

OCT: Pre- & Post-intervention



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Wakayama Medical University



Disclosure Statement of Financial Interest

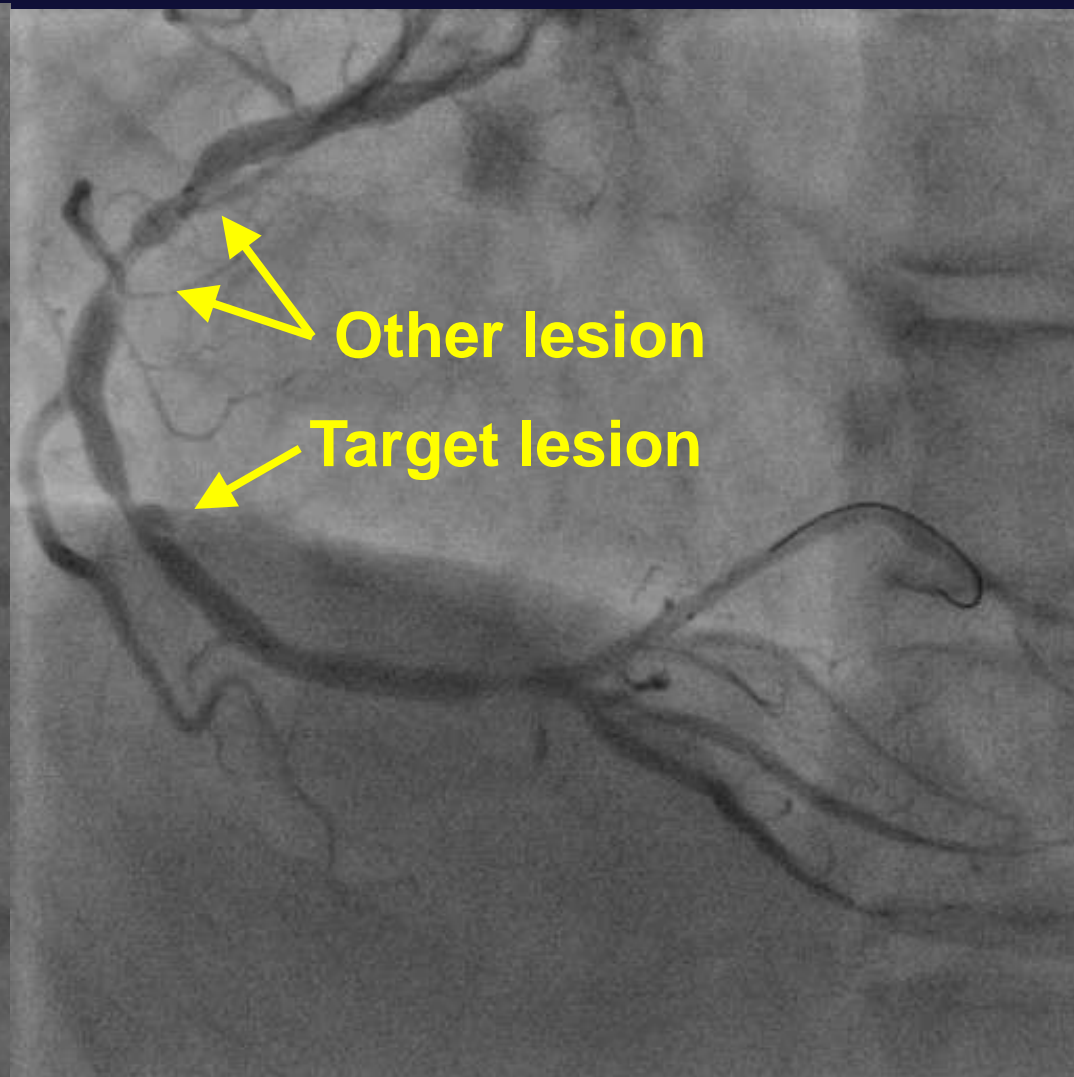
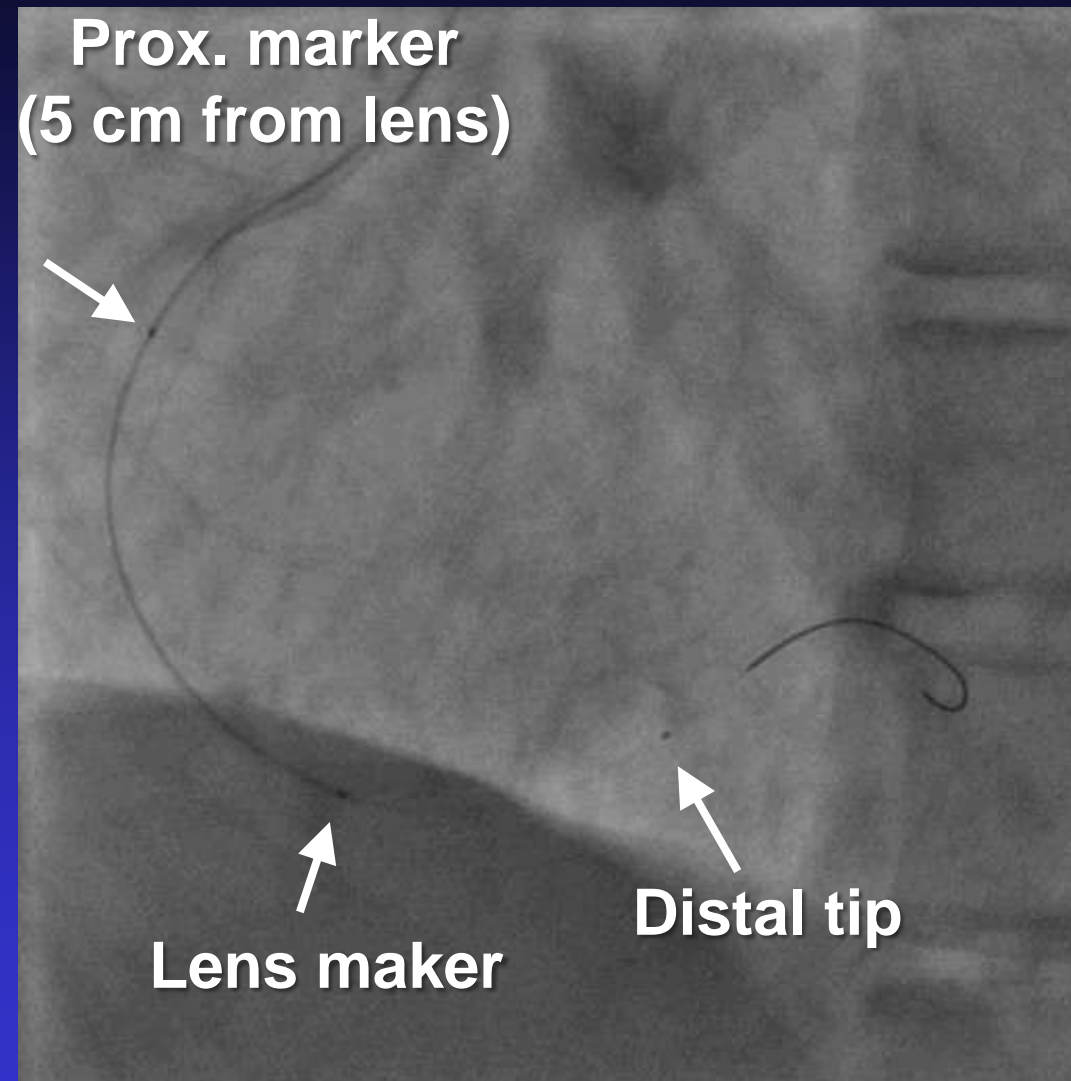
Within the past 12 months, I or my spouse/partner have had a financial interest/arrangement or affiliation with the organization(s) listed below.

Affiliation/Financial Relationship

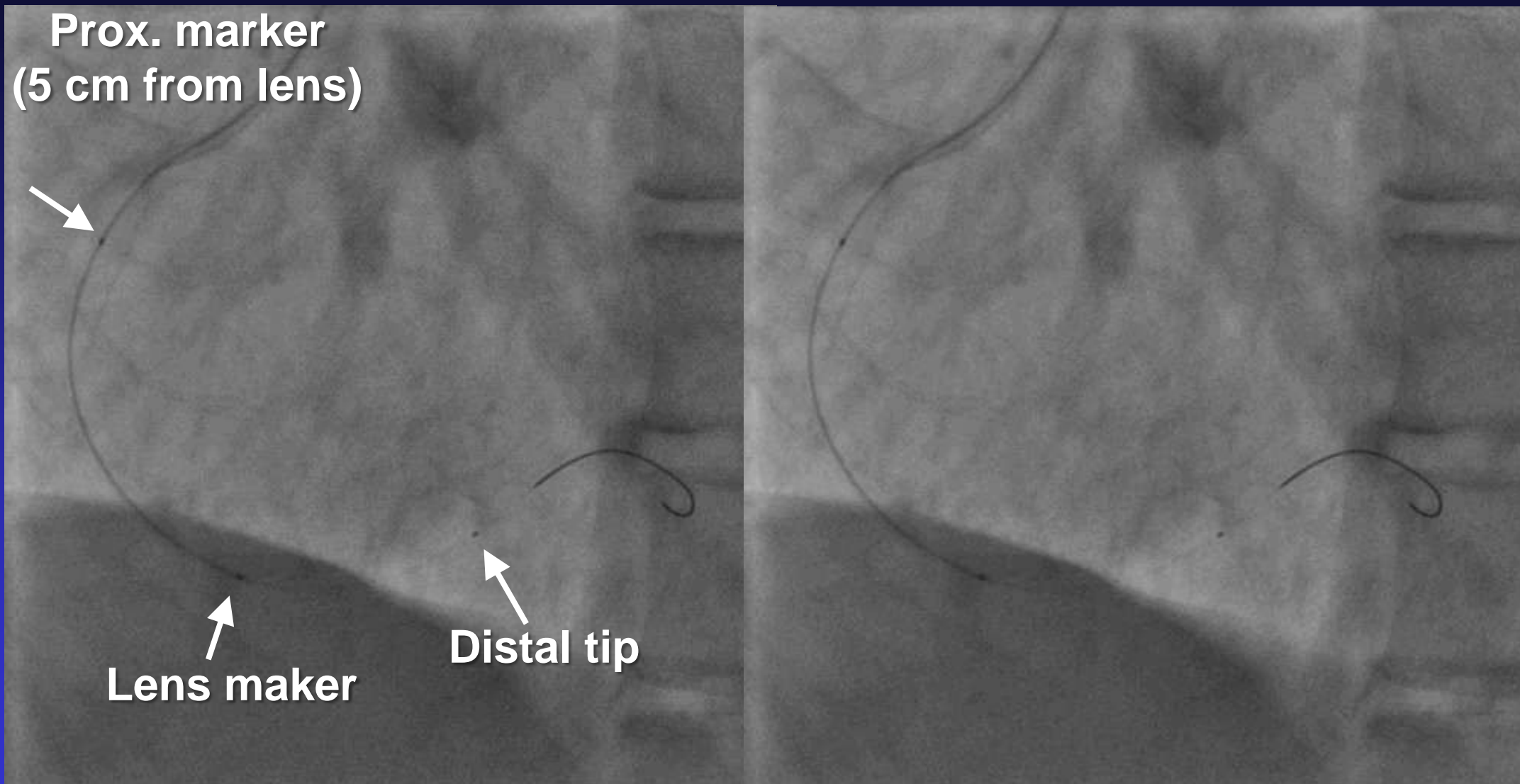
- Grant/Research Support : Abbott Vascular Japan
Boston Scientific Japan
Goodman Inc.
Sent Jude Medical Japan
Terumo Inc.
- Consulting Fees/Honoraria : Astellas Pharmaceutical Inc.
Daiichi-Sankyo Pharmaceutical Inc.
Goodman Inc.
Sent Jude Medical Japan
Terumo Inc.



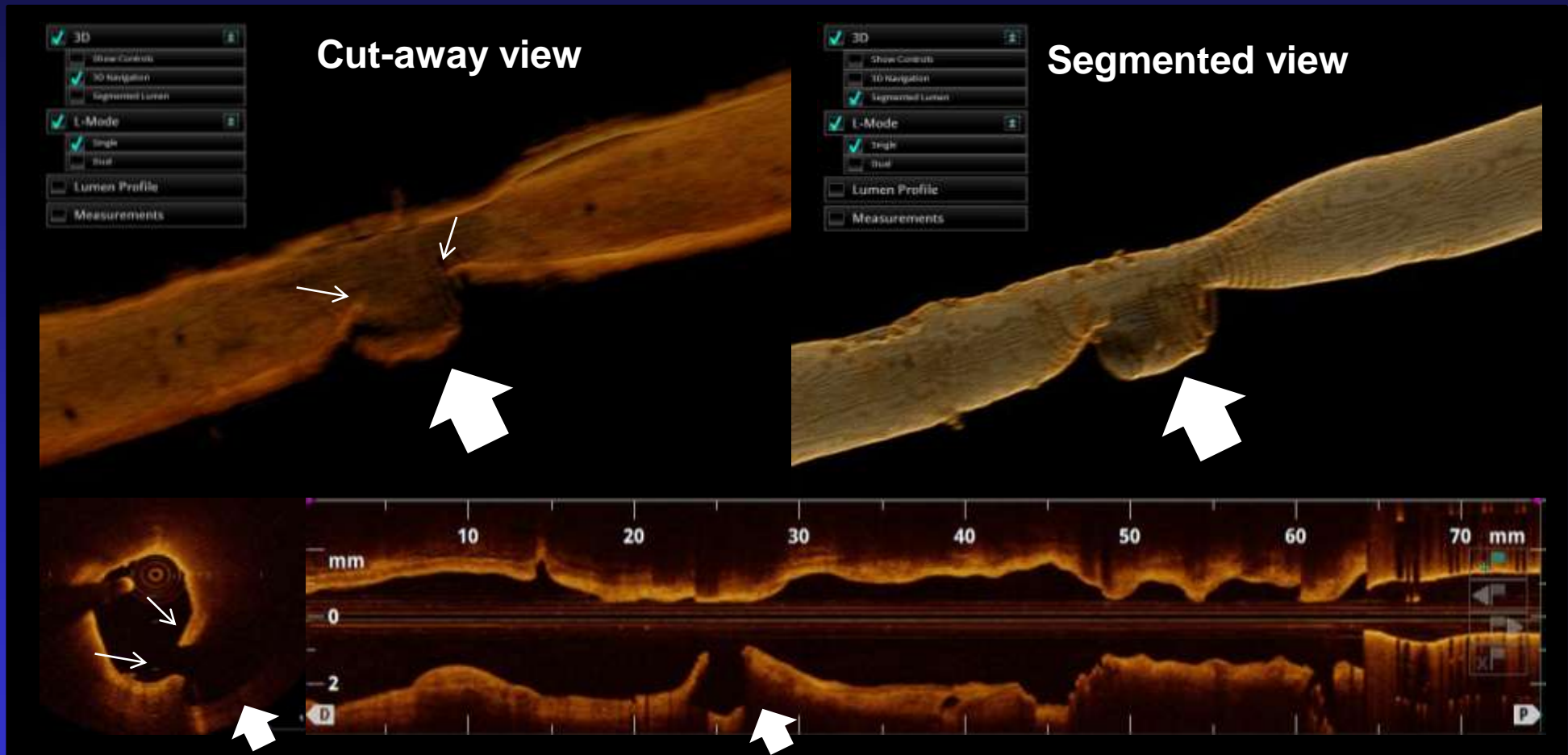
Positioning of OCT Catheter



Positioning of OCT Catheter

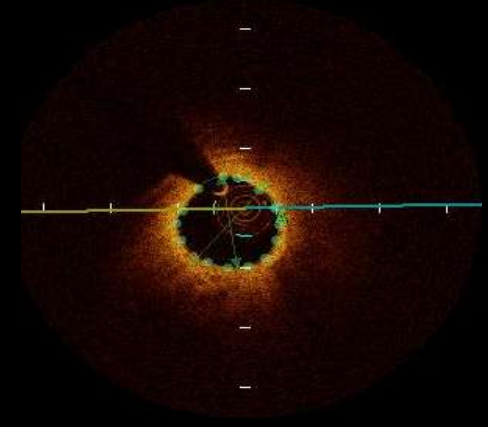
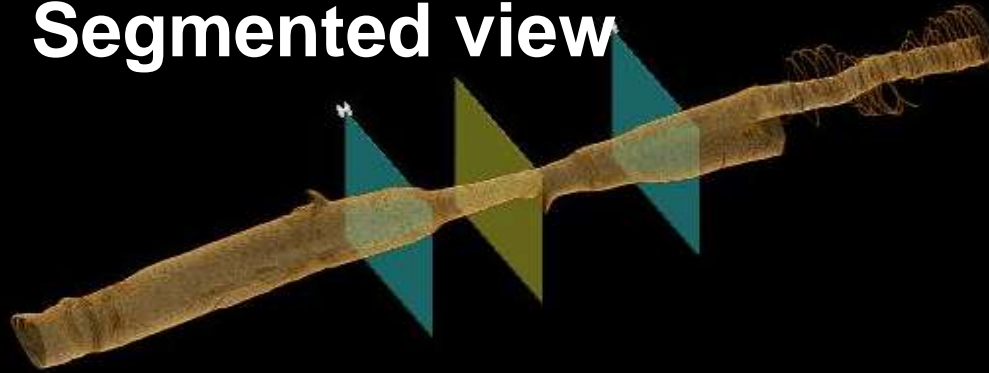


