

14th



CABG vs PCI in Multivessel disease

Balbir Singh, INDIA

Speaker's name : Balbir, SINGH, Gurgaon

I do not have any potential conflict of interest

Advances in CABG and PCI

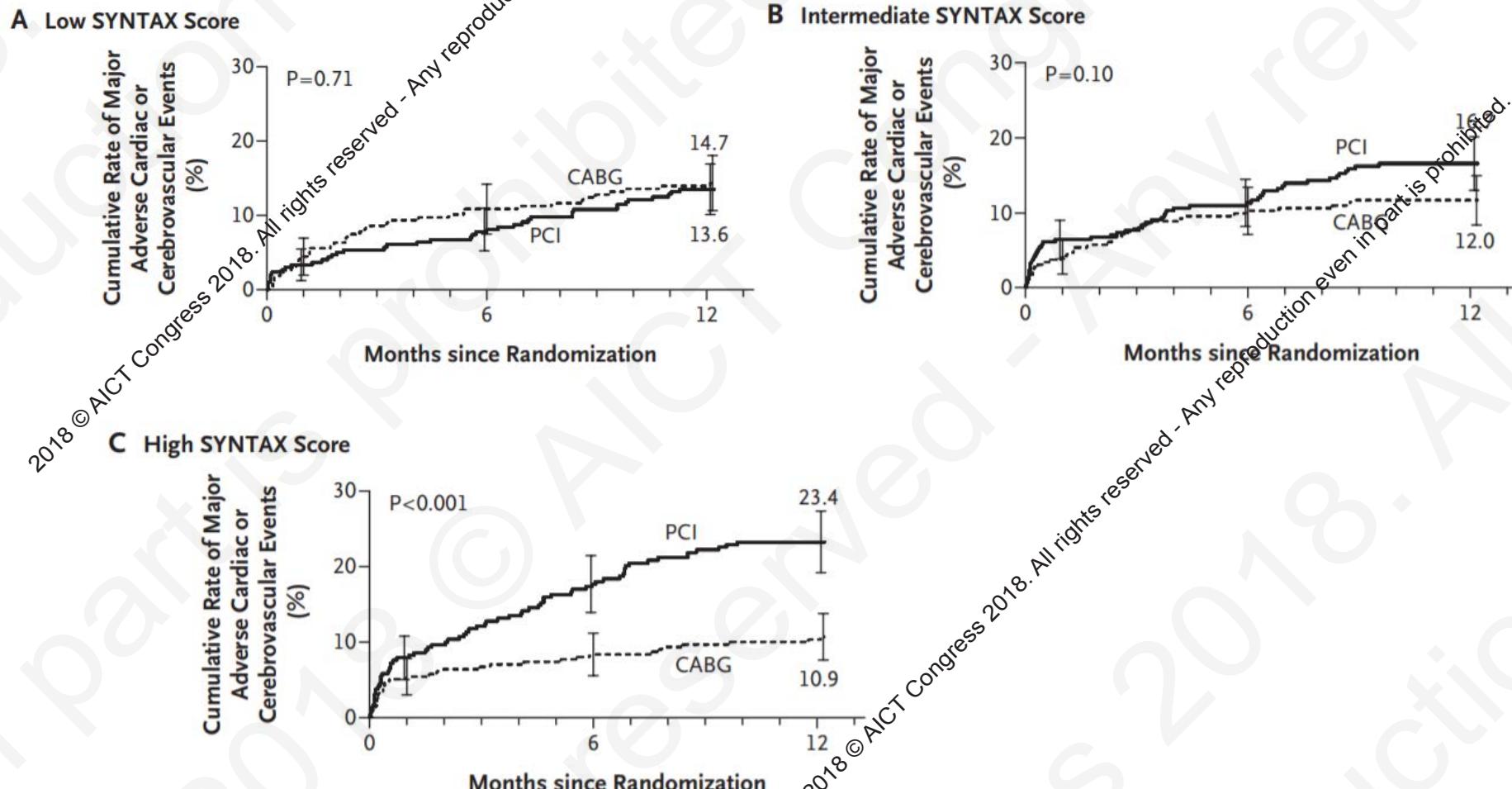
CABG

- First surgery 1960
- LIMA graft
- Complete revascularisation
- Off pump surgery
- Ventricular reconstruction surgery

PCI

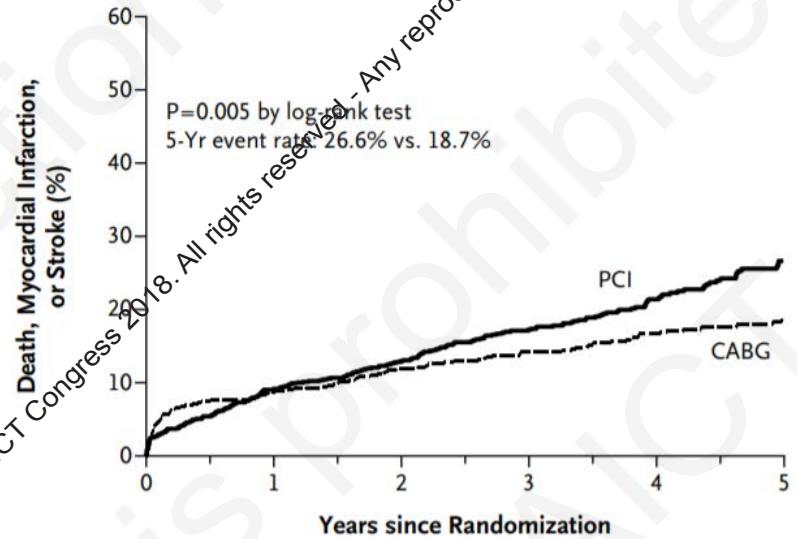
- Balloon angioplasty 1977
- Stent(BMS)
- DES
- 2 nd generation DES
- Intravascular imaging
- FFR

Percutaneous Coronary Intervention versus Coronary-Artery Bypass Grafting for Severe Coronary Artery Disease

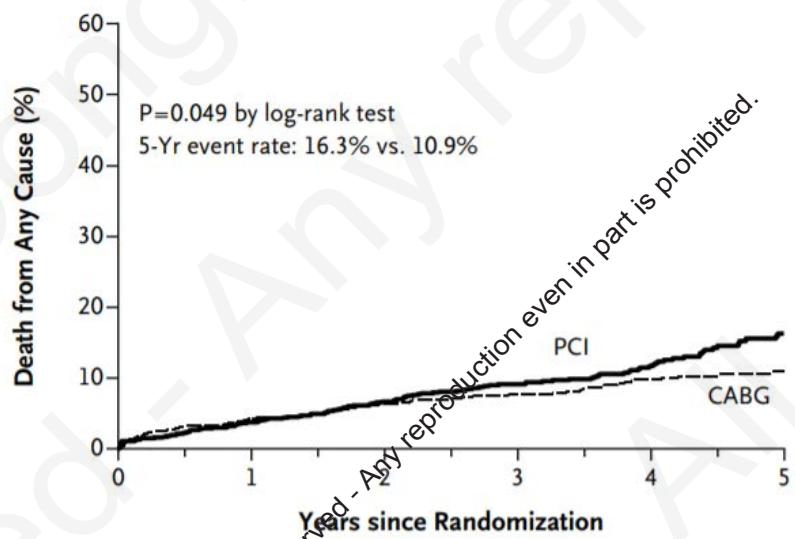


Strategies for Multivessel Revascularization in Patients with Diabetes

A Primary Outcome



B Death



No. at Risk

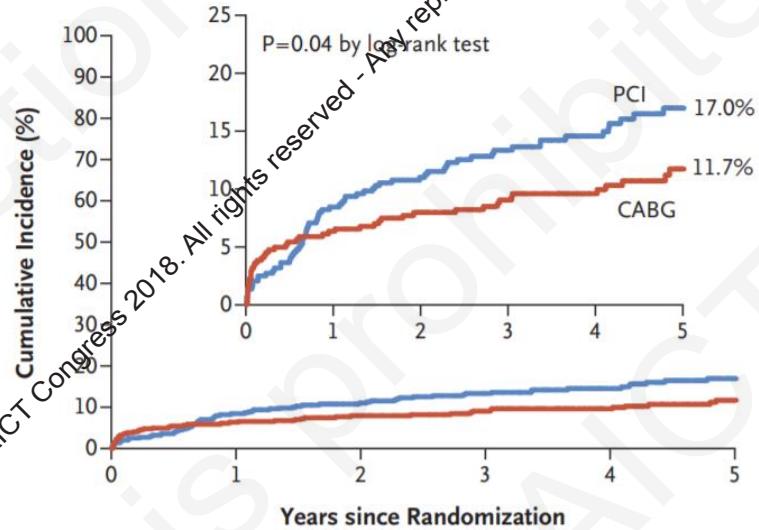
PCI	953	848	788	625	416	219
CABG	947	814	758	613	422	221

