Therapy 2014; Leipzig Registry- Schmidt A, et al. JACC Cardiovasc Interv. 2016

Results from different trials are not directly comparable. Information provided for educational purposes.

> ILLUMENATE FIH- Schroeder H et al. Catheter Cardiovasc 7Interv 2015

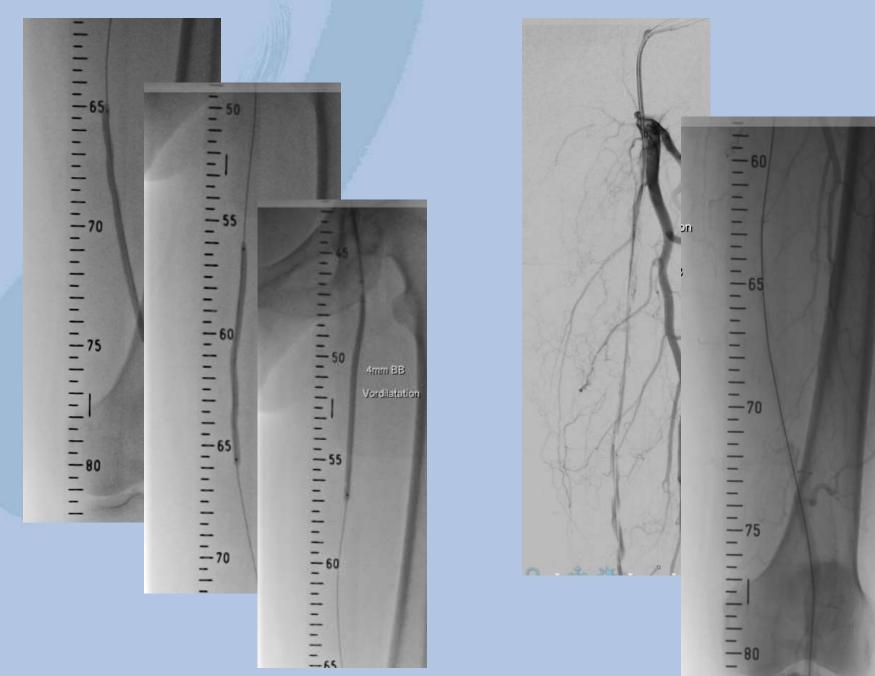
ILLUMENATE EU RCT- Schroeder et al, Circulation 2017 ILLUMENATE US RCT- Krishnan et al, Circulation 2017 In.PACT Global Registry- Ansel G. TCT 2015 Ranger All-Comer Registry- Lichtenberg, M. CIRSE 2017



# CASE 1



Results from case studies are not necessarily predictive of results in other cases. Results in other cases may vary. CAUTION: The law restricts these devices to sale by or on the order of a physician. Rx Only.



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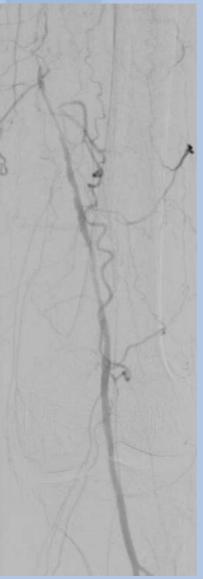


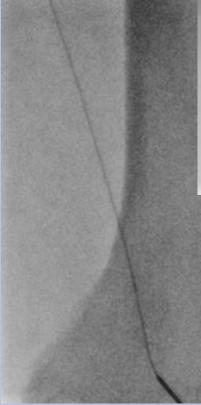
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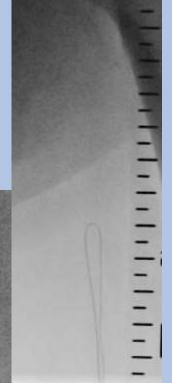