Advantages of Newly developed FD-OCT system (ILUMIEN OPTIS[®])

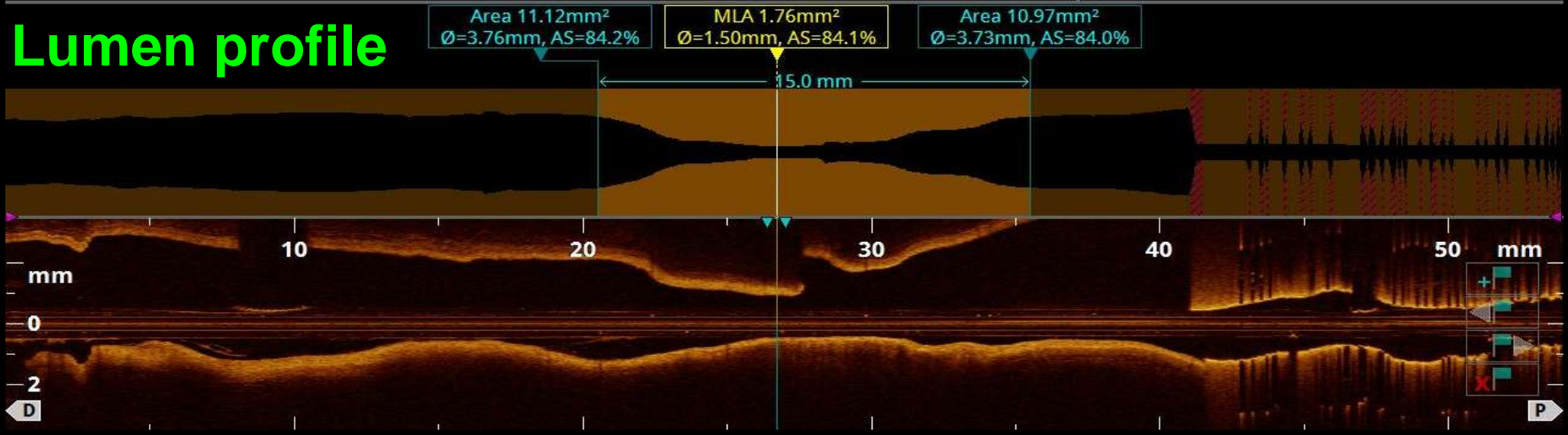


Pre-PCI assessment, #6 90%, (MultiLink 4.0 × 15mm)

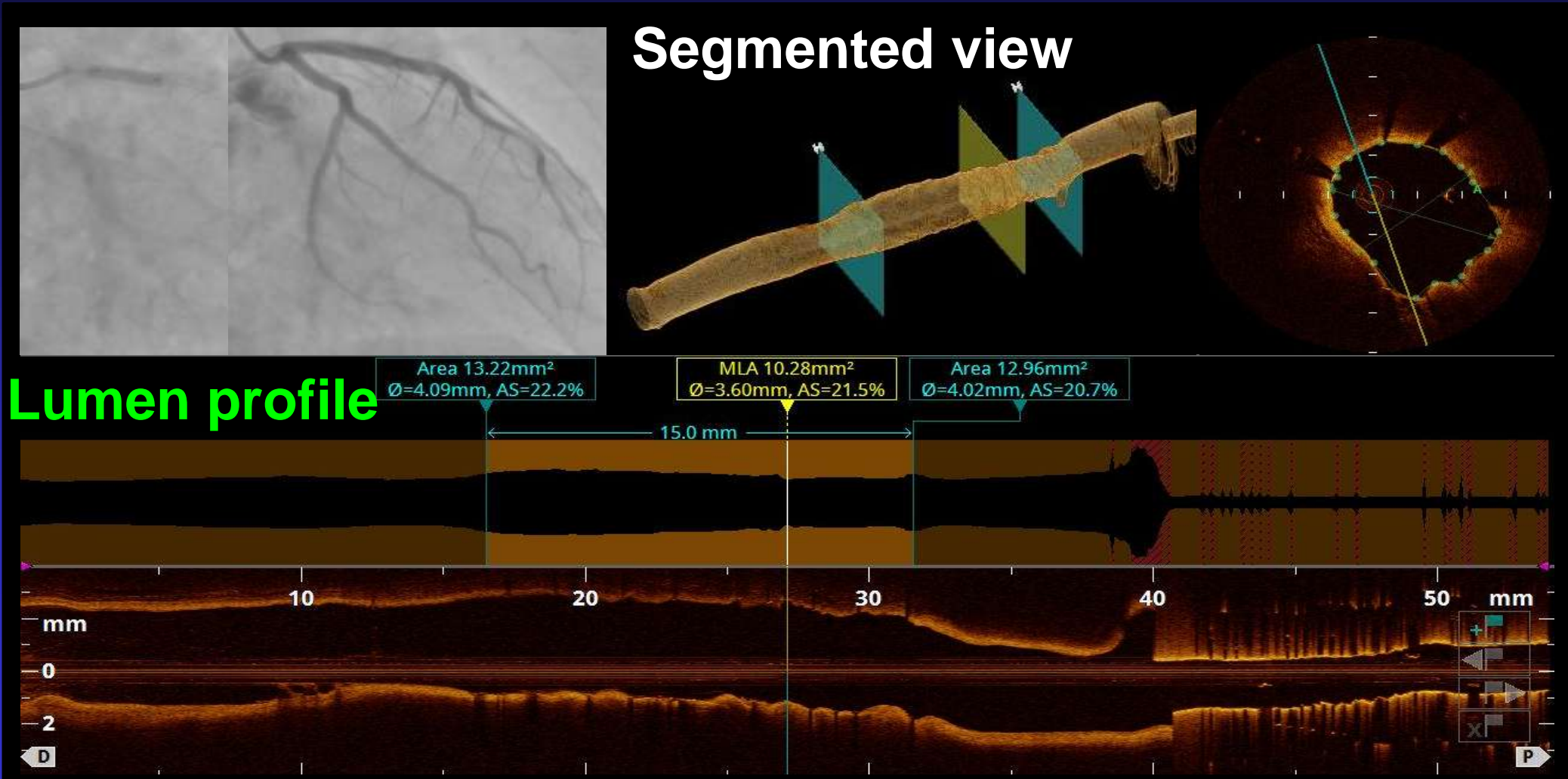
Segmented view



Lumen profile

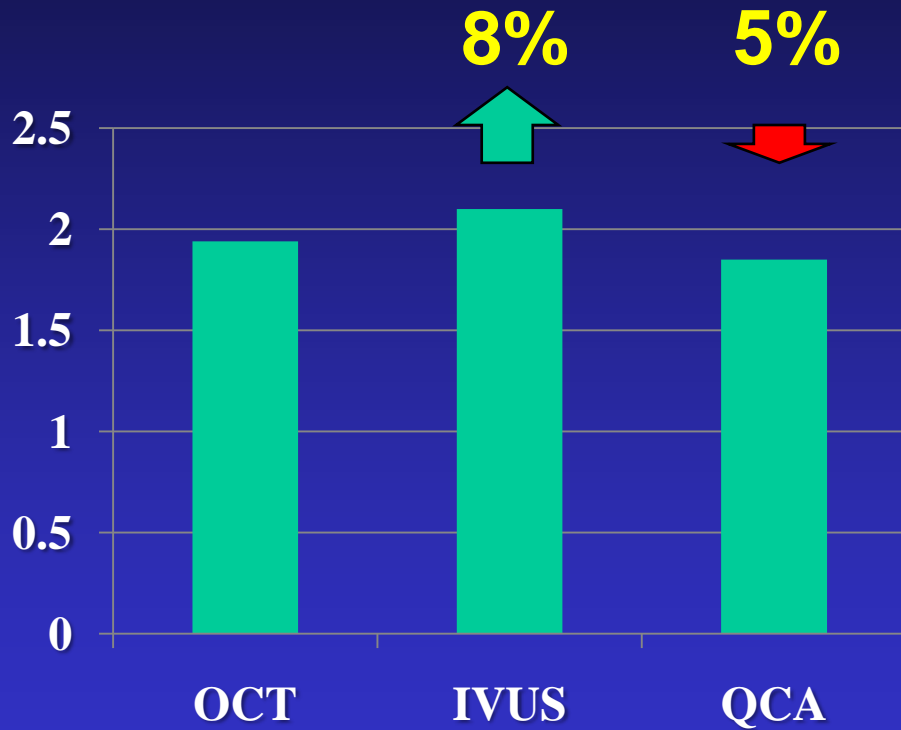


Post-PCI assessment, #6 90%, (MultiLink 4.0 × 15mm)



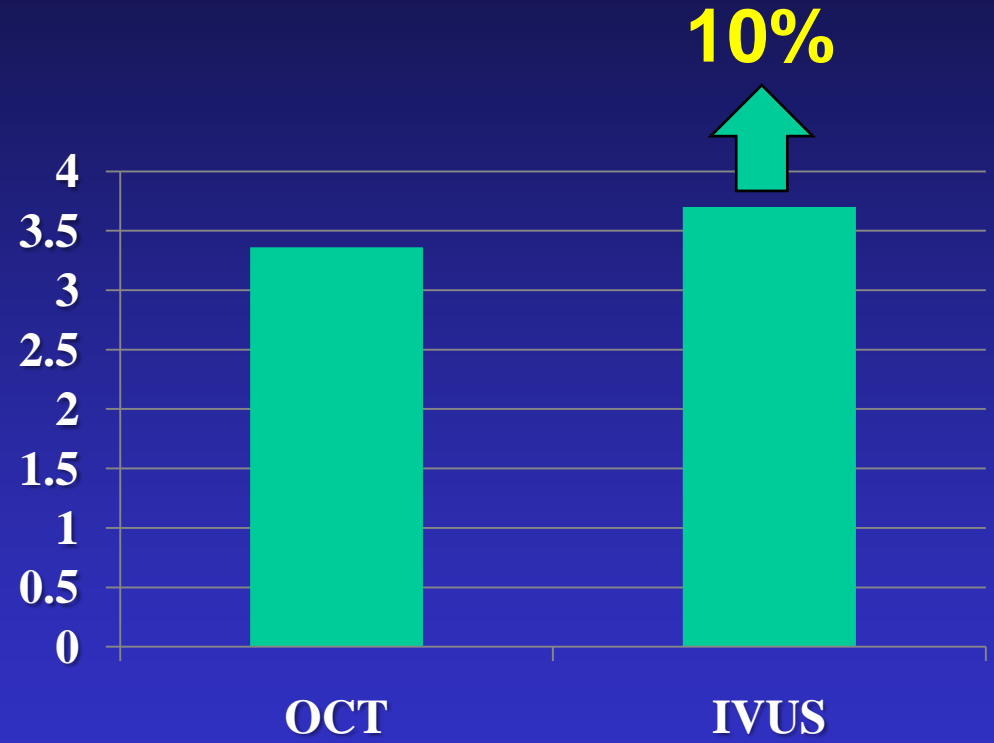
Comparison of measurements among OCT, IVUS & QCA (OPUS-CLASS study)