No. at Risk

PCI	953	845	785	685	466	243
CABG	947	806	655	449	238	

Trial of Everolimus-Eluting Stents or Bypass Surgery for Coronary Disease

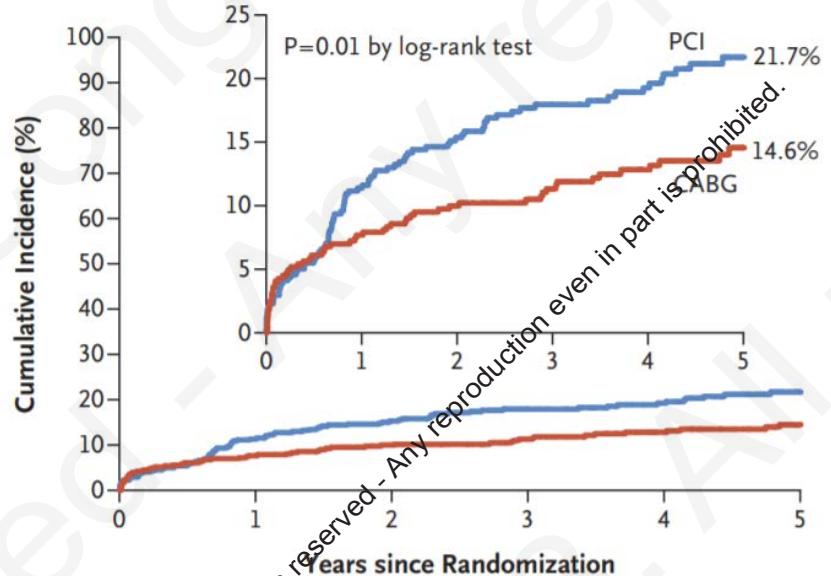
A Primary Composite End Point



No. at Risk

PCI	438	402	362	305	242	126
CABG	442	415	377	326	262	145

B Death, Myocardial Infarction, Stroke, or Repeat Revascularization



No. at Risk

PCI	438	389	341	288	229	117
CABG	442	409	368	317	250	137

BEST TRIAL

Park et al n engl j med 372;13, 2015

Recommendation for the type of revascularization (CABG or PCI) in patients with SCAD with suitable coronary anatomy for both procedures and low predicted surgical mortality.

Recommendations according to extent of CAD	CABG		PCI		Ref ^c
	Class ^a	Level ^b	Class ^a	Level ^b	
One or two-vessel disease without proximal LAD stenosis.	IIb	C	I	C	
One-vessel disease with proximal LAD stenosis.	I	A	I	A	107,108,160, 161,178,179
Two-vessel disease with proximal LAD stenosis.	I	B	I	C	108,135,137
Left main disease with a SYNTAX score ≤ 22.	I	B	I	B	17,134,170
Left main disease with a SYNTAX score 23–32.	I	B	IIa	B	17
Left main disease with a SYNTAX score >32.	I	B	III	B	17
Three-vessel disease with a SYNTAX score ≤ 22.	I	A	I	B	17,157,175,176
Three-vessel disease with a SYNTAX score 23–32.	I	A	III	B	17,157,175,176
Three-vessel disease with a SYNTAX score >32.	I	A	III	B	17,157,175,176

CABG: coronary artery bypass grafting; LAD: left anterior descending coronary artery; PCI: percutaneous coronary intervention; SCAD: stable coronary artery disease. ^aClass of recommendation. ^bLevel of evidence. ^cReferences.

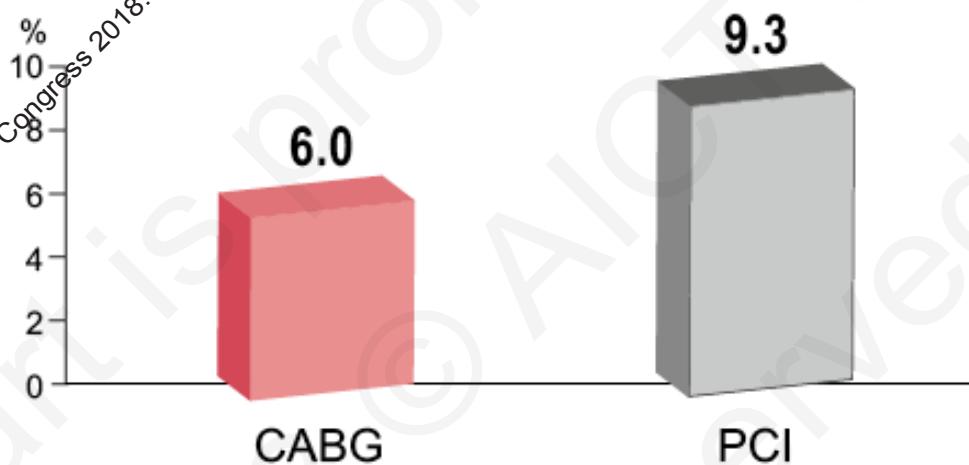
Patients
1,275

CABG 638

Median follow-up
61 months

PCI 637

Death from **any cause** (median 61 months)
HR 0.65 (95% CI 0.43-0.98)
 $p=0.039$



**Long-term death after revascularisation non-diabetic patients with MV-disease:
SYNTAX / BEST meta-analysis** Chang et al. J Am Coll Cardiol 2016;68:29-36

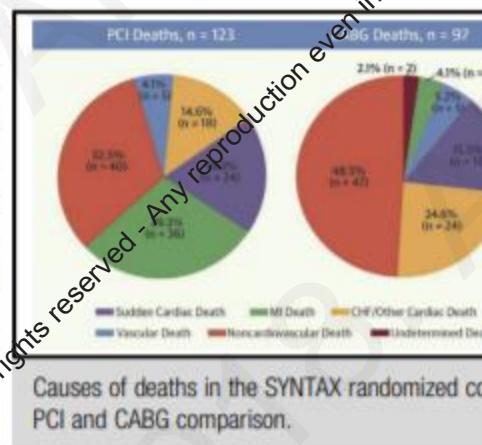
FEATURED EXPERT OPINION: ACQUIRED: VALVE

Why surgery won the SYNTAX trial and why it matters

Michael Mack, MD, Heike Baumgarten, MD, and Bruce Lytle, MD

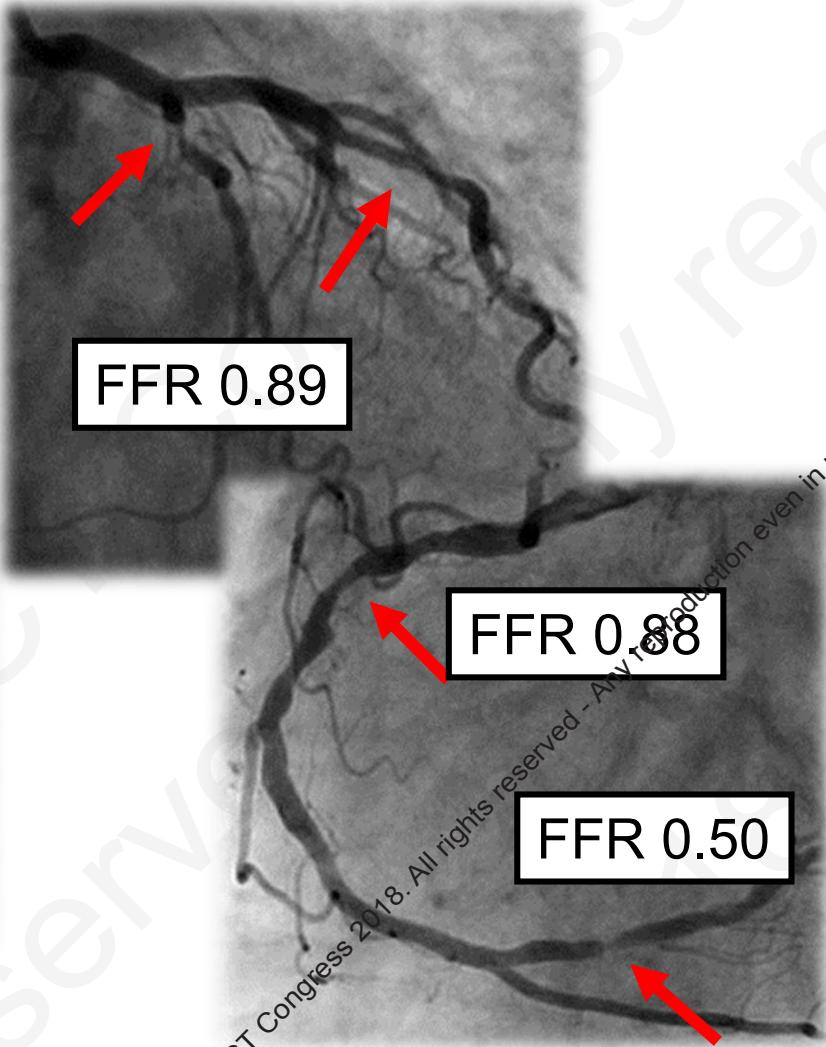
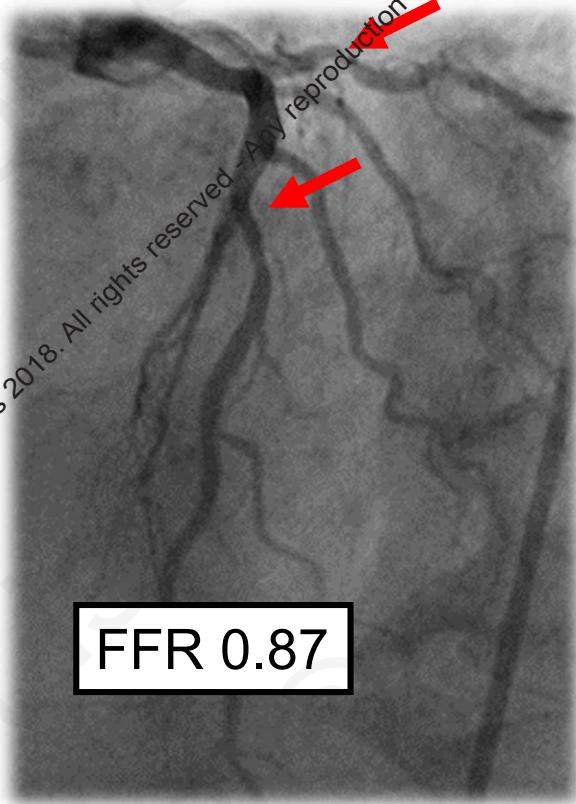
ABSTRACT