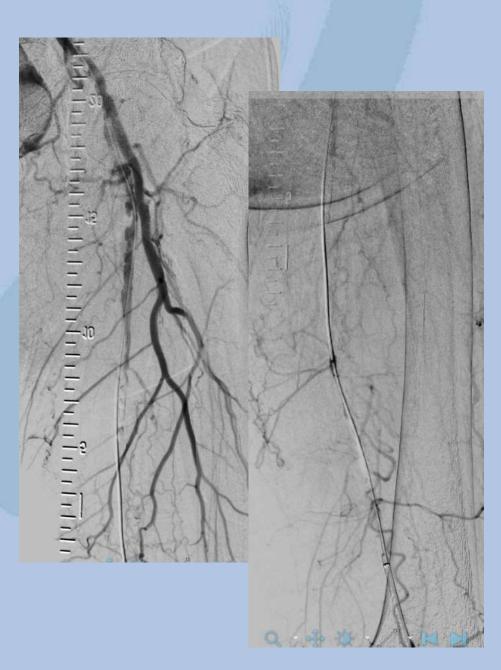
# CASE 2



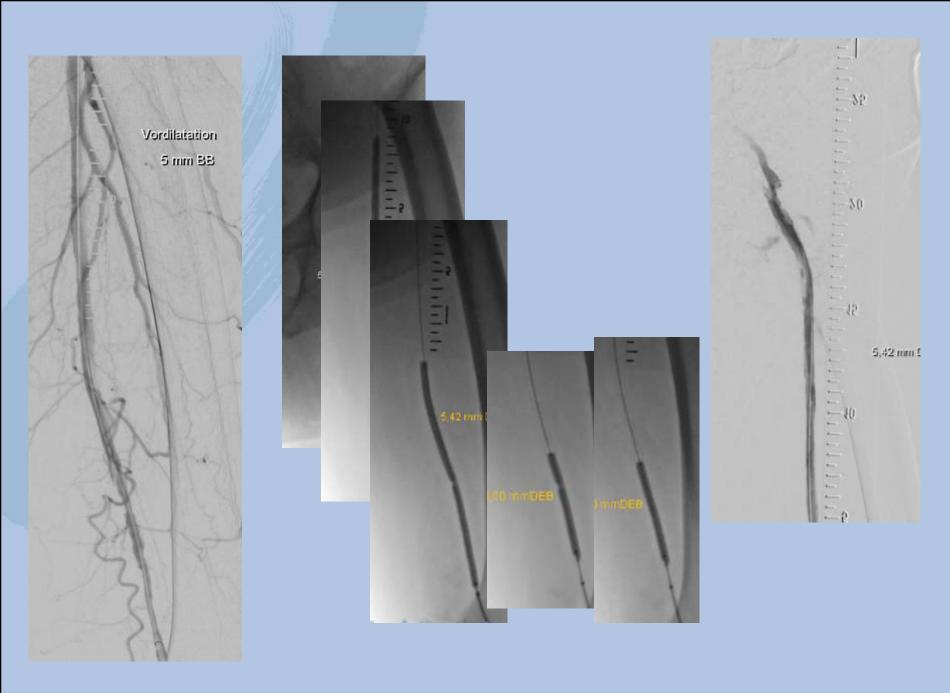


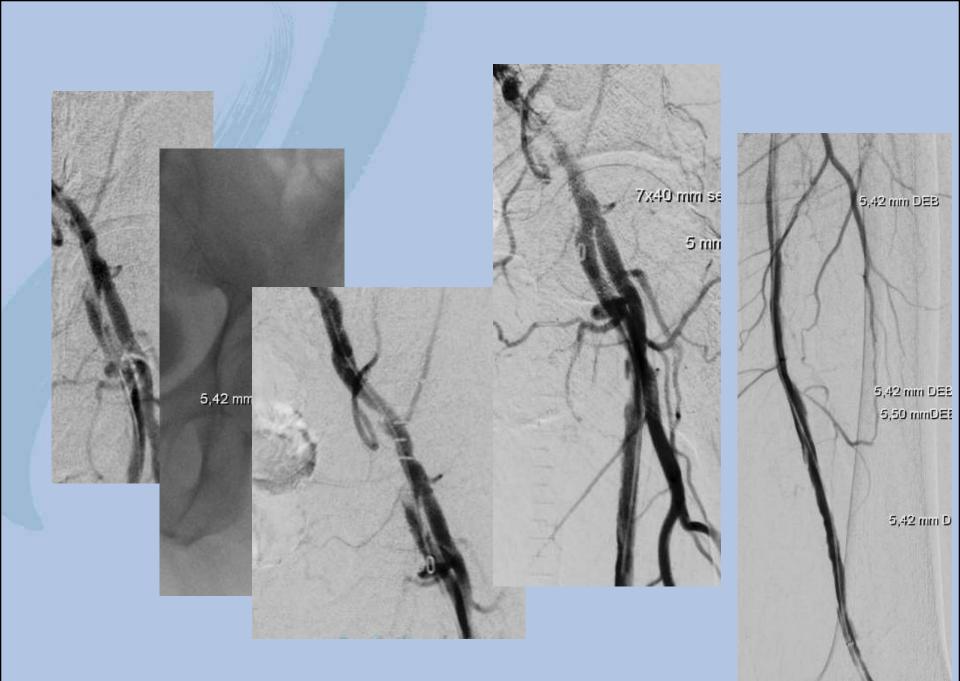


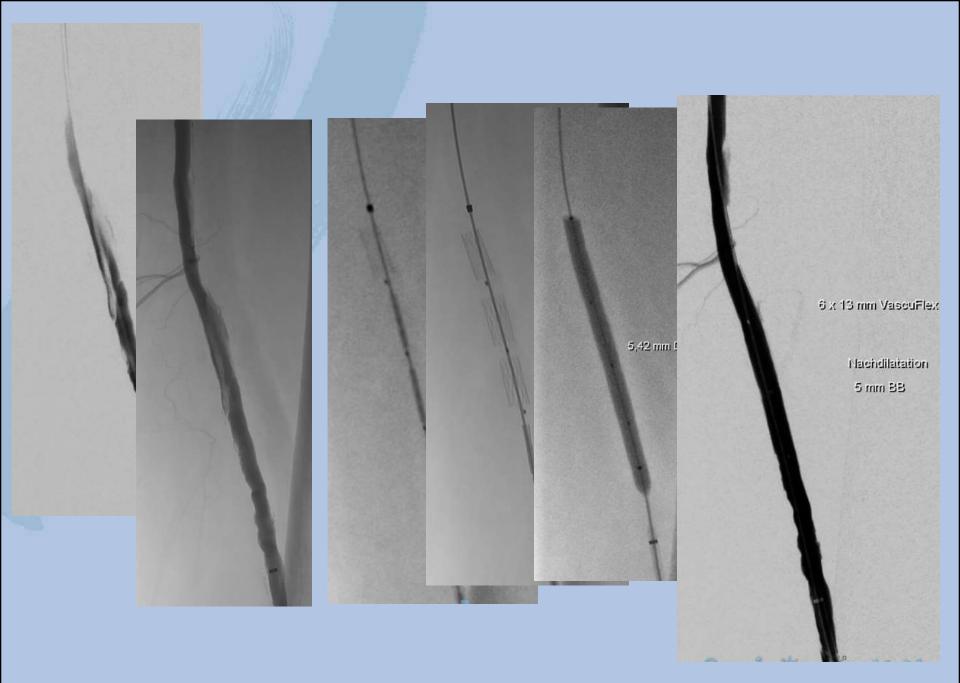




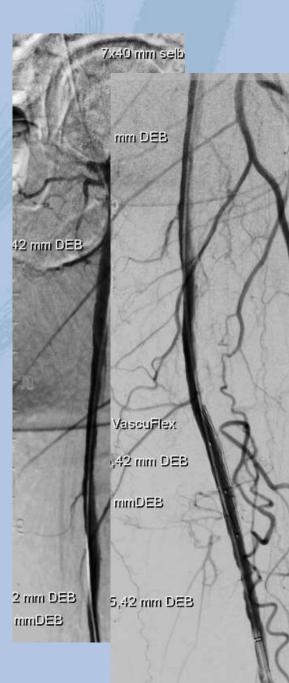


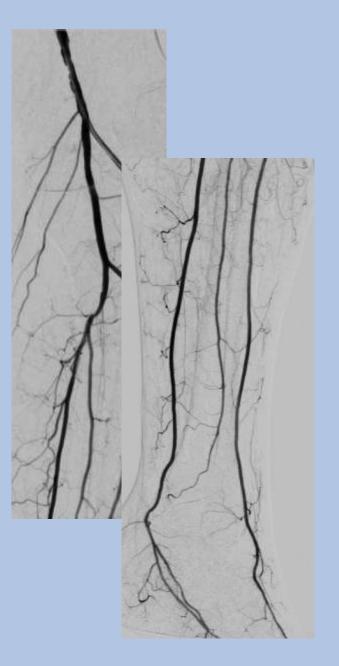












### **CTOs: Data from In.Pact Global**



Safety outcome*	CTO Imaging Cohort N=115 subjects†	Non- Stented N=63 subjects	Stented N=52 subjects
Clinically-driven TLR <sub>e</sub> t n (%)	13 <mark>(</mark> 11.3)	9 (14.3)	4 (7.7)
Thrombosis, n (%)	5 (4.3)	5 (7.9)	0 (0.0)

#### Conclusion



- DCB only = good solution, if possible
- DCB = no weapon which solves everything
- Especially in CTOs do not leave relevant rest stenosis and hope that the DCB does the rest
- DCB + (Spot)Stent = very good option



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