MLD



1.94 ± 0.70 mm **2.10 ± 0.59 mm** **1.85 ± 0.77 mm**

MLA



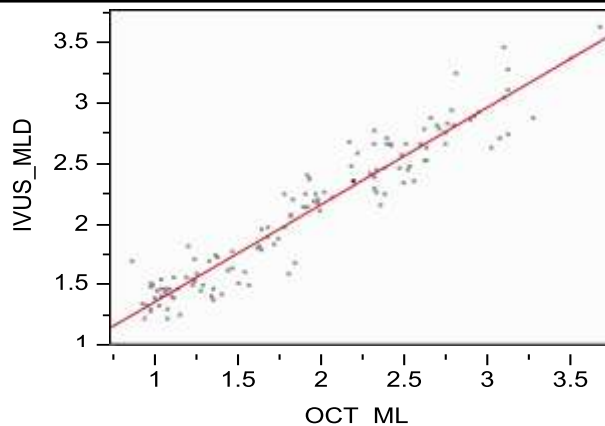
3.36 ± 2.28 mm² **3.70 ± 2.04 mm²**



MLD & MLA

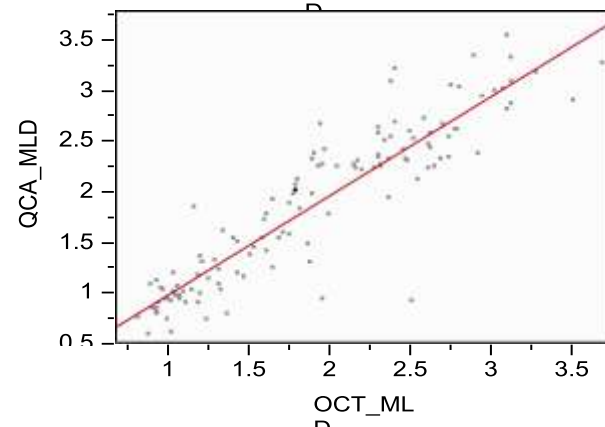
IVUS vs. OCT

$R^2=0.92$, $P<0.001$



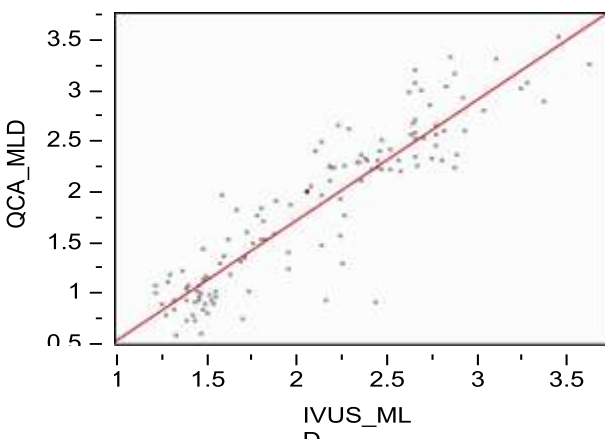
QCA vs. OCT

$R^2=0.83$, $P<0.001$



QCA vs. IVUS

$R^2=0.82$, $P<0.001$



$OCT = 2.11 \pm 0.82mm$

$IVUS = 2.27 \pm 0.61mm$

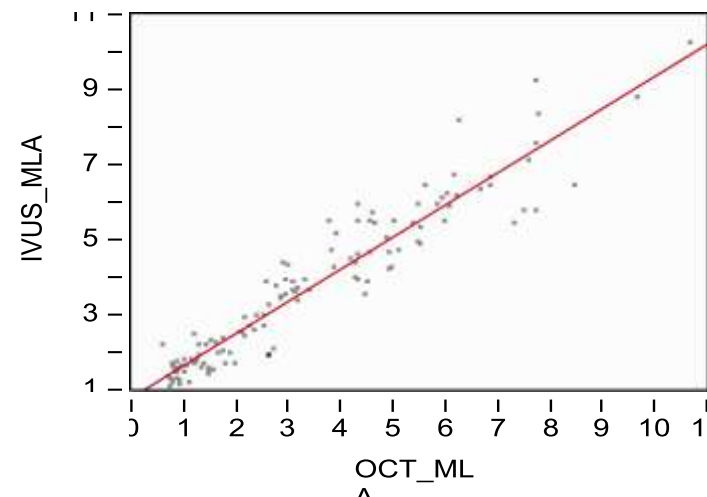
$QCA = 2.03 \pm 0.71mm$

*Kubo T, et al,
JACC Cardiovasc Img. 2013;6:1095-1104*

IVUS vs. OCT

$R^2=0.91$, $P<0.001$

$y = 0.83 + 0.85x$

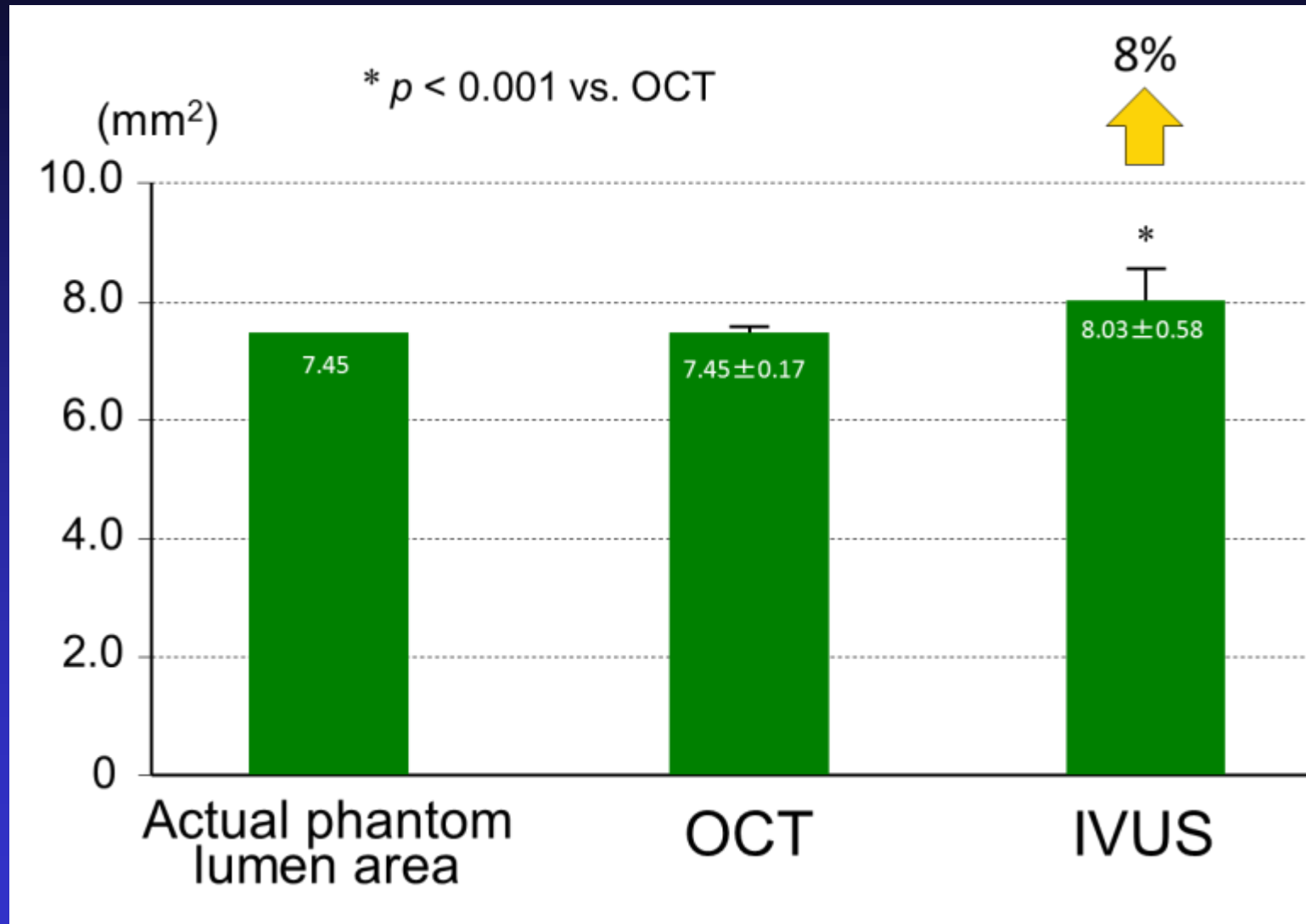


$OCT = 3.36 \pm 2.28mm^2$

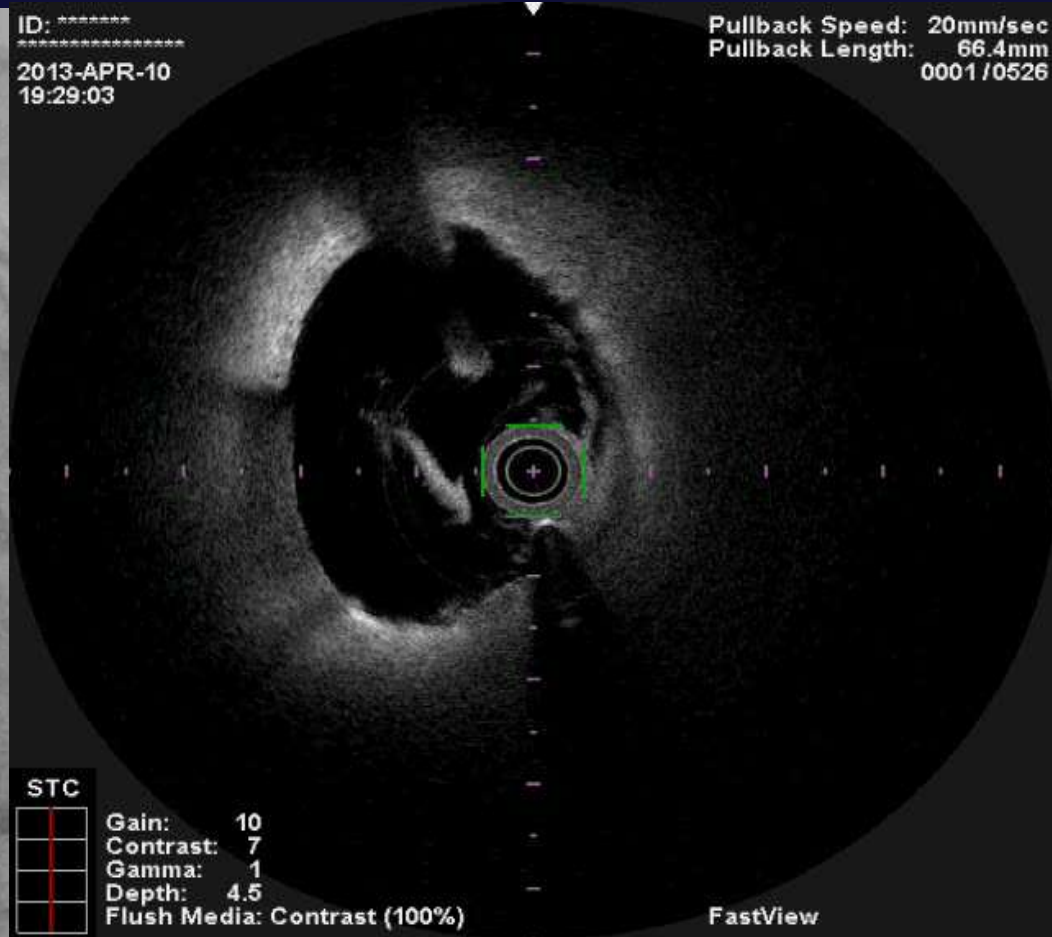
$IVUS = 3.70 \pm 2.04mm^2$



Accuracy of the measurement: MLA



Pre-PCI FD-OCT

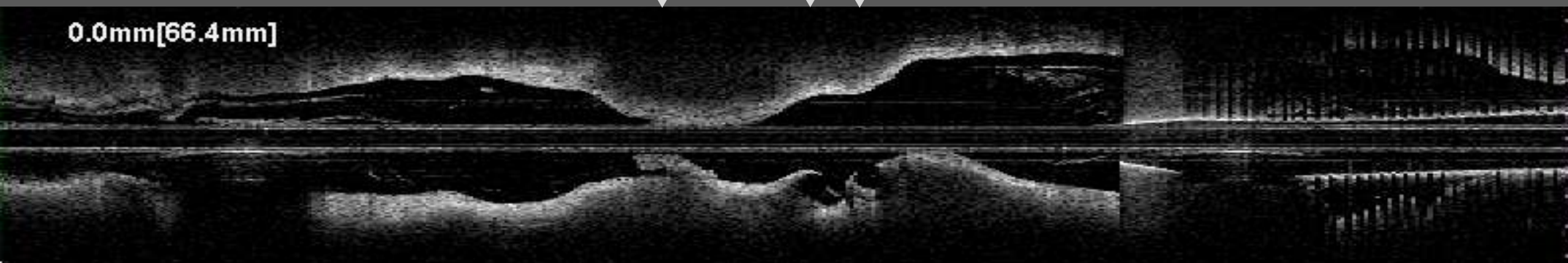
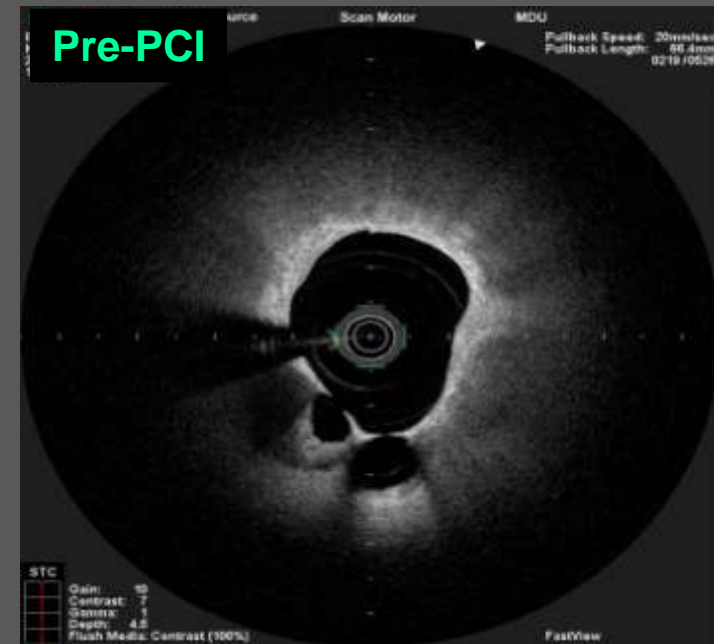
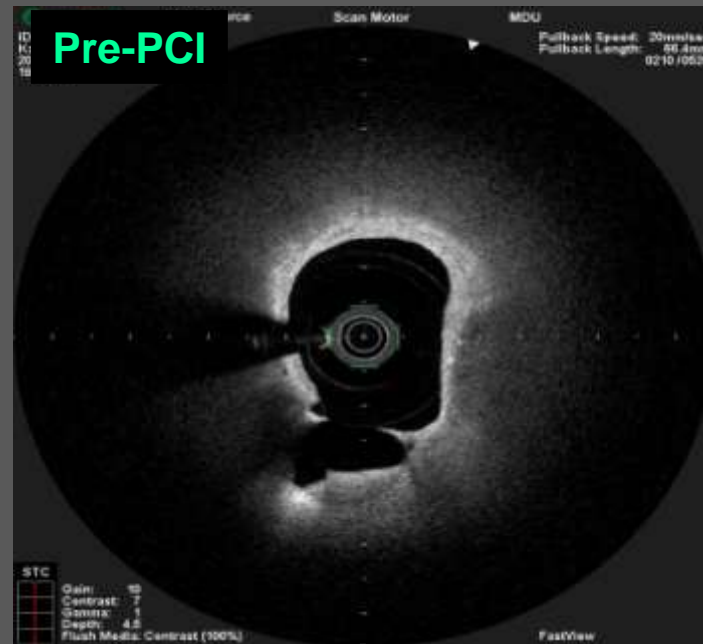
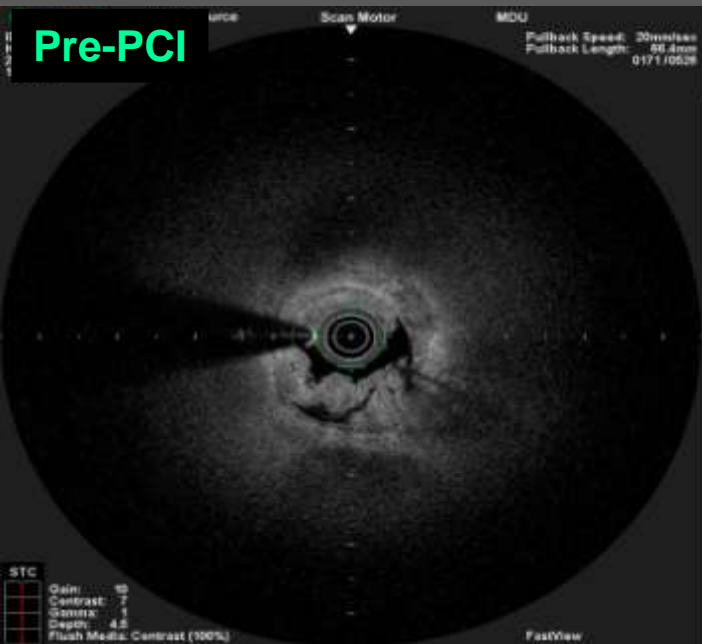