The Synergy Between Percutaneous Coronary Intervention With Taxus and Cardiac Surgery (SYNTAX) trial cause of death analysis shows that cardiac death due to myocardial infarction is 10 times higher with percutaneous coronary intervention than coronary artery bypass grafting in the higher-risk patients. There was a clear advantage for surgery in the prevention of death in both the intermediate and high SYNTAX score groups with 3-vessel disease and in the high SYNTAX score group with left main disease, and that incremental advantage is statistically significant and widening with time, which should be transparently communicated to patients. (J Thorac Cardiovasc Surg 2016;152:1237-40)



State of art PCI

- Physiology
- Imaging
- Bifurcation lesions (using the EBC guidelines)
- CTO management
- Management of calcified lesions
- Quality of stents



FFR in stable lesions

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FAME 2

ORIGINAL ARTICLE

Fractional Flow Reserve–Guided PCI for Stable Coronary Artery Disease

Benjamin Dreyer, M.D., Ph.D., William F. Fearon, M.D., Nico H.J. Pijls, M.D., Ph.D., François Bertrand, M.D., Ph.D., Rôm Tanami, M.D., Ph.D., Zsolt Piroth, M.D., Bojan Jelic, M.D., Sven Mathias Winckler, M.D., Gilles Rioufol, M.D., Ph.D., Alis Witt, M.D., Ph.D., Peter Kala, M.D., Philip MacCarthy, M.D., Thomas Engstrom, M.D., Keith Oldroyd, M.D., Kostas Mamoulakis, M.D., Gautham Manoharan, M.D., Peter Virdee, M.D., Ole Fløbert, M.D., Nick Curwin, B.M., Ph.D., Jane B. Johnson, R.N., B.S.N., Andreas Limacher, Ph.D., Esther Nienhuijs, Ph.D., and Peter Jüni, M.D., for the FAME 2 Trial Investigators*

Compare-Acute

ORIGINAL ARTICLE

Fractional Flow Reserve–Guided Multivessel Angioplasty in Myocardial Infarction

Pieter C. Smits, M.D., Ph.D., Mohamed Abdel-Wahab, M.D., Frans-Joost Verheyen, M.D., Bianca M. Boxma-de Klerk, Ph.D., Ketil Lunde, M.D., Carl E. Selsværgh, M.D., Zsolt Piroth, M.D., David Horak, M.D., Adrian Włodarczak, M.D., Paul J. Ong, M.D., Rainer Hambrecht, M.D., Oskar Angeris, M.D., Gert R. Schmidt, M.D., Ph.D., and Elmir Omerovic, M.D., for the Compare-Acute Investigators*

DANAMI-3-PRIMULTI

Complete revascularisation versus treatment of the culprit lesion only in patients with ST-segment elevation myocardial infarction and multivessel disease (DANAMI-3—PRIMULTI) an open-label, randomised controlled trial

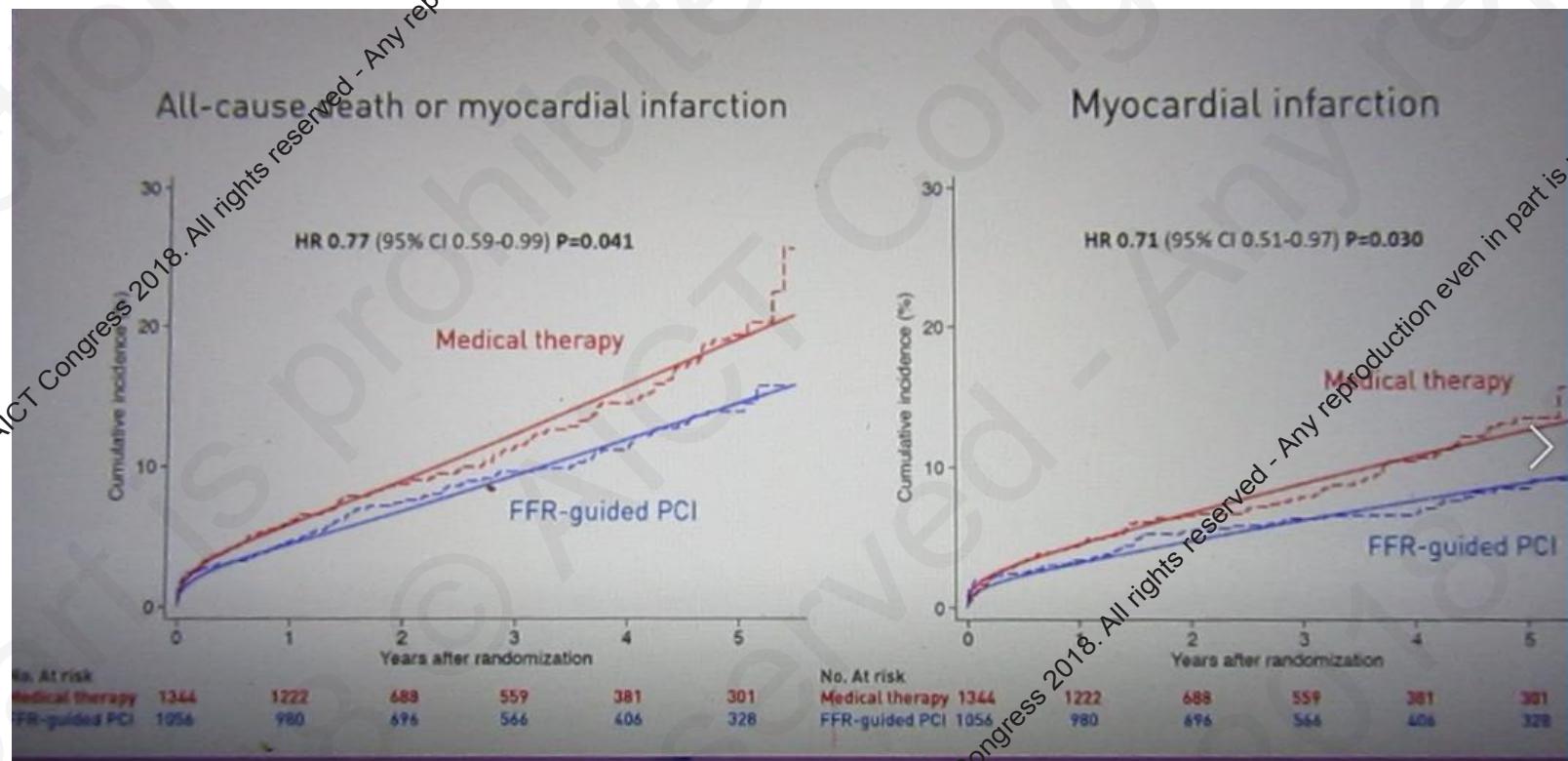
Thomas Engstrom, Henning Kalrauk, Staffan Holmberg, Dan J. Hofvind, Lars Klaesgaard, Lars Holmér, Erik Jorgenson, Lars Petersson, Lars

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Pooled analysis

Total number of patients included in this pooled analysis:

- 2400 patients
 - 1056 patients randomized to **FFR-guided PCI**
 - 1344 patients randomized to **Medical therapy**
- Primary endpoint: cardiac death or myocardial infarction
- Secondary endpoints: composite of all-cause death or myocardial infarction and the individual components: myocardial infarction, cardiac death, and all-cause death
- Subgroup analyses: clinical presentation, patients with at least 1 FFR-positive lesion, age, sex, diabetes