OFDI at culprit site

Thrombus

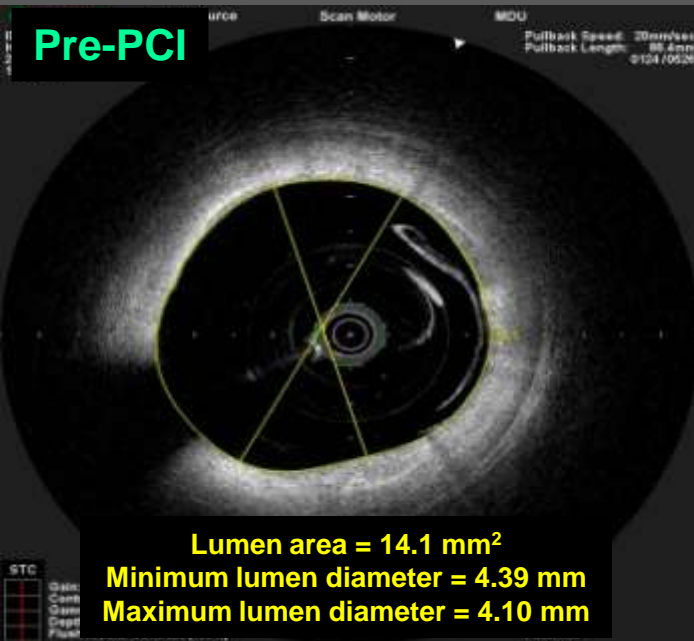
Plaque rupture

Plaque rupture

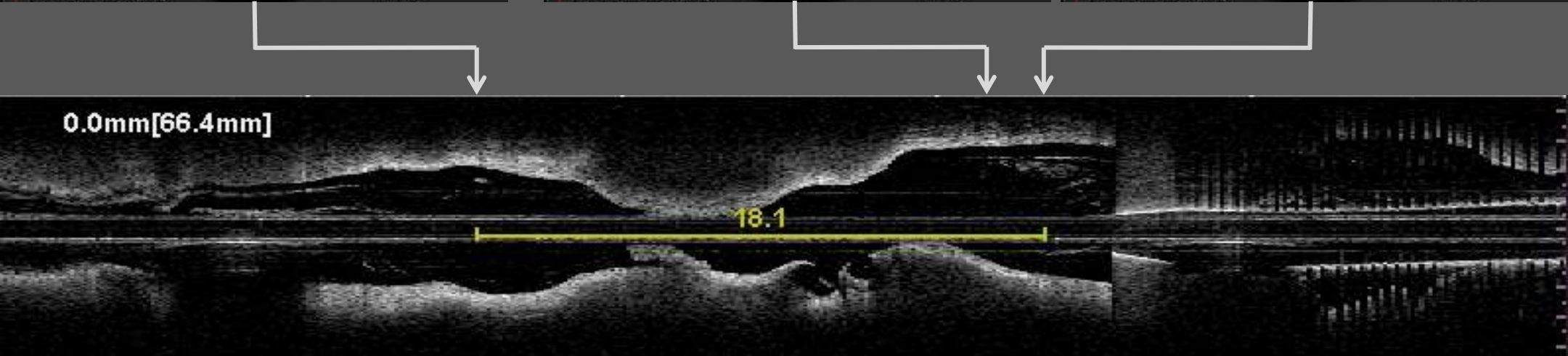
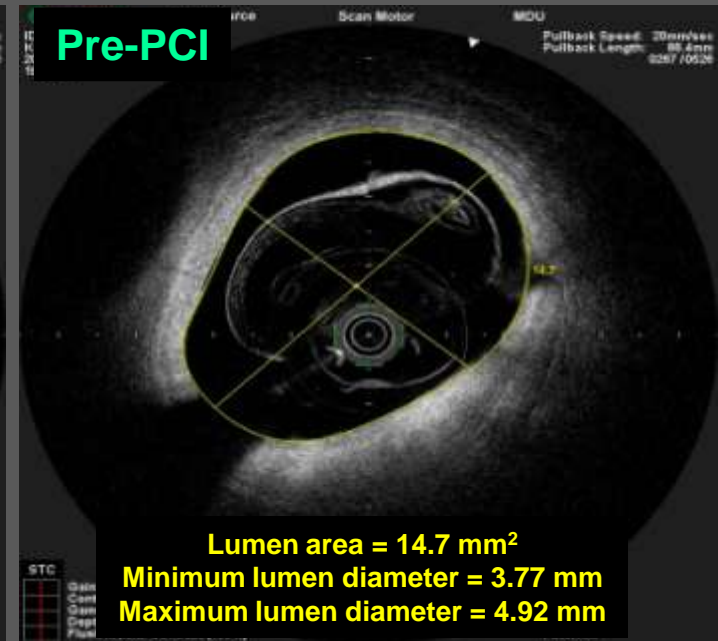
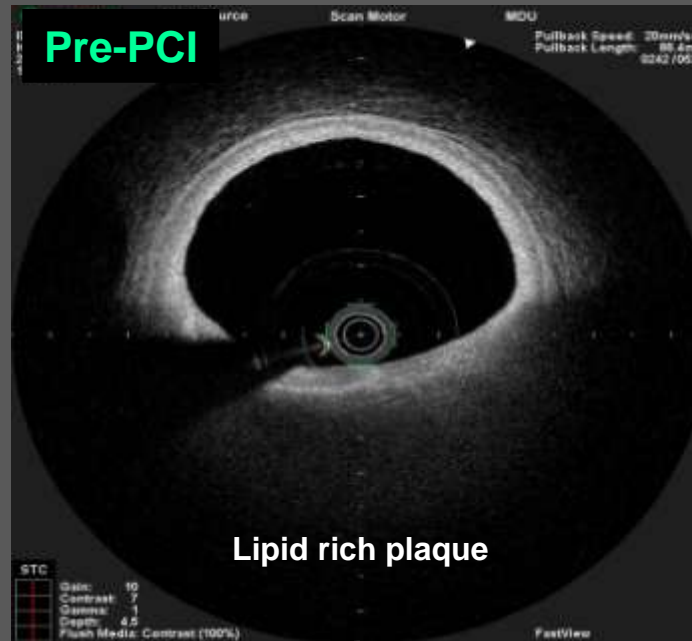


OFDI at reference site (How to select landing zone)

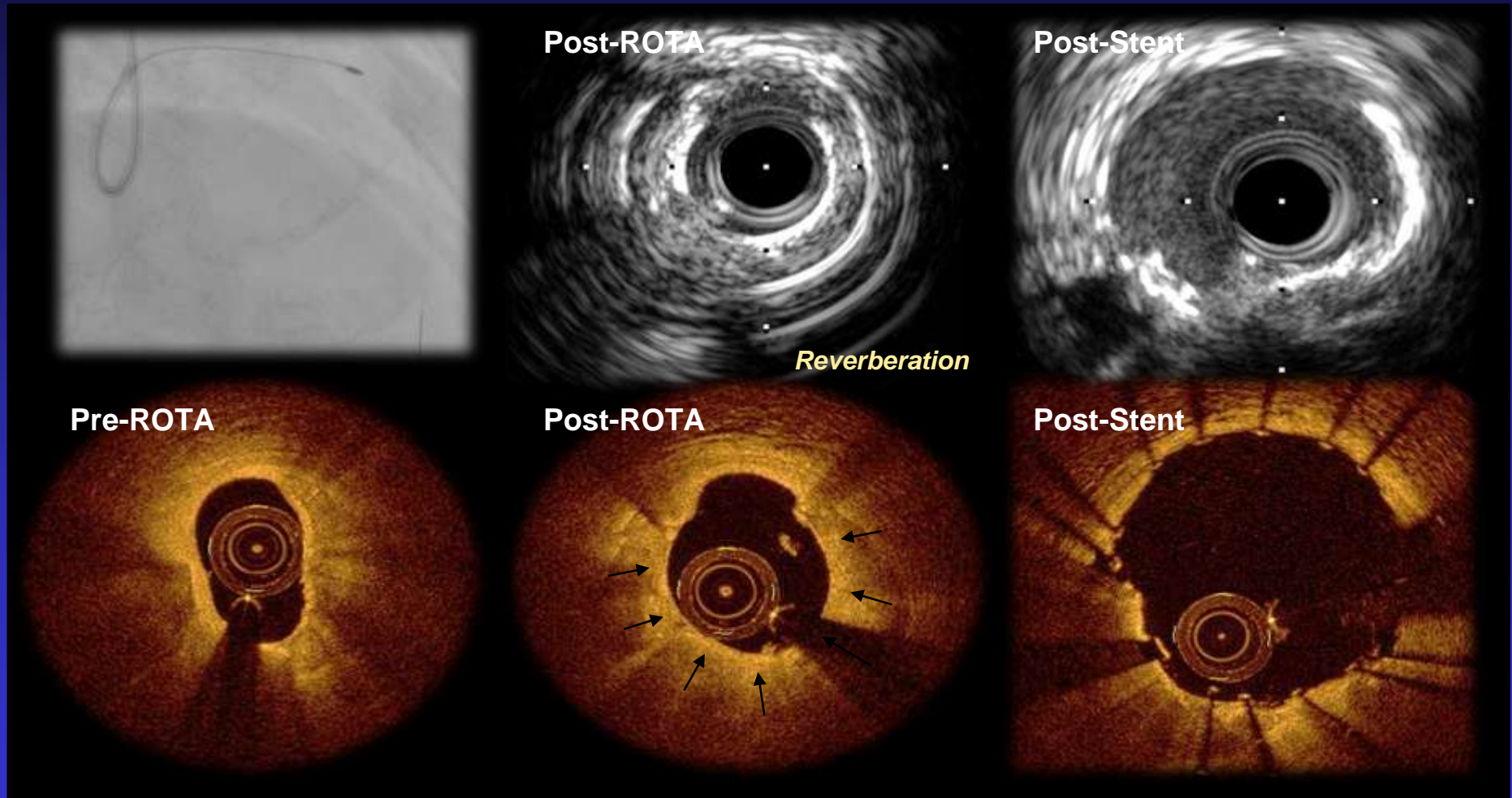
Distal reference



Proximal reference



Rotational atherectomy



Markedly calcified lesion

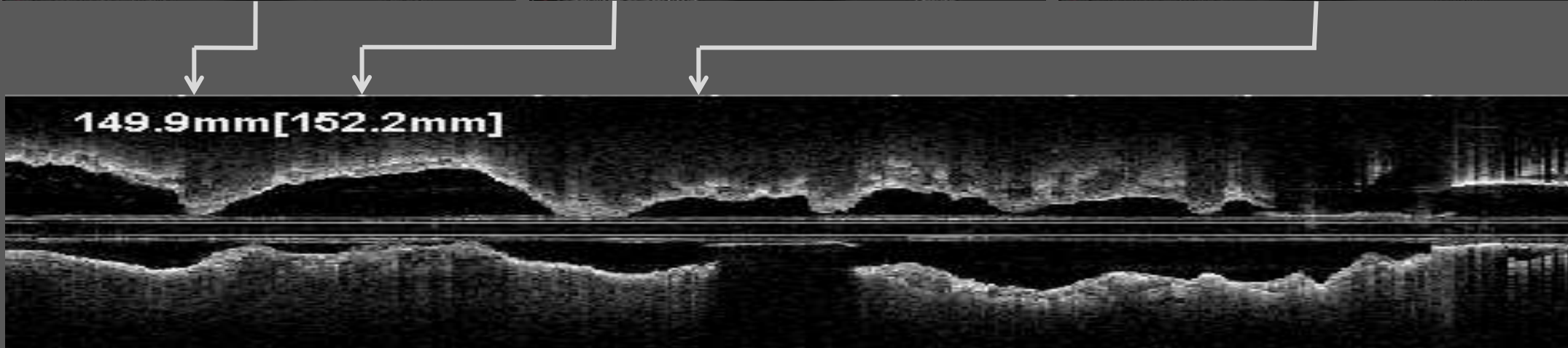
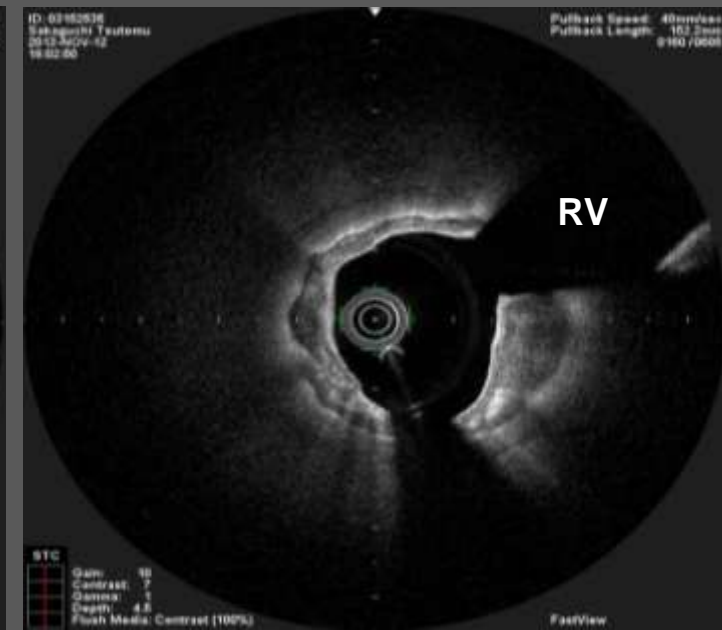
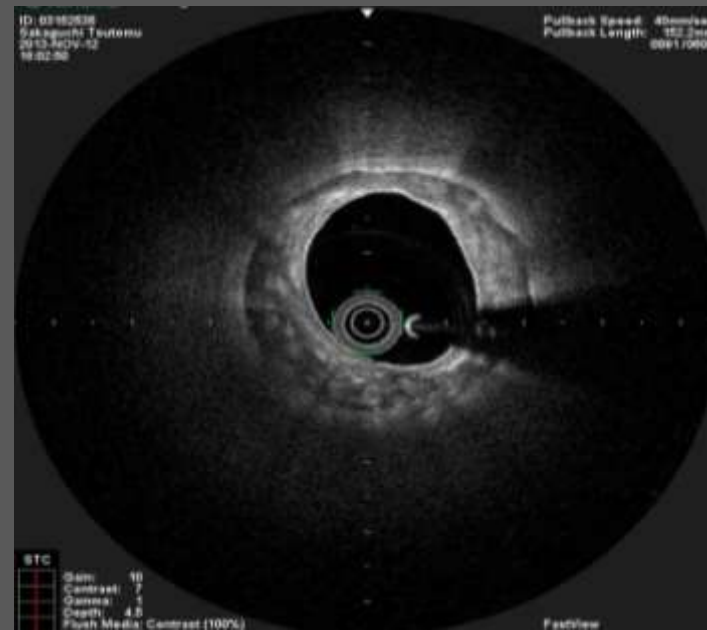
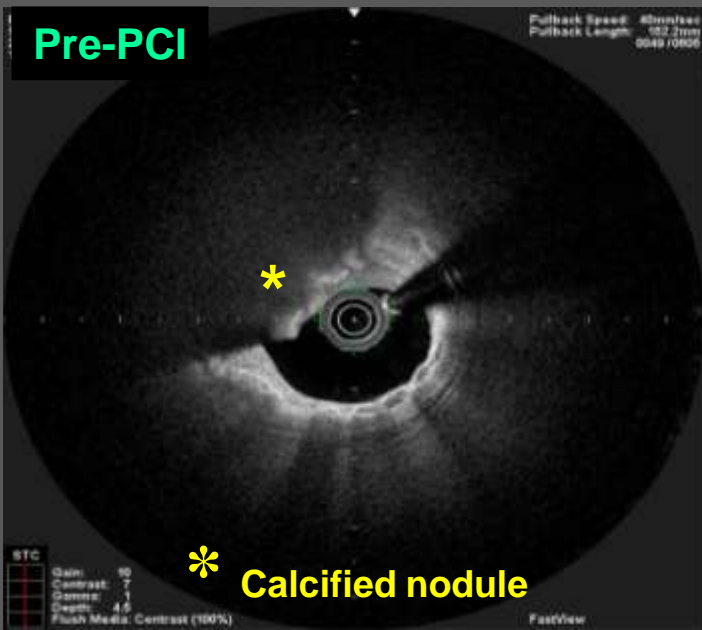


Pre-PCI FD-OCT (Markedly calcified lesion)

Minimum lumen area site

Severe calcification

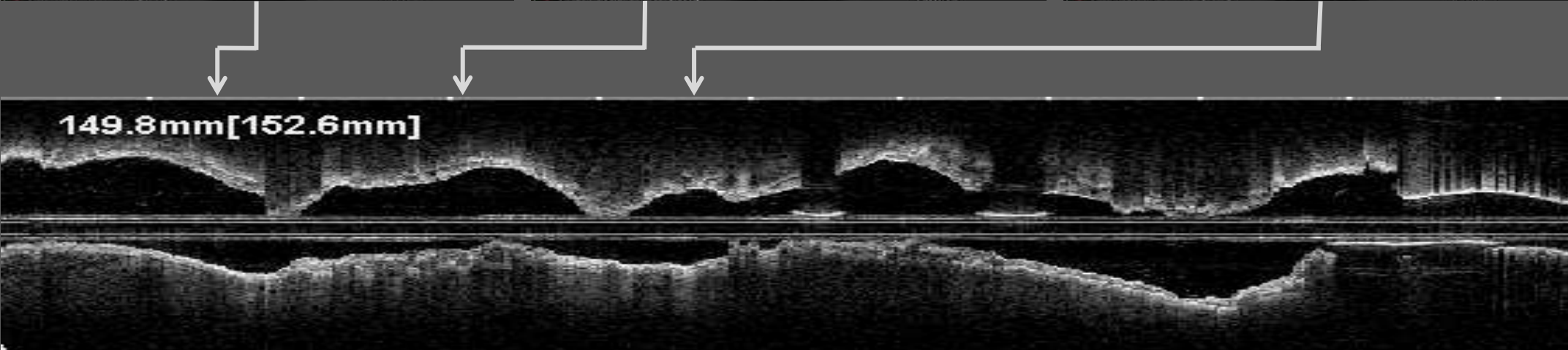
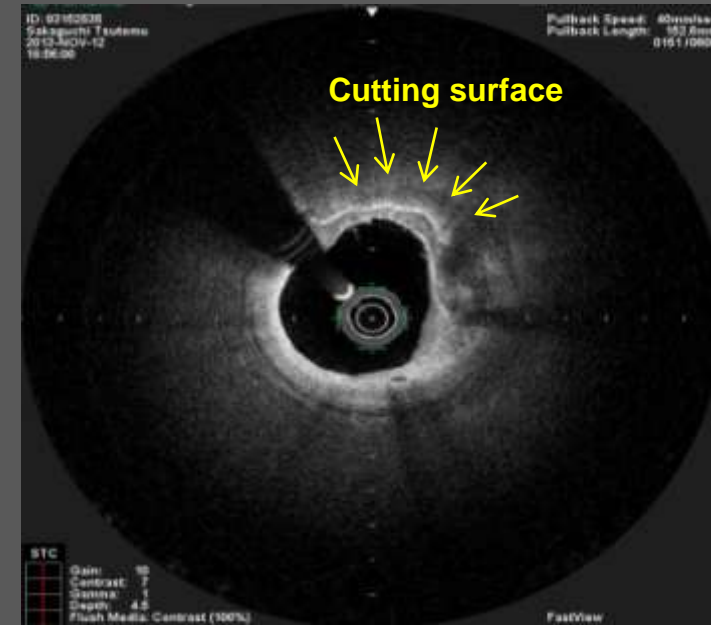
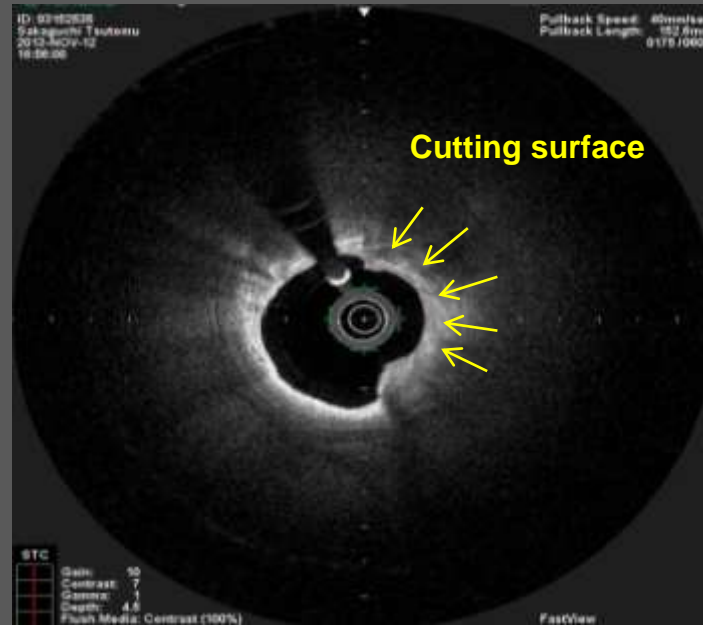
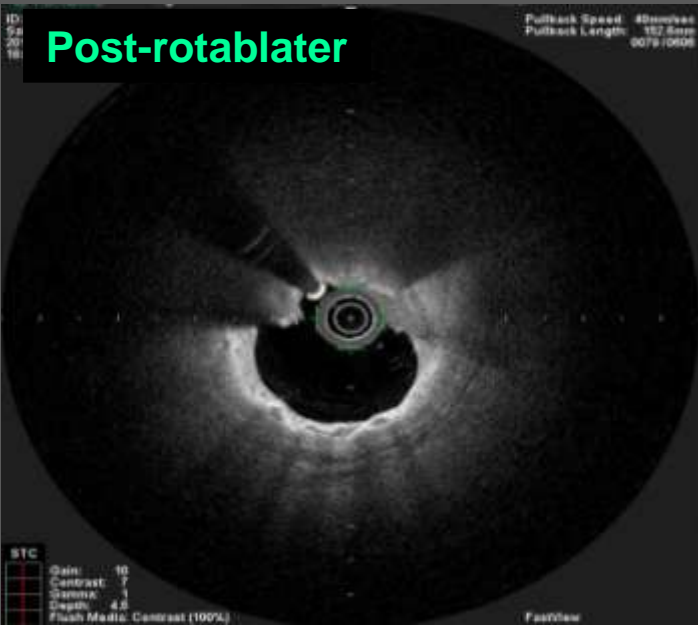
Severe calcification



Minimum lumen area site

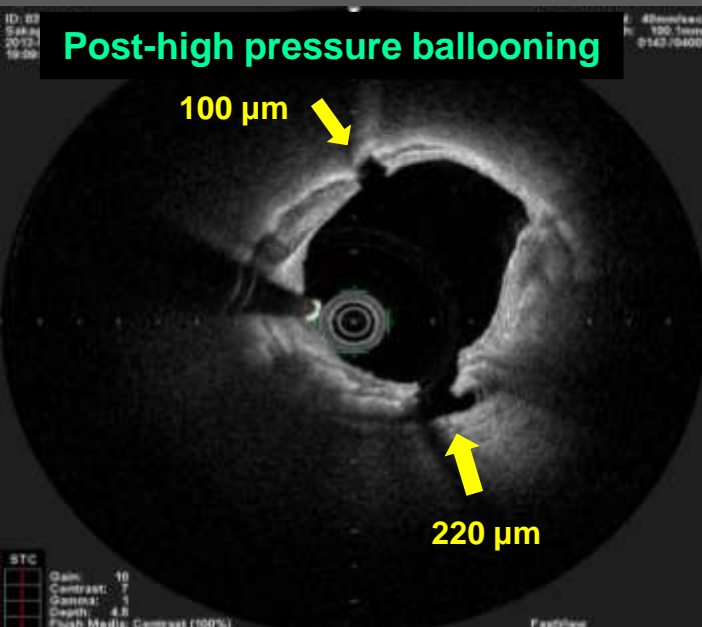
Calcification

Calcification

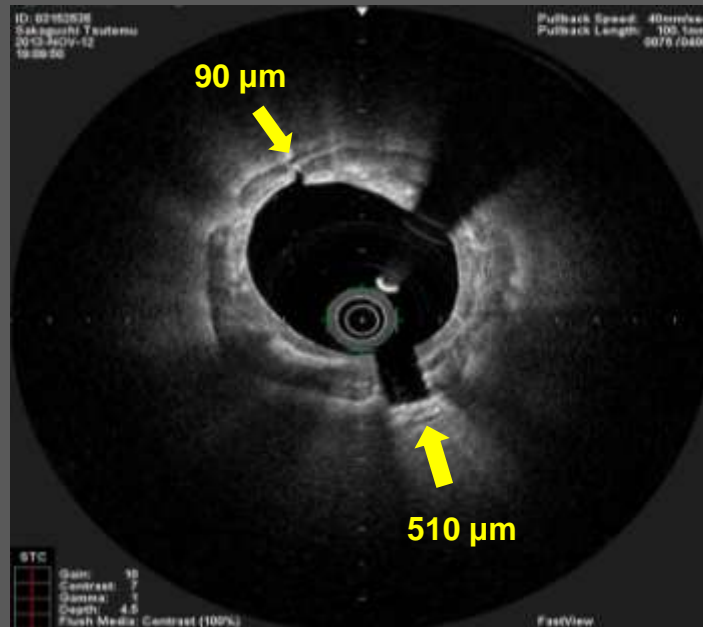


Broken calcium plate

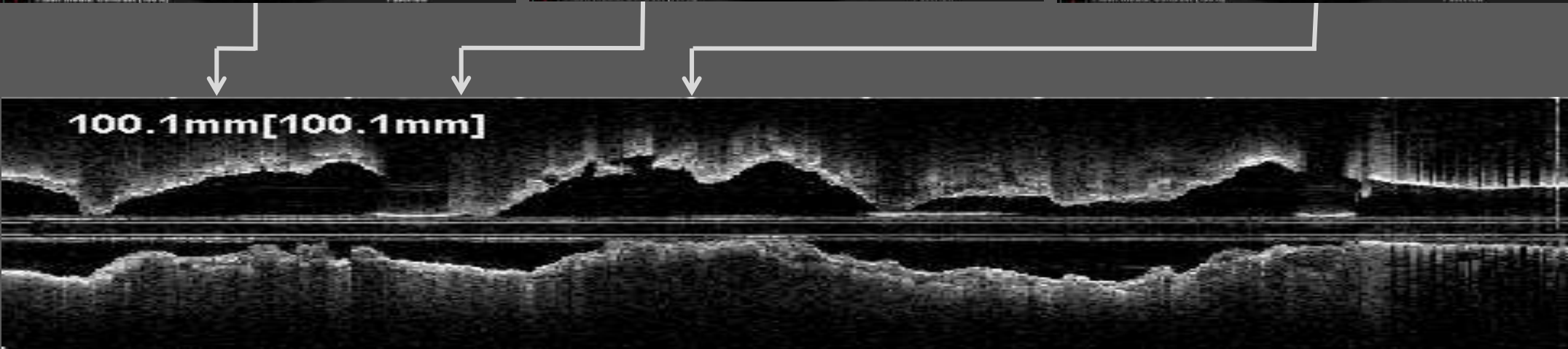
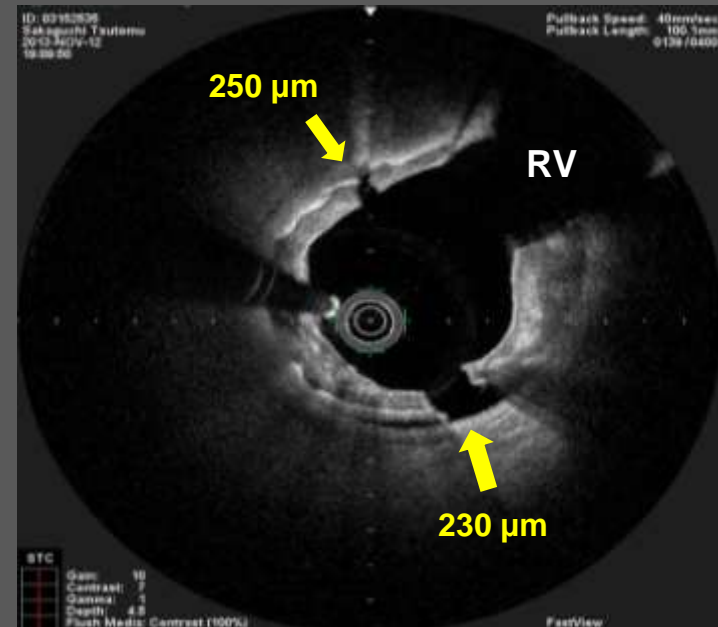
Post-high pressure ballooning