CONCLUSIONS

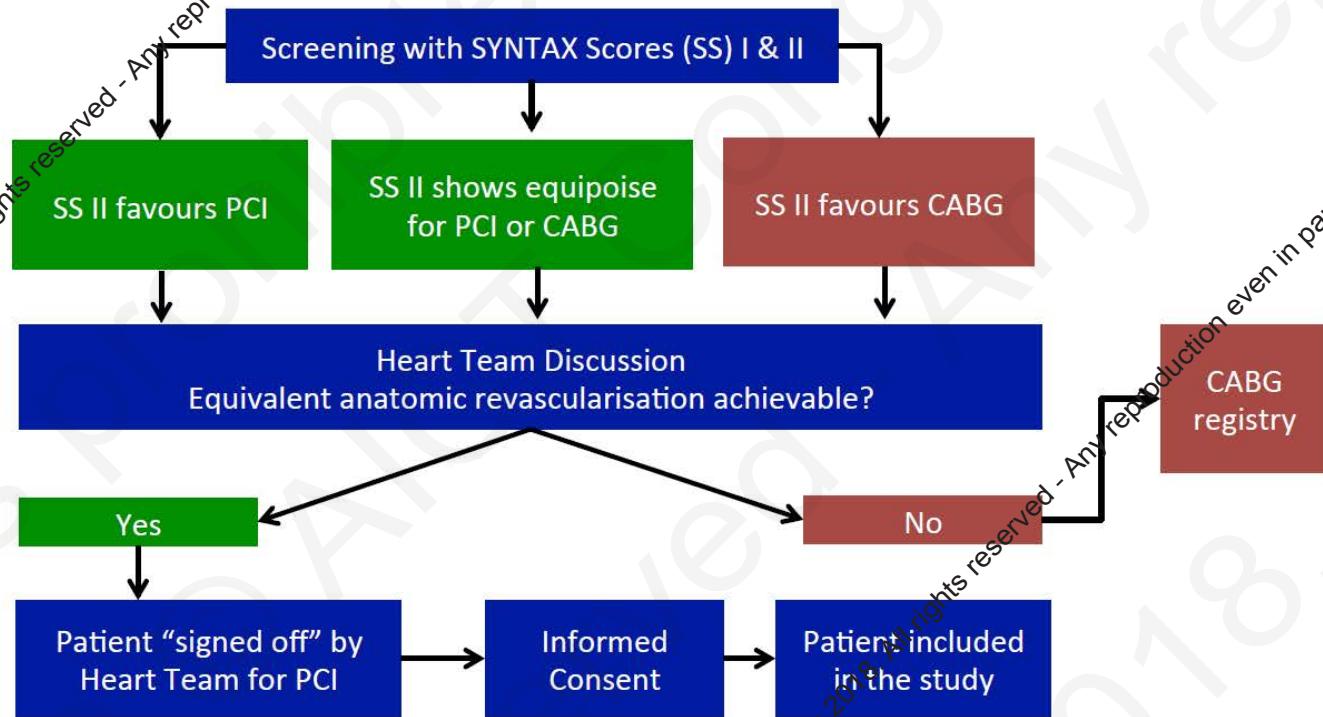
1. Initial FFR-guided PCI strategy resulted in a sustained clinical benefit, as compared with medical therapy alone, with regard to the composite primary end point of death, myocardial infarction, or urgent revascularization at 5 years.
2. Patients without hemodynamically significant stenoses had a favorable long-term outcome with medical therapy alone.

LIMITATIONS OF OCULOSTENOTIC REFLEX

- The potential benefit of revascularization depends on the presence and extent of myocardial ischemia.
- Performing PCI on nonischemic stenoses is not beneficial and is probably harmful.
- Thus, careful selection of ischemia-inducing stenoses is essential for deriving the greatest benefit from revascularization in patients with stable coronary artery disease.



Study flowchart: patient inclusion

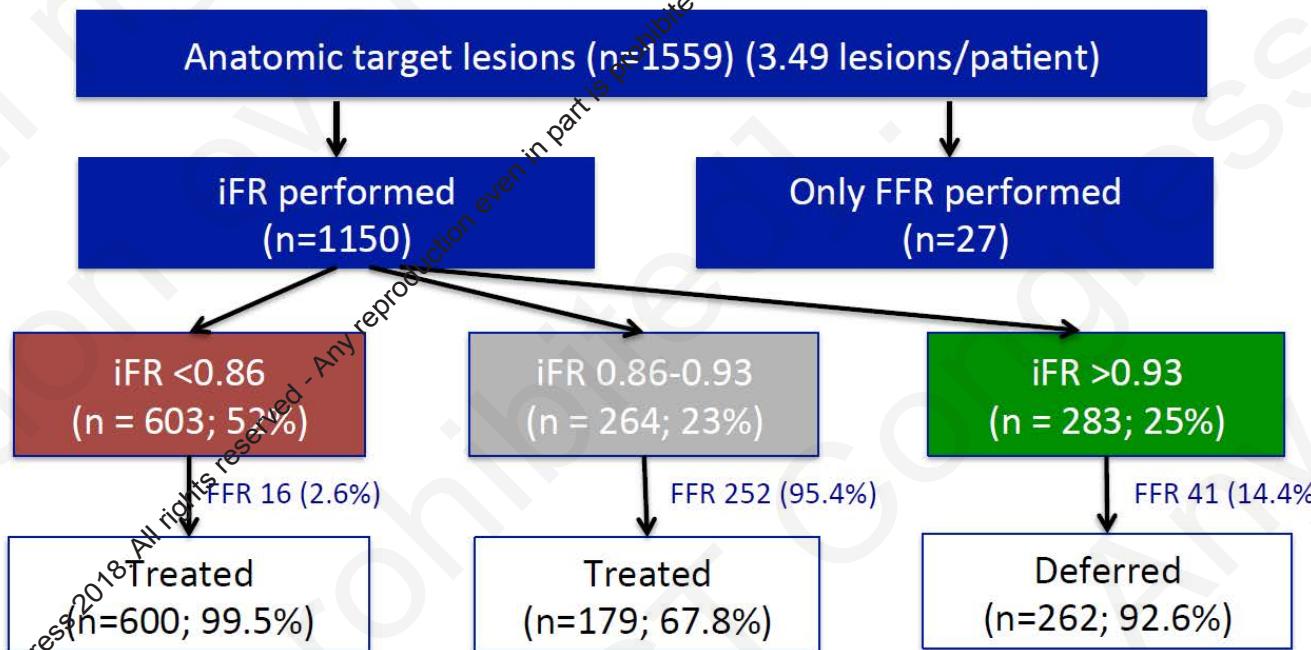




SYNTAX Score II

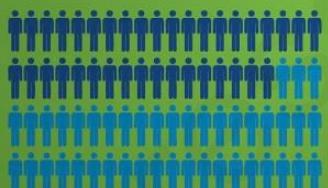
	SYNTAX II	SYNTAX I PCI arm	P value
Components of the SYNTAX Score II			
Age	66.7 ± 9.7	66.7 ± 9.1	0.99
Gender (Male)	93.2%	93.0%	0.93
Cr Clearance (ml/min)	82.0 ± 26.9	87.3 ± 28.5	0.008
Ejection Fraction (%)	58.1 ± 8.3	61.8 ± 11.3	<0.001
Peripheral Vascular Disease	7.7%	9.5%	0.37
COPDs	10.8%	12.7%	0.42
Anatomic SYNTAX Score	20.3 ± 6.4	22.8 ± 8.7	<0.001
SYNTAX Score II PCI			
	30.2 ± 8.6	30.6 ± 8.7	0.528
Predicted 4-yr mortality PCI (%)	8.9 ± 8.8%	9.2 ± 8.7%	0.640
SYNTAX Score II CABG			
	29.1 ± 10.4	29.1 ± 9.6	1.0
Predicted 4-yr mortality CABG (%)	9.0 ± 9.3	8.5 ± 8.1	0.440