Broken calcium plate

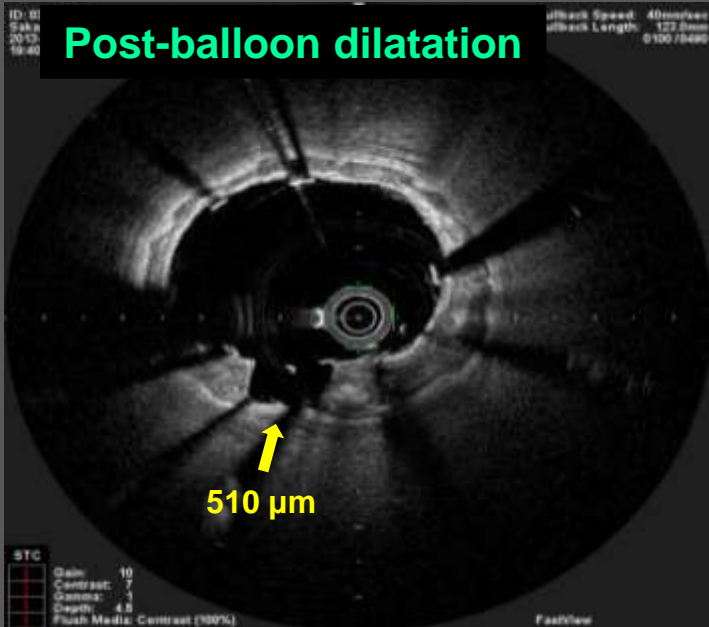


Broken calcium plate

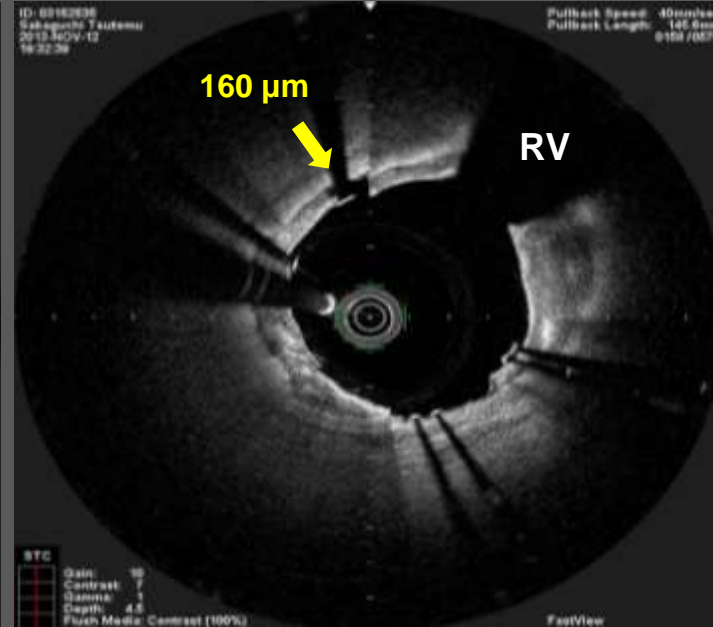


Broken calcium plate

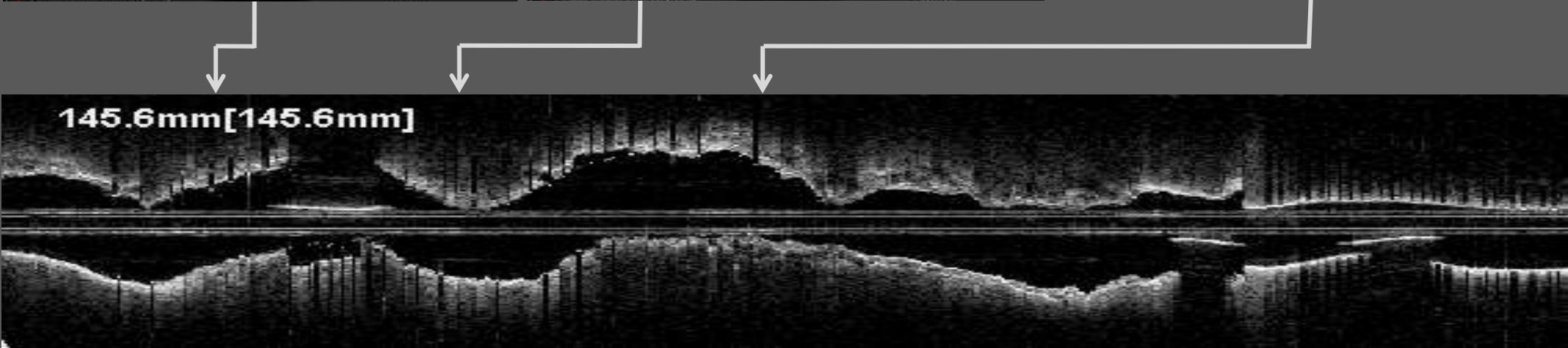
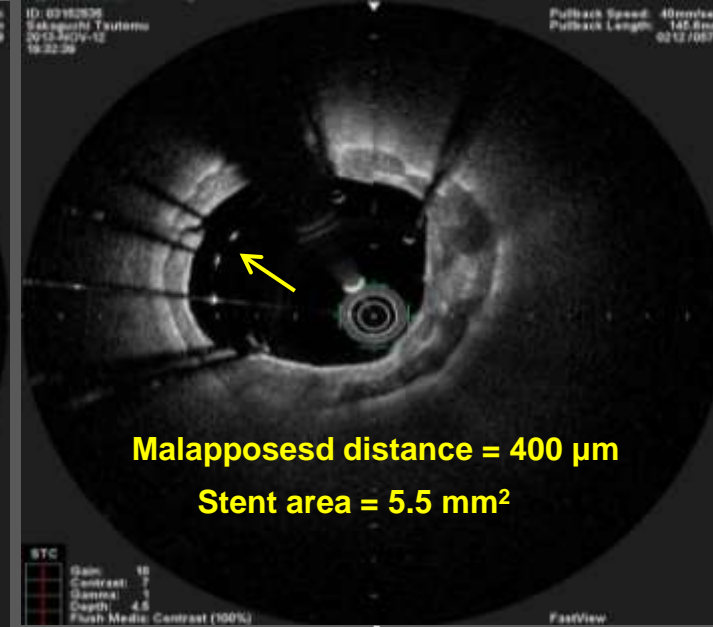
Post-balloon dilatation



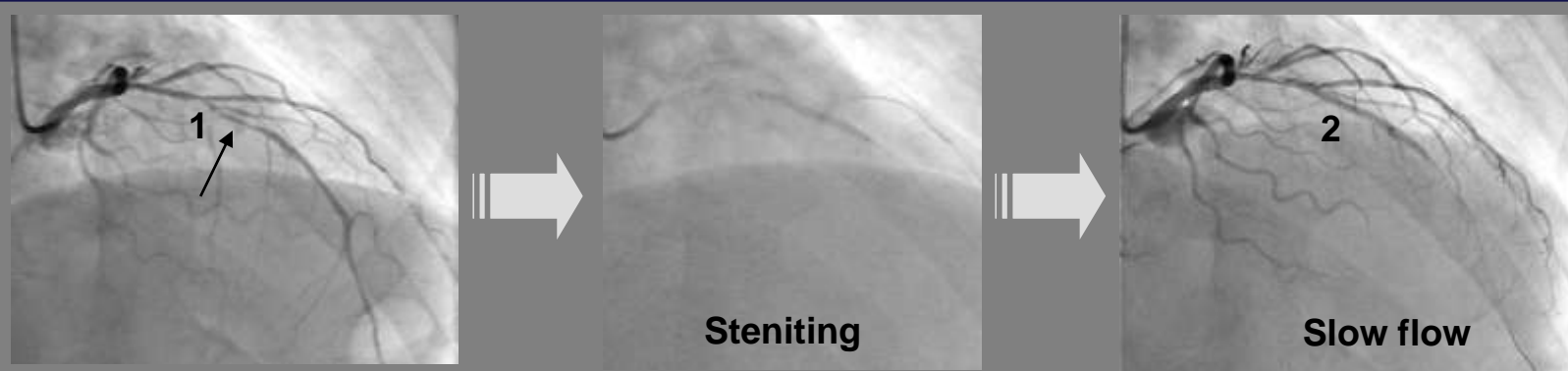
Broken calcium plate



Stent malapposition



Prediction of angiographic slow flow



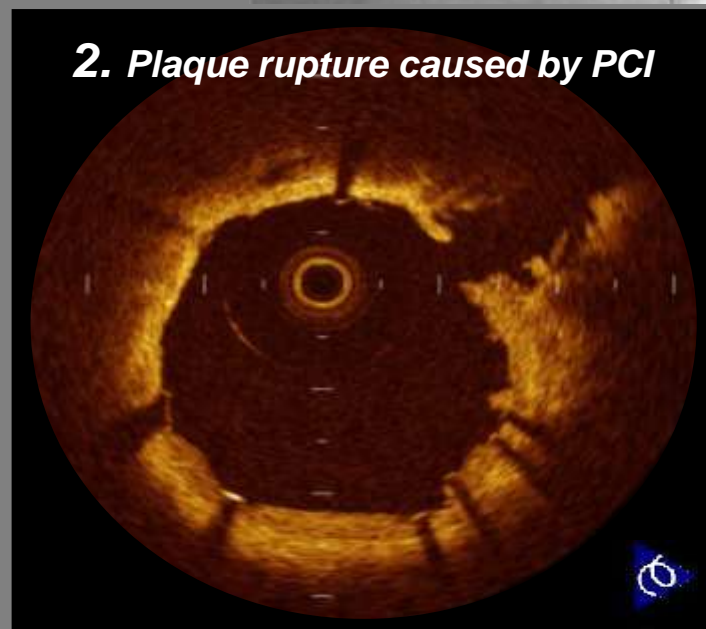
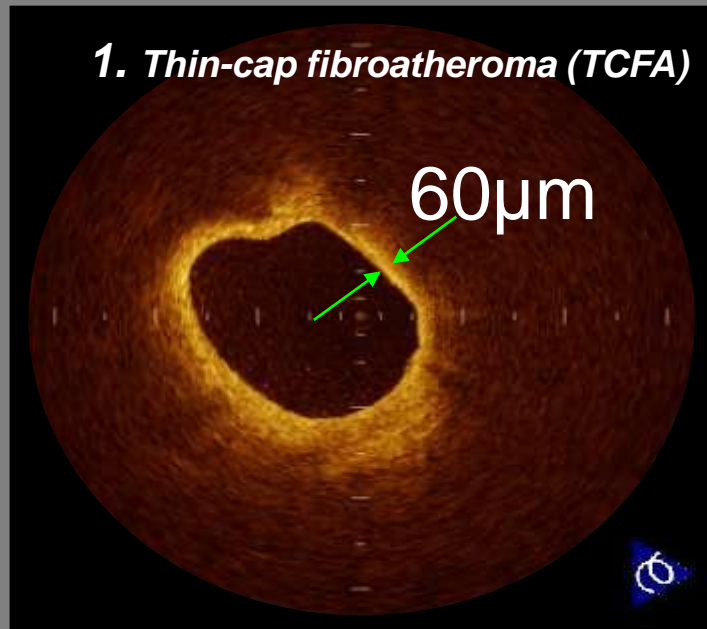
A 73-year-old male underwent PCI for the treatment of mid-LAD lesion (arrow).

In OCT image at pre-intervention, the culprit lesion presented lipid-rich plaque with thin-fibrous cap (TCFA).

After stenting, angiogram showed slow flow, and OCT disclosed plaque rupture behind stent.

TCFA is easy to be ruptured by PCI and has a high risk for coronary slow flow.

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Prediction of No-reflow Post-PCI

	No-reflow n=14	Reflow n=69	p-Value
Plaque rupture, %	71	48	0.053
Thrombus, %	79	80	0.567
TCFA, %	50	16	0.034
Lipid-arc, degree*	166	44	0.012

Tanaka A, Kubo T, Akasaka T et al. Eur Heart J 2009;30:1348-55

Prediction of Microvascular Obstruction

	OR	95% CI	P
ST-elevation myocardial infarction	48.05	2.85–809.11	0.007
TCFA at culprit	5.43	1.27–23.32	0.023
Thrombectomy	0.014	0.001–0.35	0.009
Diameter stenosis, %	1.1	1.02–1.19	0.011

Ozaki, Kubo, Akasaka et al. Circulation Img 2011;4:620-7

There is not enough data demonstrating the efficacy of distal protection during PCI.

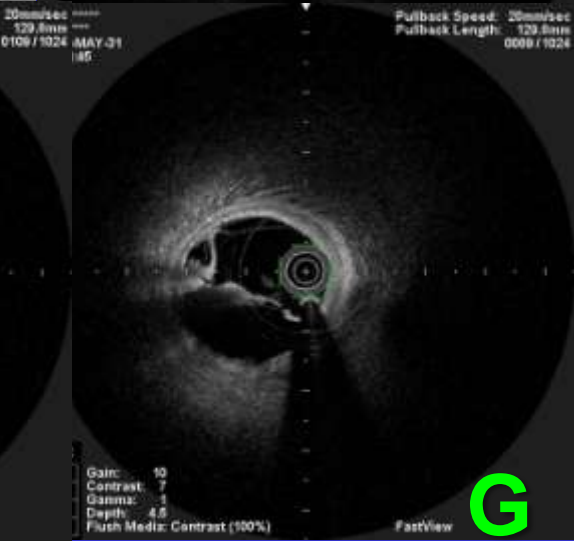
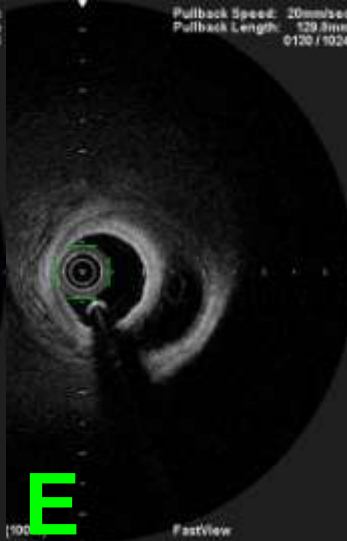
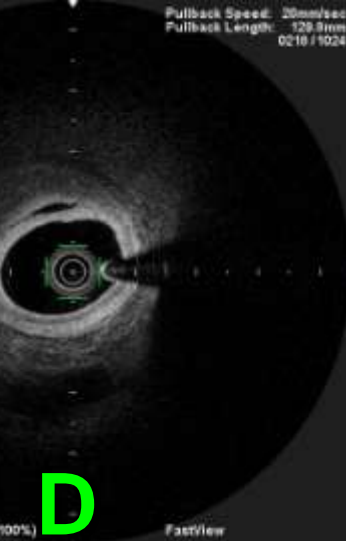
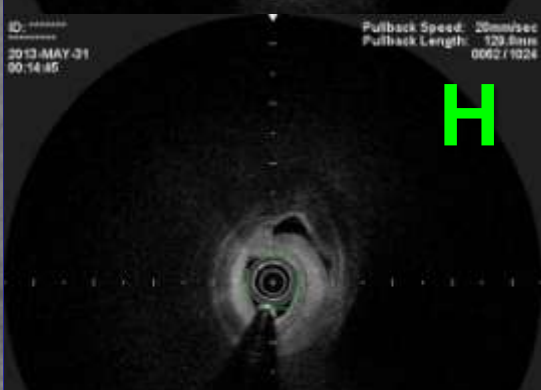
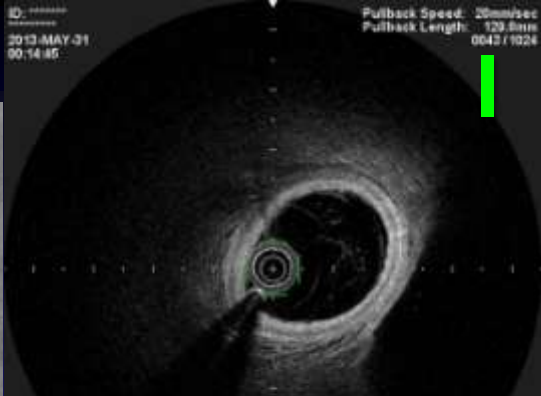
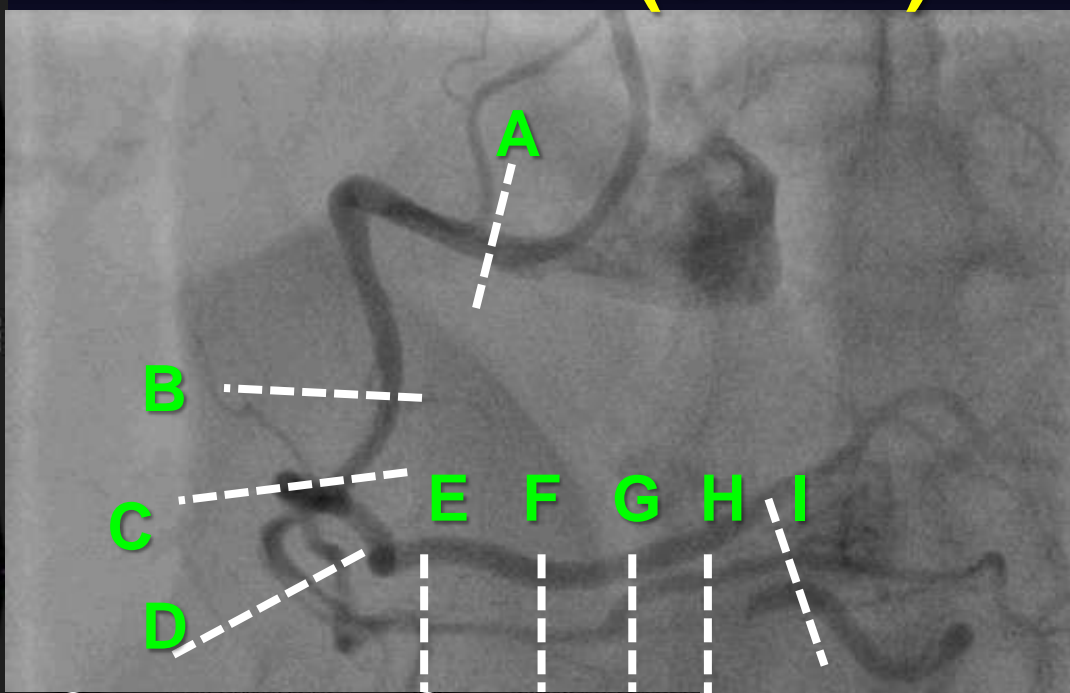
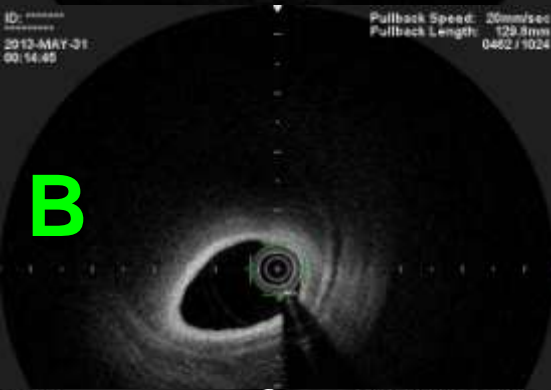
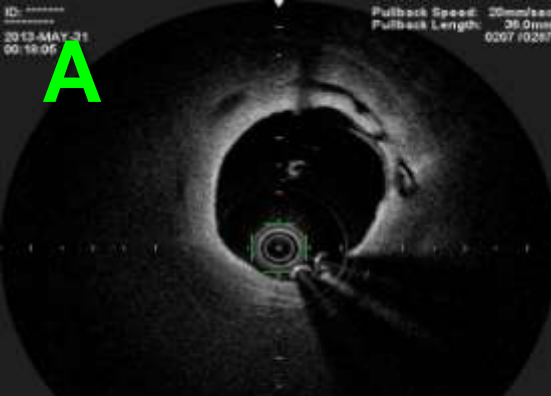
Wakayama Medical University