Physiological stenosis interrogation

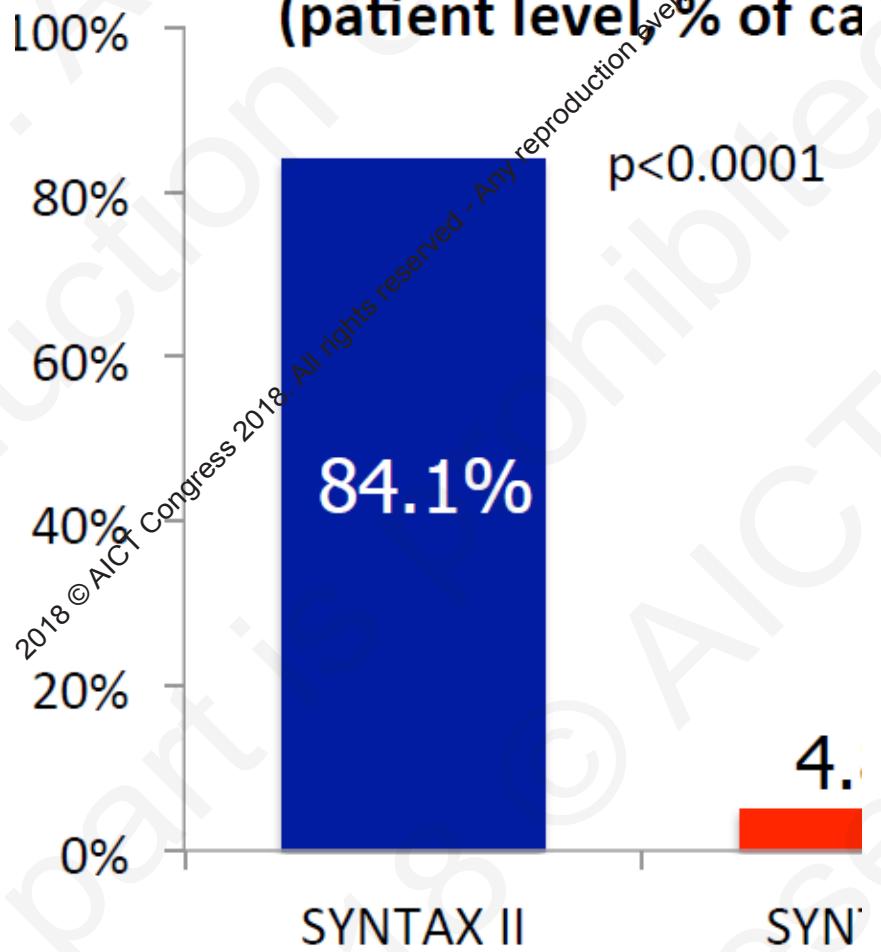


PHYSIOLOGY

The right patients were treated for the right reasons when physiology (FFR/iFR) is utilized.



IVUS use in SYNTAX II and (patient level) % of ca



STENT OPTIMIZATION

IVUS helps to optimize stent placement as a part of contemporary PCI.



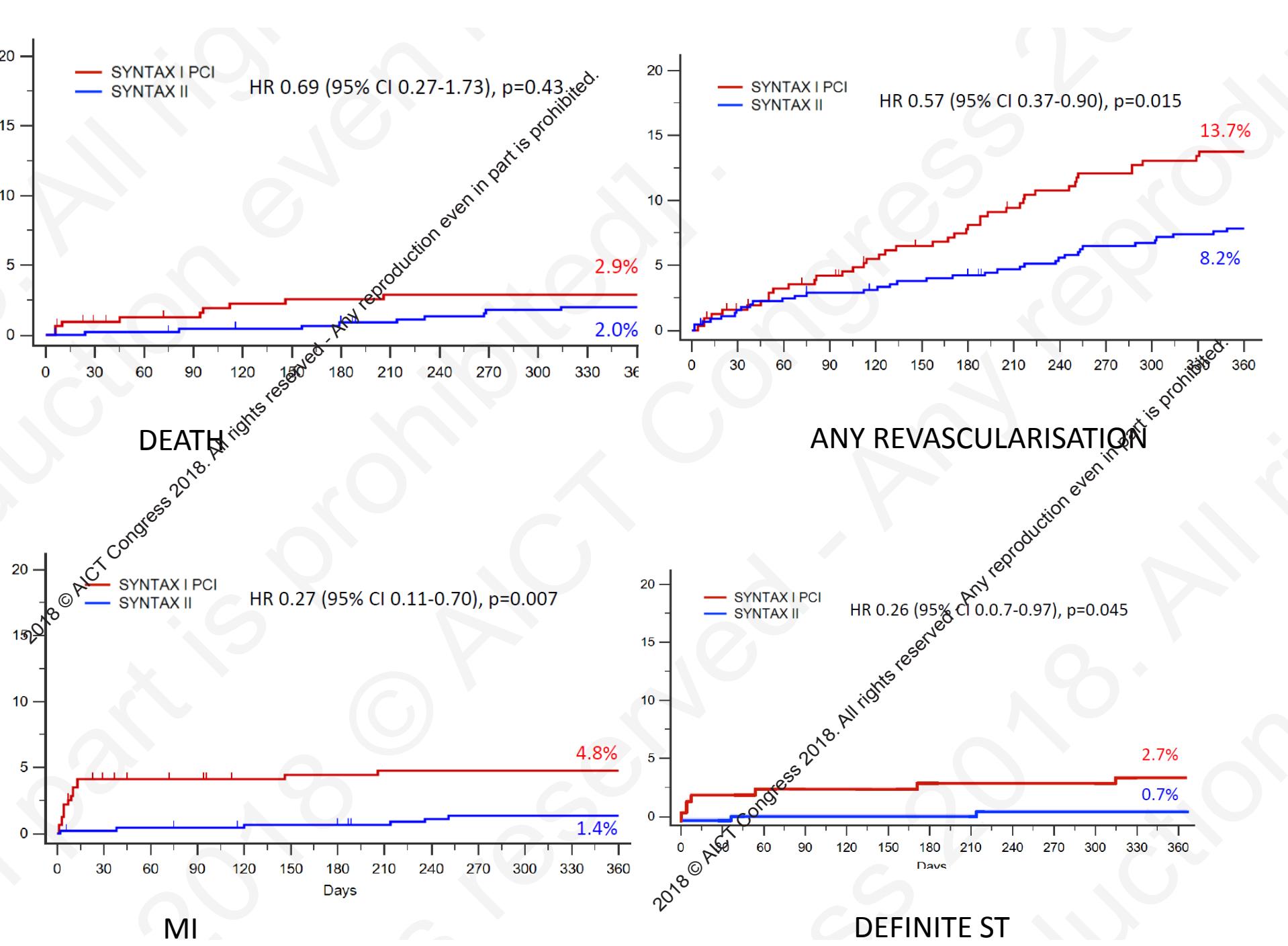
CTO success in SYNTAX 2 Trial

➤ CROSSING

Contemporary CTO PCI in SYNTAX II Trial demonstrated a significantly higher procedural success rate compared to those in SYNTAX I.

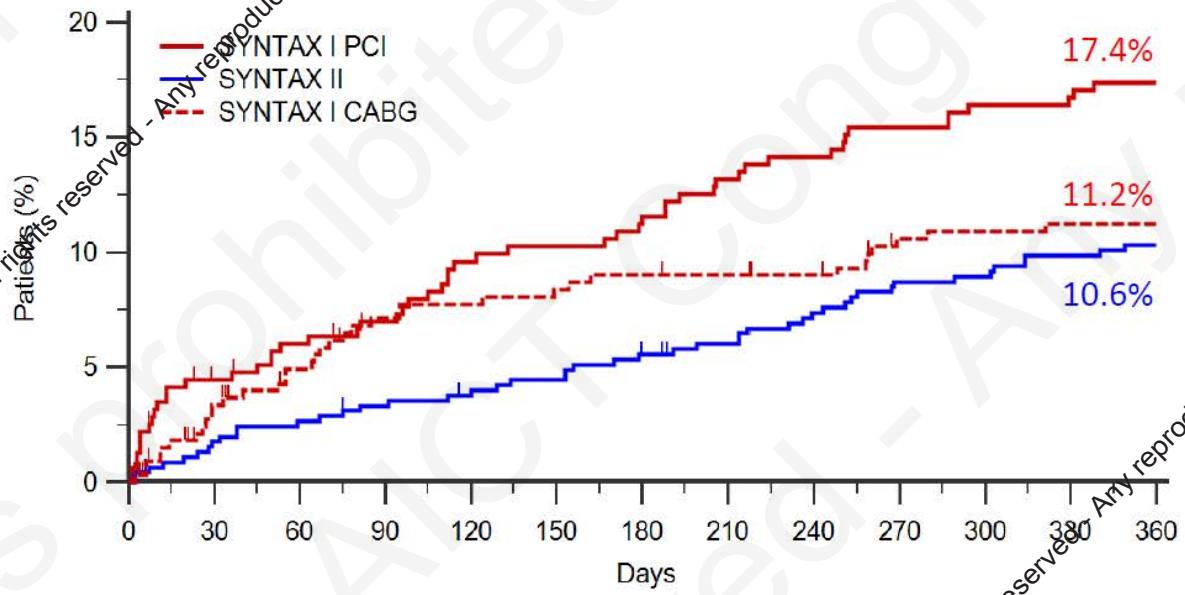
PCI with CTO procedural success rates jumped from **53% in SYNTAX I to 87% in SYNTAX II**. Representing a **64% increase in successful CTO treatment**.