CAG (SCAD)



CAG & OCT (SCAD)



Diagnosis of Spontaneous Coronary Artery Dissection by Optical Coherence Tomography

Fernando Alfonso, MD, PHD, Manuel Paulo, MD, Nieves Gonzalo, MD, PHD, Jaime Dutary, MD, Pilar Jimenez-Quevedo, MD, PHD, Vera Lennie, MD, Javier Escaned, MD, PHD, Camino Bañuelos, MD, Rosana Hernandez, MD, PHD, Carlos Macaya, MD, PHD

Madrid, Spain

Objectives	This study sought to assess the diagnostic value of optical coherence tomography (OCT) in patients with suspected spontaneous coronary artery dissection (SCAD).
Background	SCAD is a rare but challenging clinical entity.
Methods	Following a prospective protocol, OCT was performed in 17 consecutive patients with a clinical and angiographic suspicion of SCD from a total of 5,002 patients undergoing coronary angiography. A conservative management strategy was followed.
Results	OCT ruled out the diagnosis of SCAD in 6 patients with coronary artery disease (atherosclerotic plaques and/or intracoronary thrombus). In 11 patients (age 48 ± 9 years, 9 female), OCT confirmed the presence of SCAD. A double-lumen or intramural hematoma image was visualized in all cases. However, only 3 patients presented an intimal "flap" on angiography. OCT readily identified the intimal rupture site ($n = 7$), the thickness ($348 \pm 84 \mu\text{m}$) and length ($31 \pm 9 \text{ mm}$) of the intimomedial membrane, the area of the true ($1.1 \pm 0.5 \text{ mm}^2$) and false lumen ($5.9 \pm 2.1 \text{ mm}^2$), the associated intramural hematoma ($n = 9$), and thrombi in the true or false lumens ($n = 11$). Most of these findings were angiographically silent. After stenting ($n = 4$), OCT disclosed adequate stent coverage, expansion, and apposition, but also residual intramural hematoma at the stented site (abluminal) and at the distal vessel.
Conclusions	<u>OCT provides unique insights in patients with SCAD that allow an early diagnosis and adequate management. Most of these findings are undetectable by angiography.</u> (J Am Coll Cardiol 2012;59:1073-9) © 2012 by the American College of Cardiology Foundation



How to use OCT-guided PCI

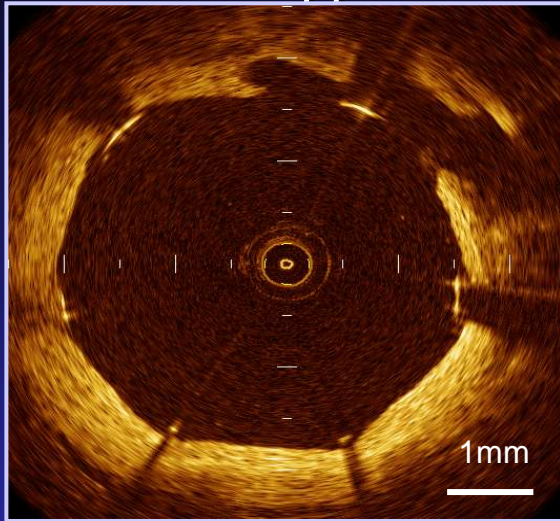
● Pre PCI Assessment

- Image acquisition is very fast and easy.
- Precise measurements might be possible automatically.
- Lesion morphology can be assessed in detail.
Easy to plan PCI strategies, easy to decide stent landing zone,
easy to identify unexpected lesions, etc.

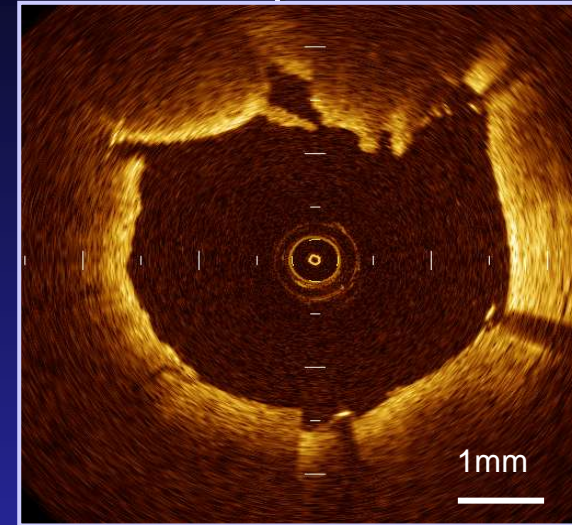


Inadequate lesion morphologies after stenting

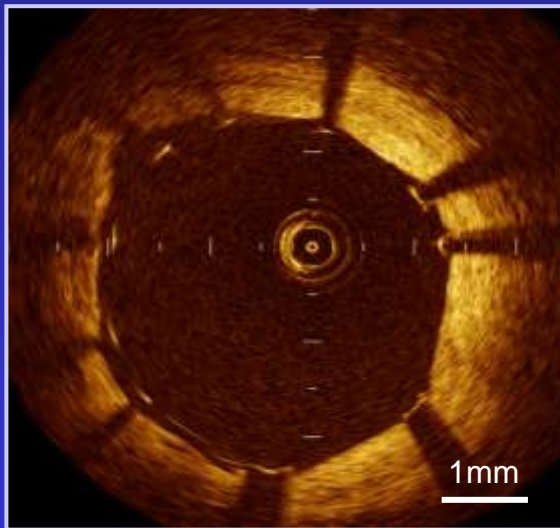
Stent malapposition



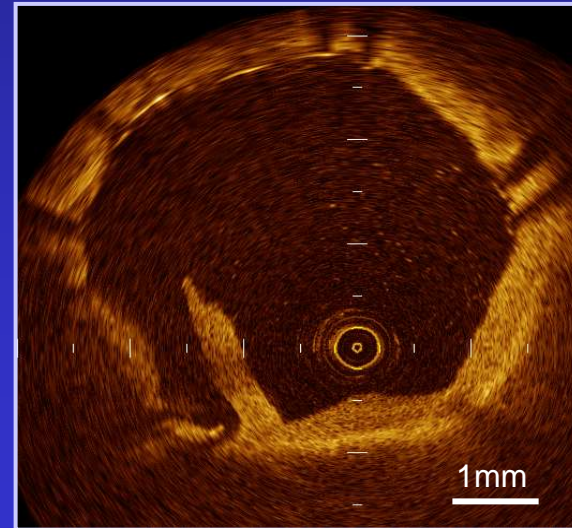
Tissue protrusion



Incomplete stent apposition

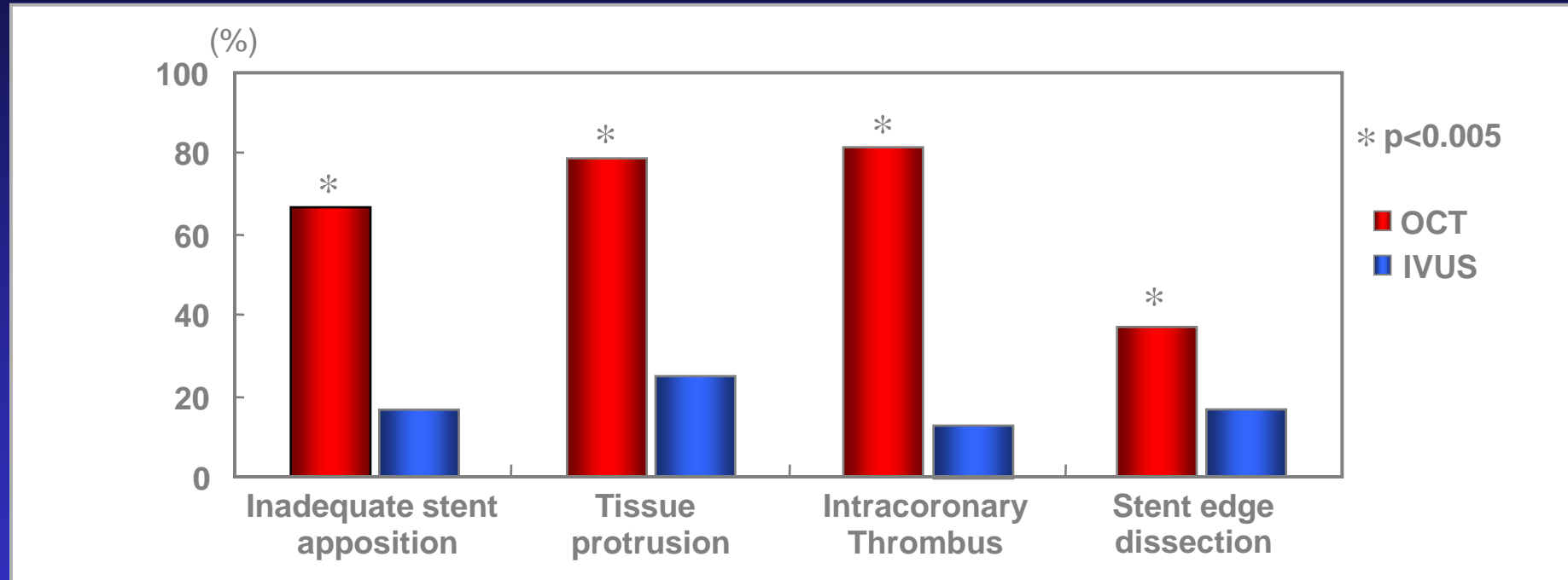


Stent edge dissection



Comparison of the ability for monitoring stent deployment between OCT and IVUS

55 patients were examined by OCT and IVUS to evaluate lesion morphologies after stent implantation.

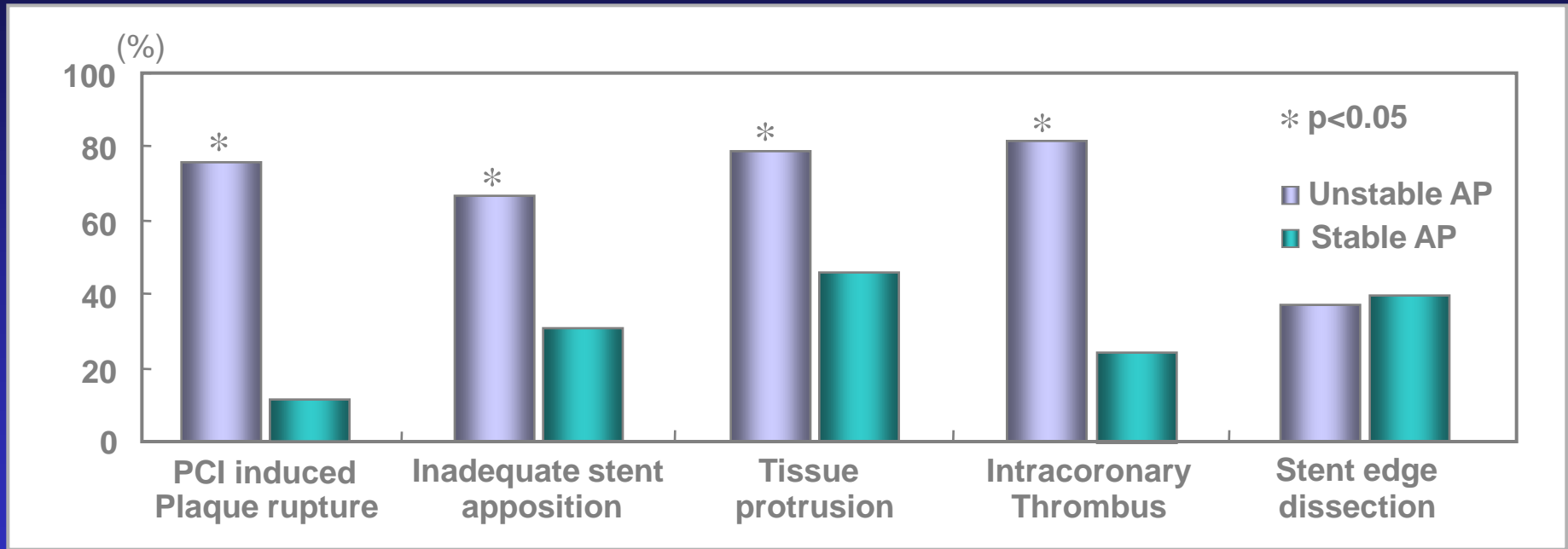


Conclusion: OCT can provide more detailed morphological information after stenting than IVUS.