MACCE SYNTAX II and SYNTAX I PCI / CABG



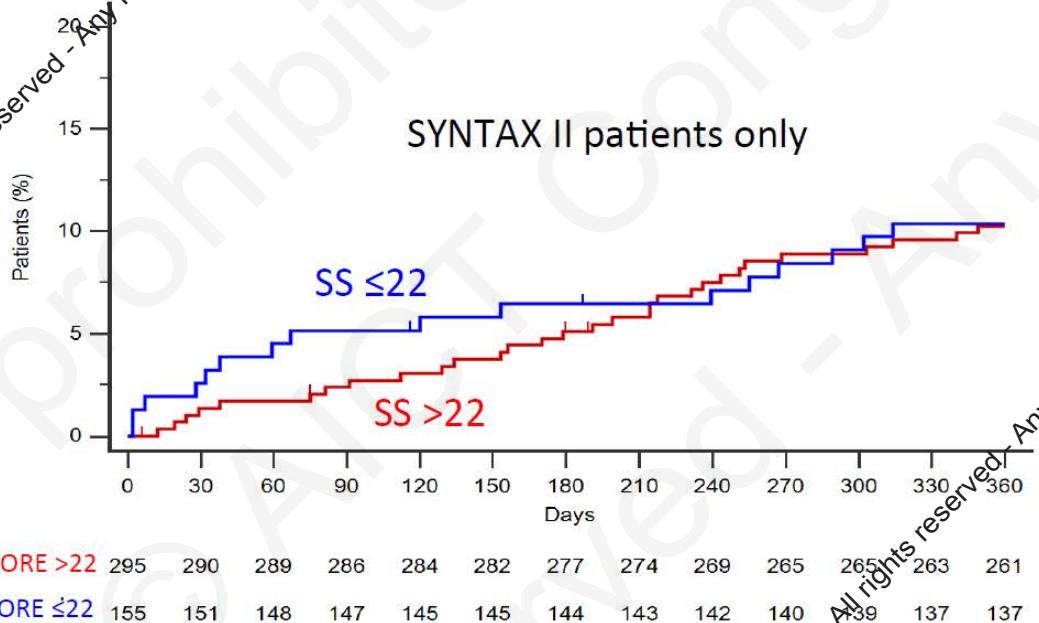
SYNTAX I PCI	315	298	292	288	280	278	274	269	266	262	259	258	256
SYNTAX II	450	441	437	433	429	427	421	417	411	405	400	398	
SYNTAX I CABG	334	313	304	295	293	291	289	288	287	279	278	277	277

Advances in the SYNTAX 2 trial

1. New stratification tool – clinical variables
2. Use of iFR and FFR
3. Second generation DES – Synergy
4. IVUS guided stent optimisation (MUSIC criteria)
5. Contemporary CTO revascularisation techniques
6. Guideline based OMT ($LDL < 1.8$)

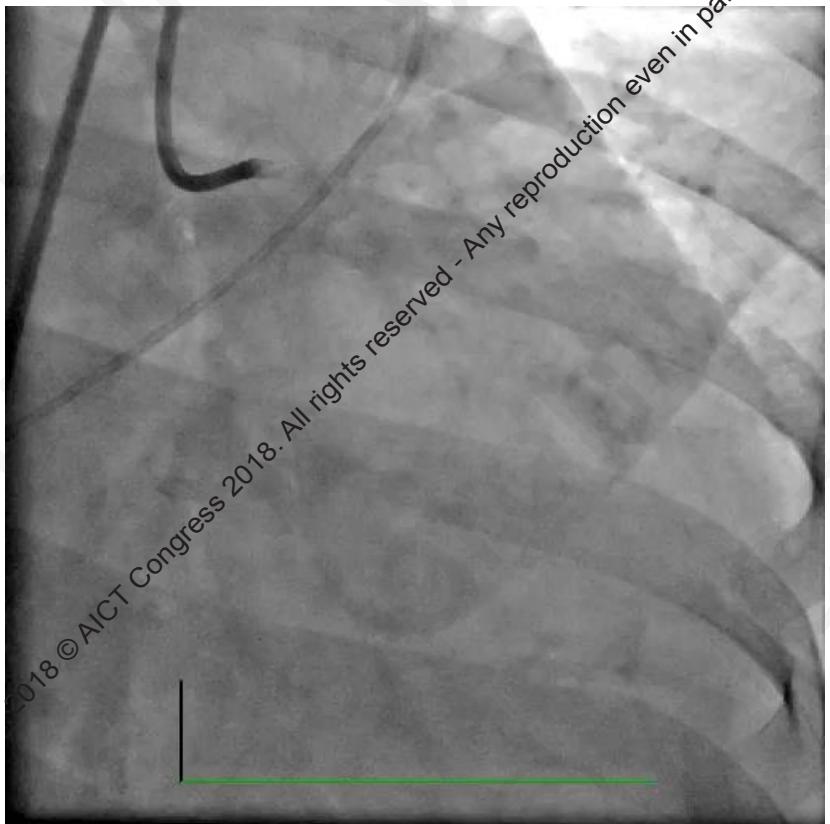


SYNTAX II MACCE in SS I ≤ 22 and >22



European Heart Journal (2017) 38, 3124–3134

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Case

64 yrs old lady

Type 2 diabetes on insulin for >20 yrs

HbA1c- 8.8

Severe Diabetic Neuropathy ,
Retinopathy

Symptoms – 2 HF admissions in 3
months

LVEF – 25% , PASP -48mm, mild MR

PET scan – no infarction , large areas of
Hibernation

S Creatinine 1.2 mg%, Urine
microalbuminuria present

SYNTAX Score overview

The SYNTAX Score has been calculated successfully for this patient.

Please note that this anatomy was not covered in the SYNTAX Trial.

Print results or save as PDF
The fields below are optional



Patient ID: MM0025040

Print

Name: UD

Save

Date of birth: 1952



Continue with next patient
Current data will be lost, please print/save first

New patient

SYNTAX Score II

SYNTAX II

Decision making -between CABG and PCI- guided by the SYNTAX Score II to be endorsed by the Heart Team.

PCI

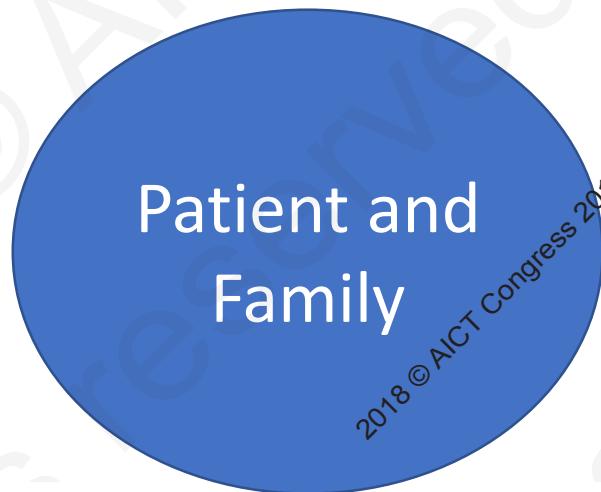
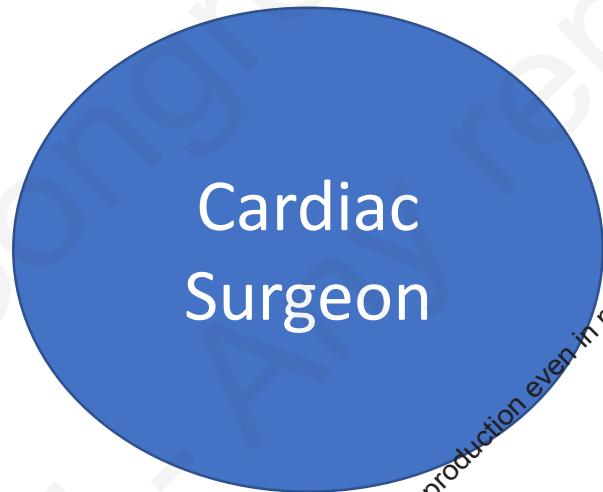
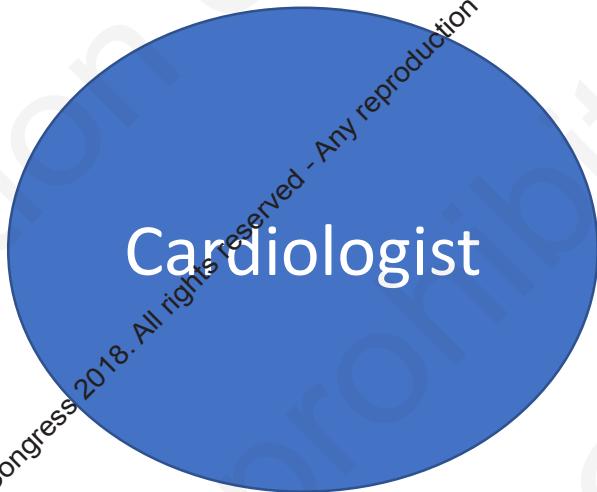
SYNTAX Score II: 53.8
PCI 4 Year Mortality: 40.7 %

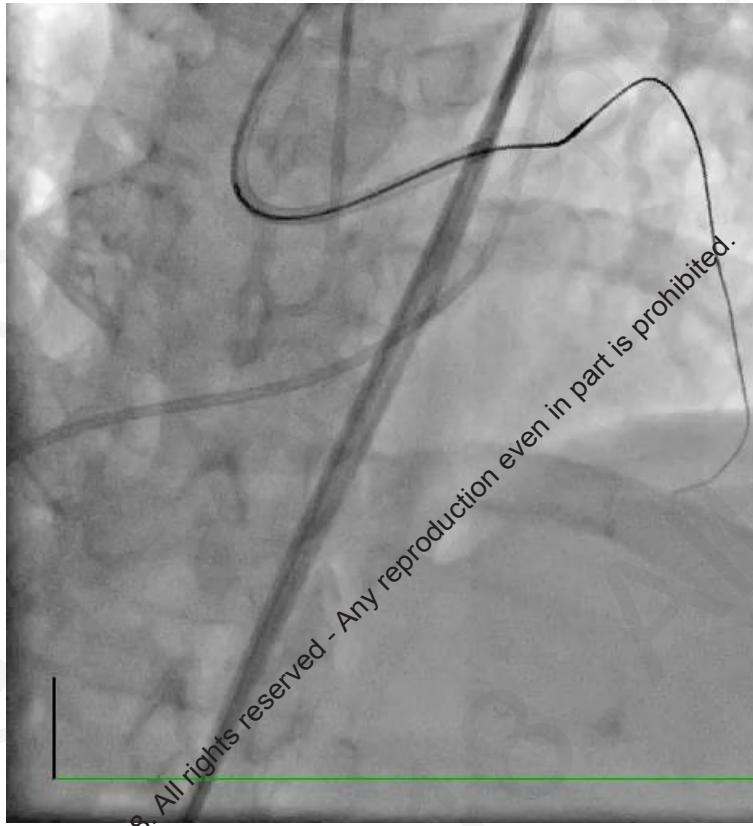
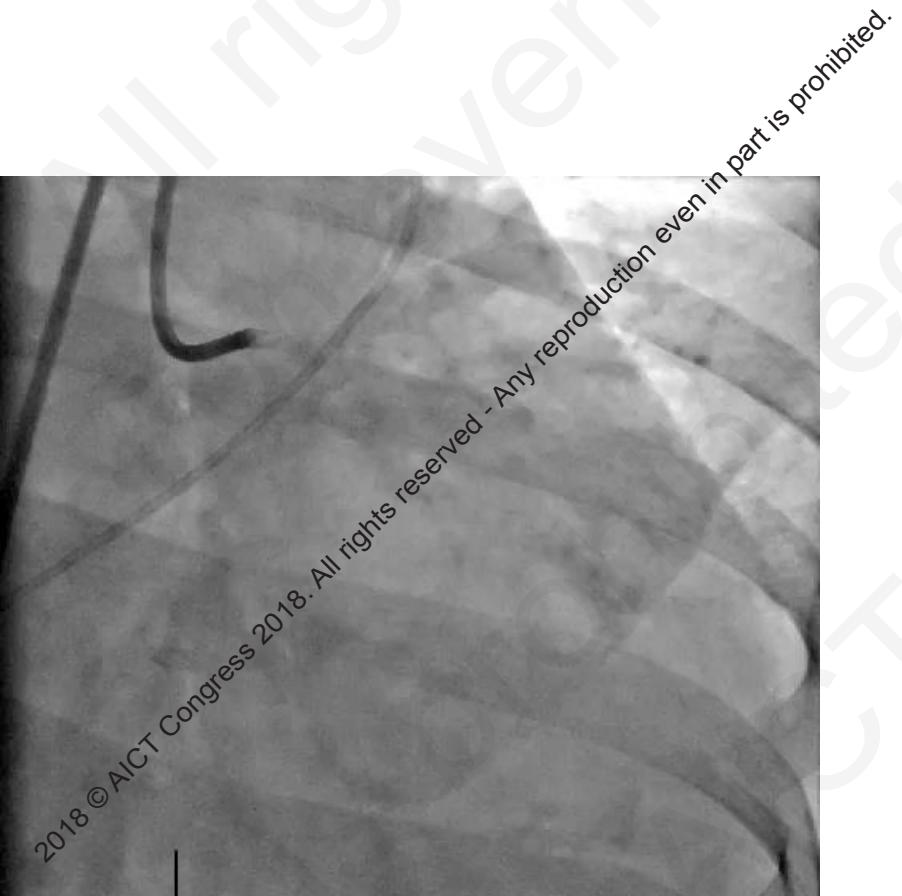
CABG

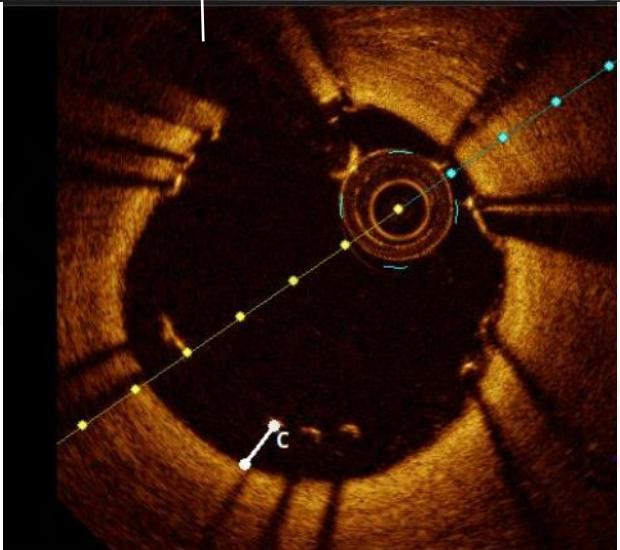
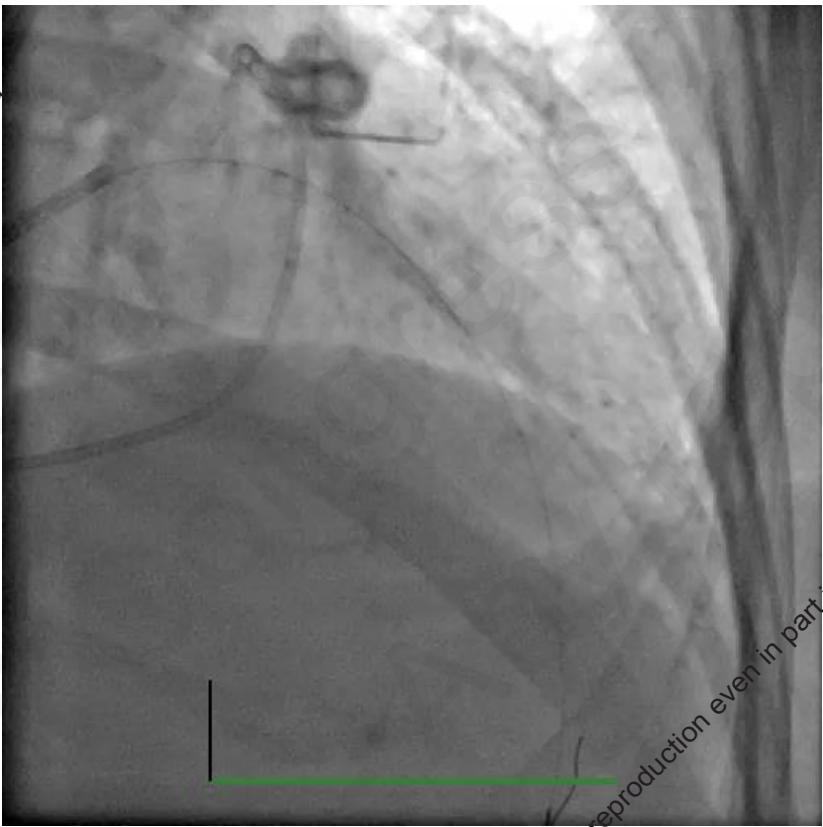
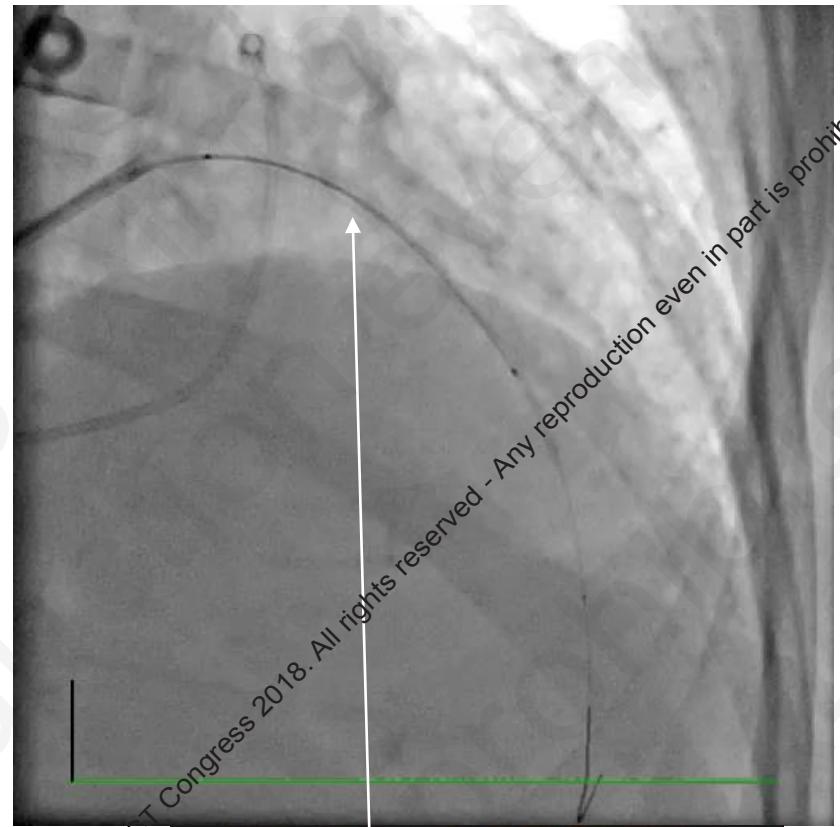
SYNTAX Score II: 26.4
CABG 4 Year Mortality: 5.4 %

Treatment recommendation: CABG

Heart Team Approach

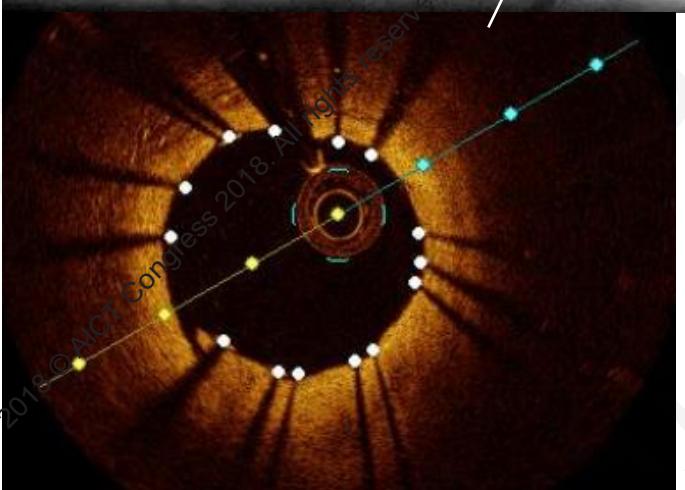
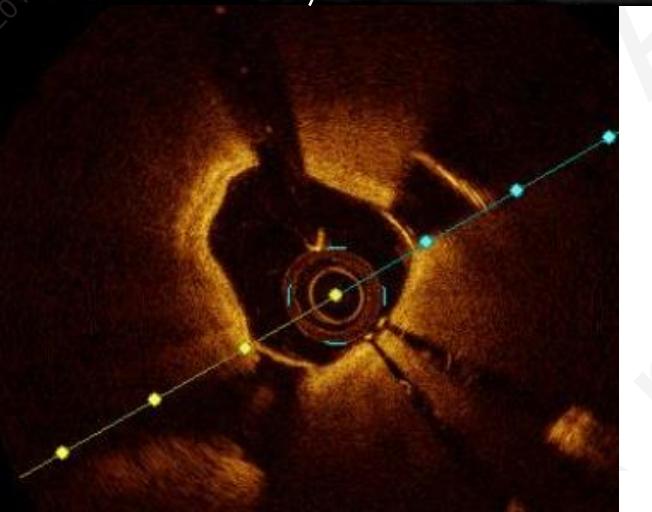
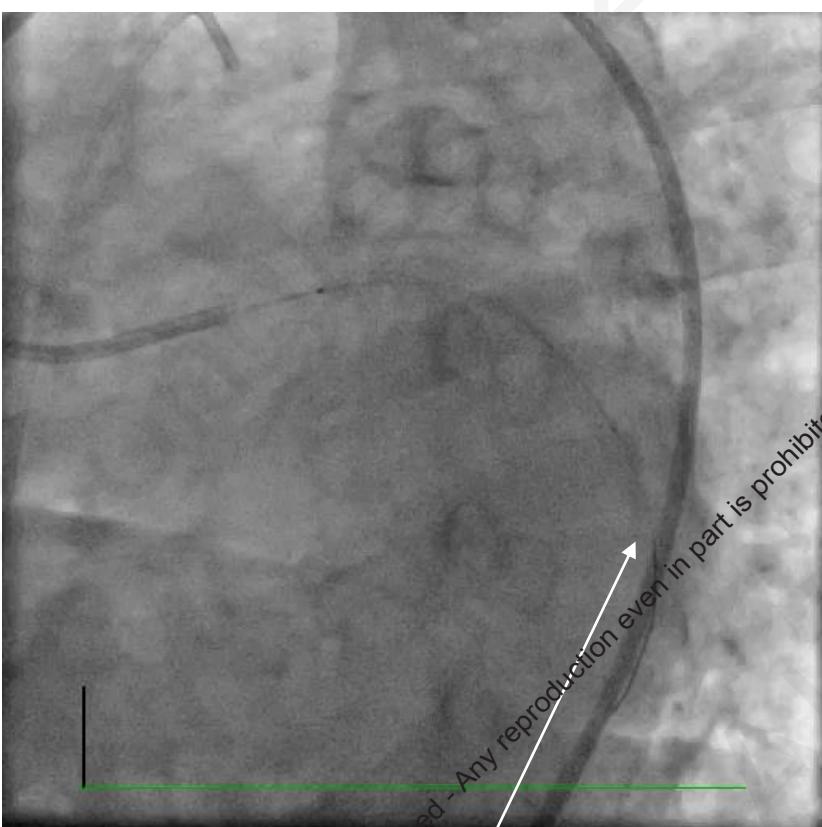
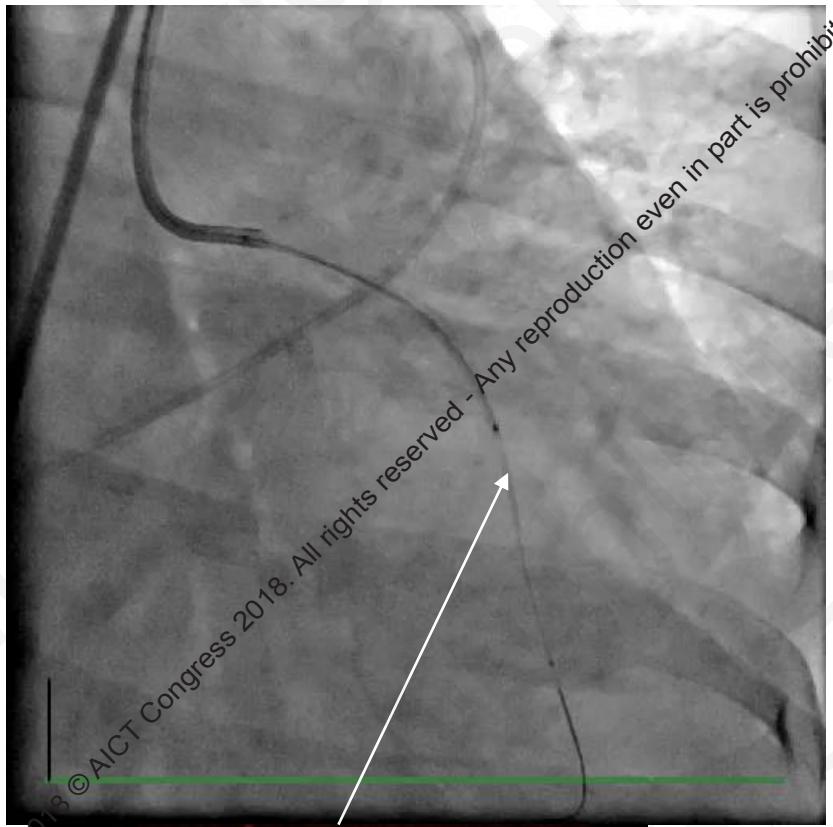






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Follow up at 6 m

- LVEF – 45%
- Asymptomatic
- Diabetes still controlled
- OMT = ARNI , beta blockers , ivabradine high dose statins , aspirin and ticagrelor
- Diabetes is still uncontrolled

CONCLUSION

Multivessel PCI on its 40th anniversary: finally a match for CABG?

Gregg W Stone 

European Heart Journal, Volume 38, Issue 42, 7 November 2017, Pages 3135–3138,



THANK YOU ↗

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ASIAN INTERVENTIONAL CARDIOVASCULAR THERAPEUTICS
THE OFFICIAL CONGRESS OF APSIC



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