Vascular response after stent implantation between unstable and stable AP

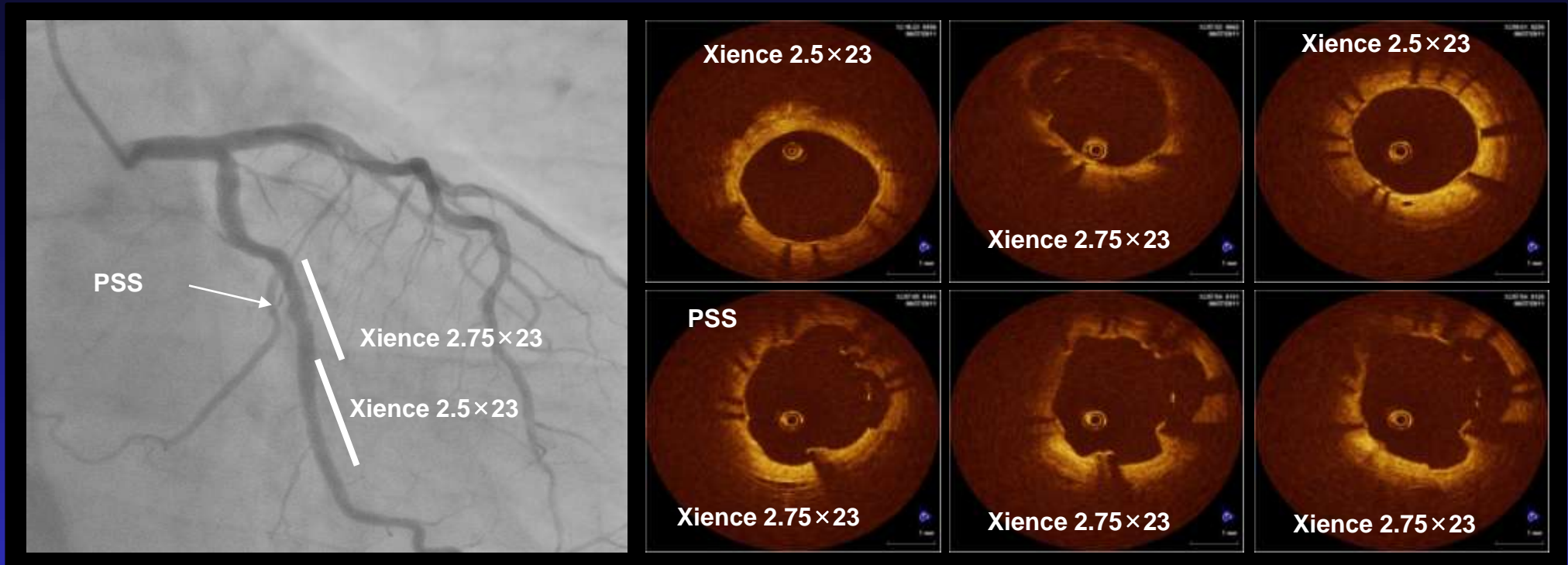
24 unstable and 31 stable AP patients were examined by OCT to evaluate lesion morphologies after stent implantation.



Conclusion: The inadequate lesion morphologies after stenting were observed more frequently in unstable AP patients.



PSS, #13 CTO (2010/10/26)

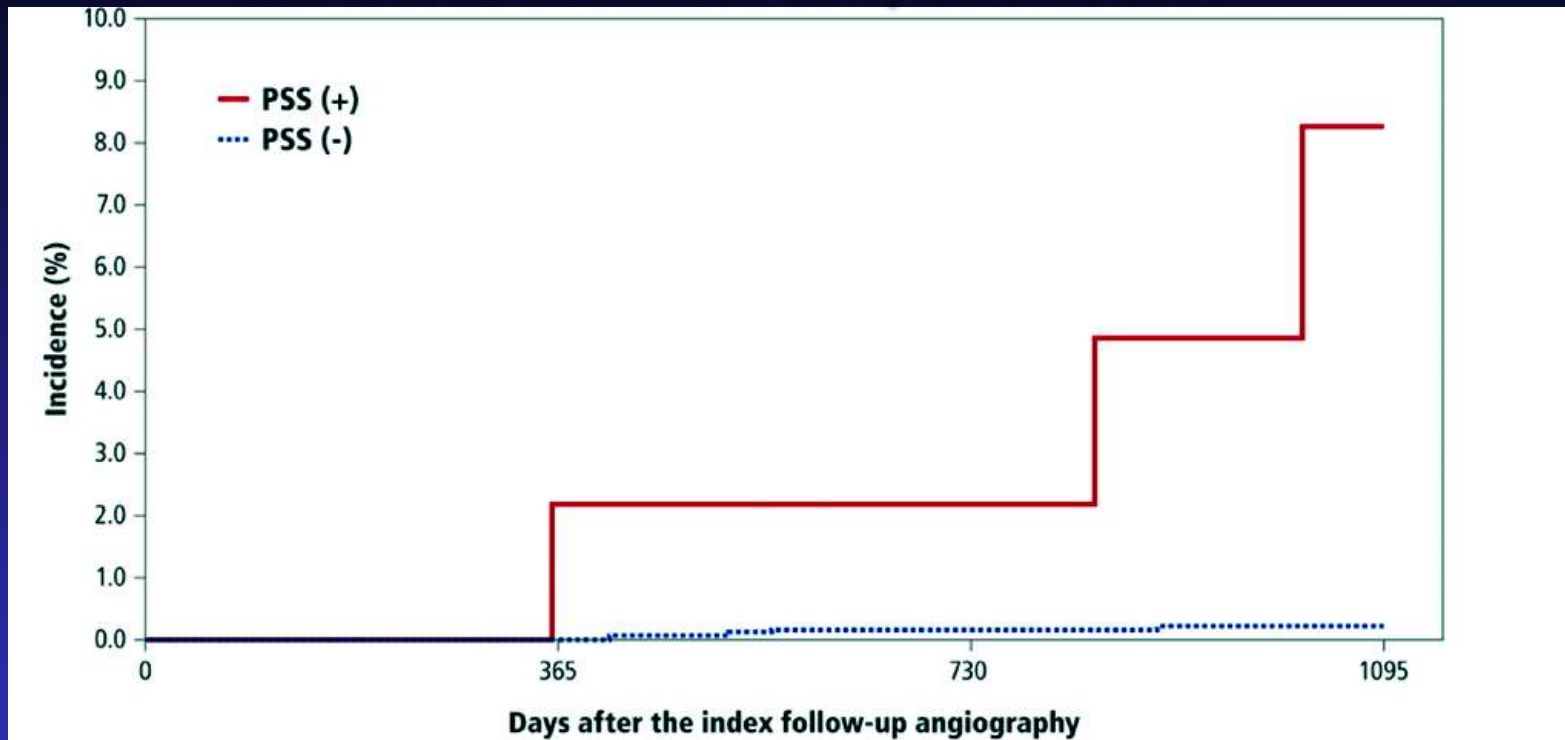


PSS by CAG is demonstrated as mal-apposition by OCT.

Persistent incomplete apposition & late acquired mal-apposition should be considered as the cause of mal-apposition in late phase.

Cumulative incidence of ST after the index follow-up CAG

Imai M et al. Circulation 2011;123:2382-2391

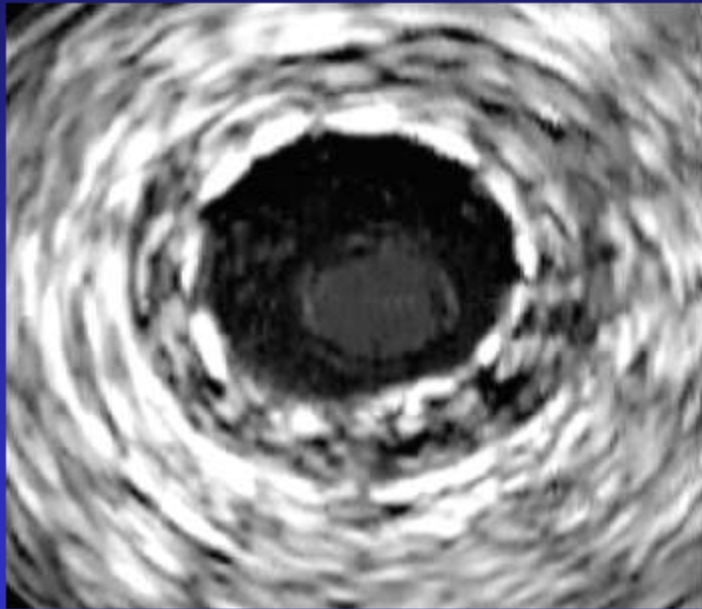


Days	0	365	730	1095
PSS (+) N of lesions at risk	51	46	40	26
N of lesions with events	0	1	1	3
Cumulative incidence	0%	2.1%	2.1%	8.2%
PSS (-) N of lesions at risk	2761	2532	1847	580
N of lesions with events	0	0	3	4
Cumulative incidence	0%	0%	0.13%	0.2%

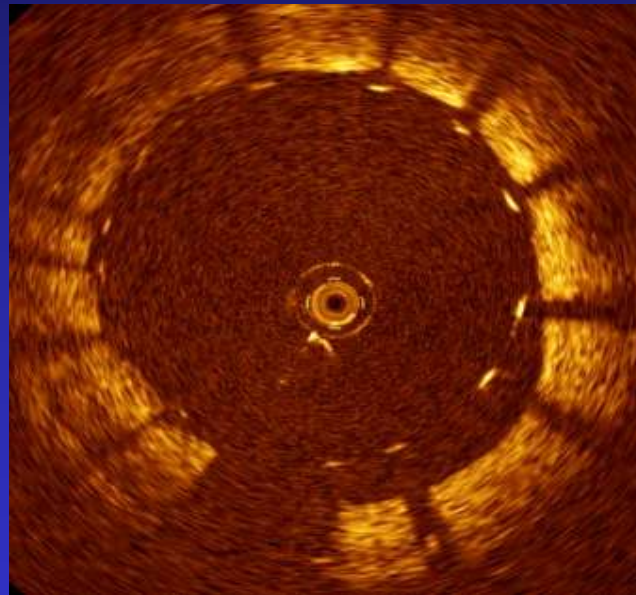


Incomplete stent apposition

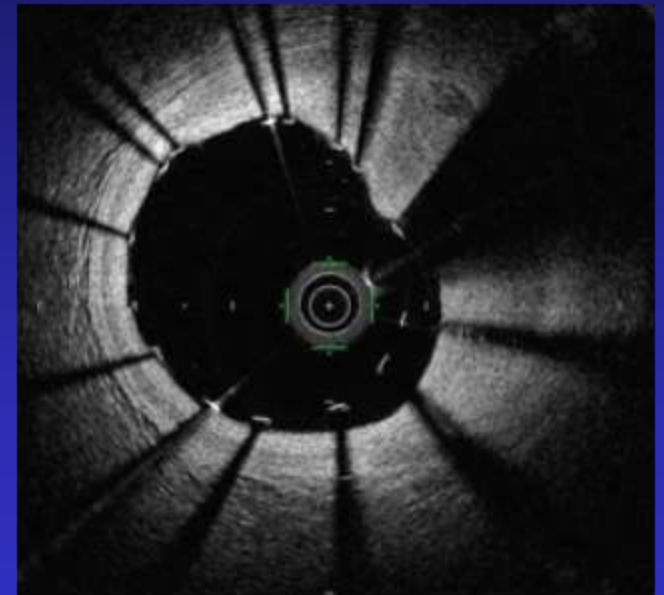
Incomplete stent apposition can be easily demonstrated by OFDI/FD-OCT, even if it is not identified by IVUS.



IVUS



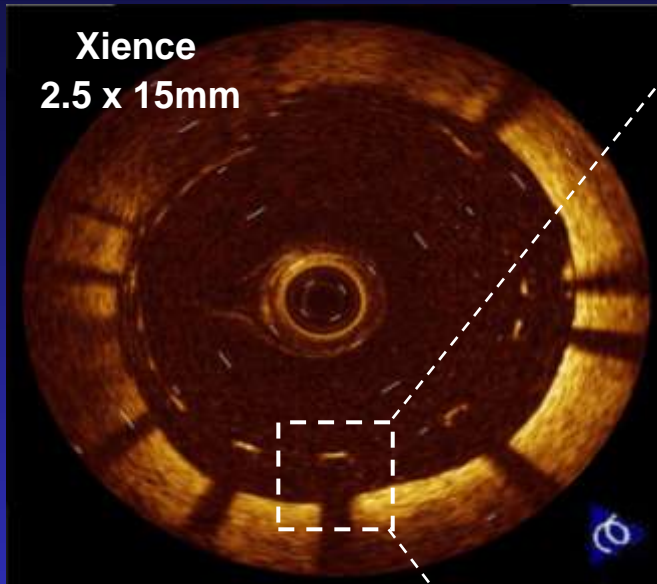
FD-OCT



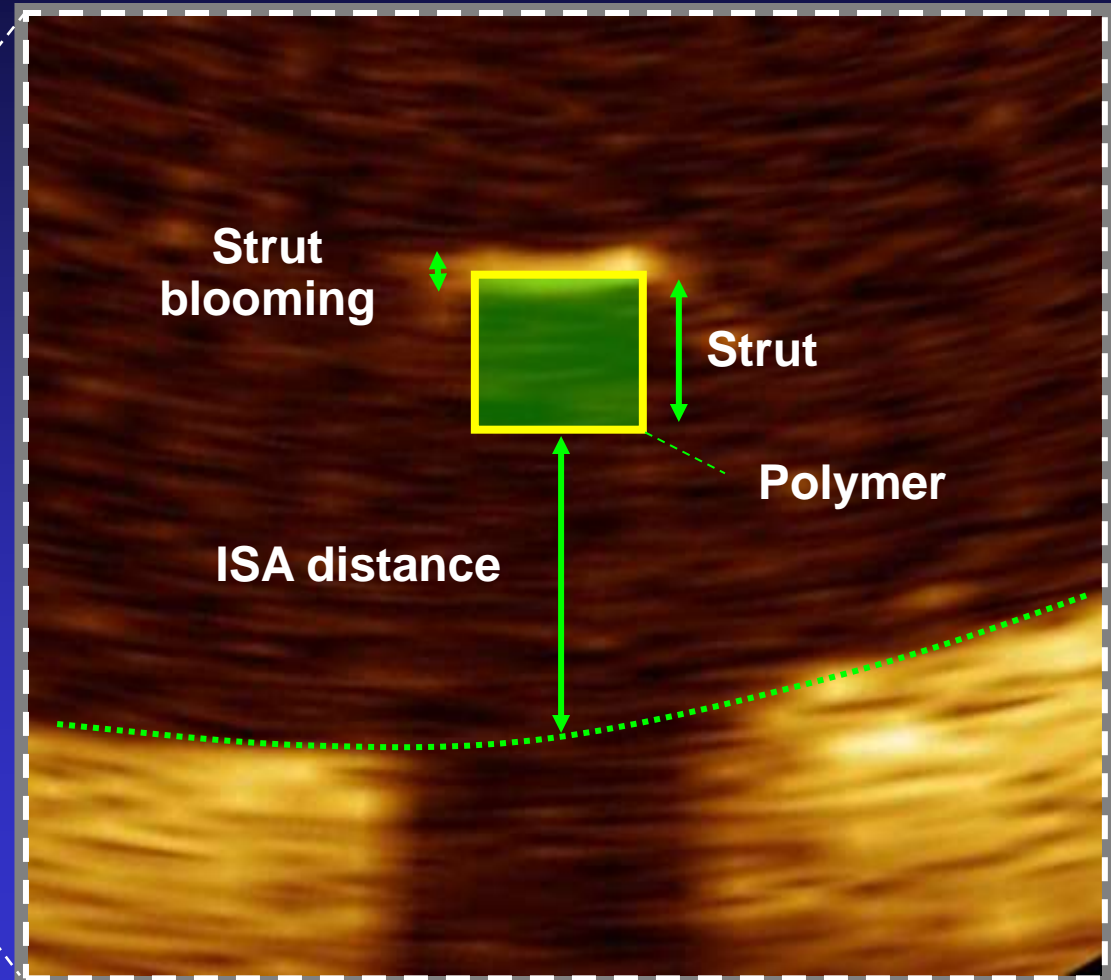
OFDI



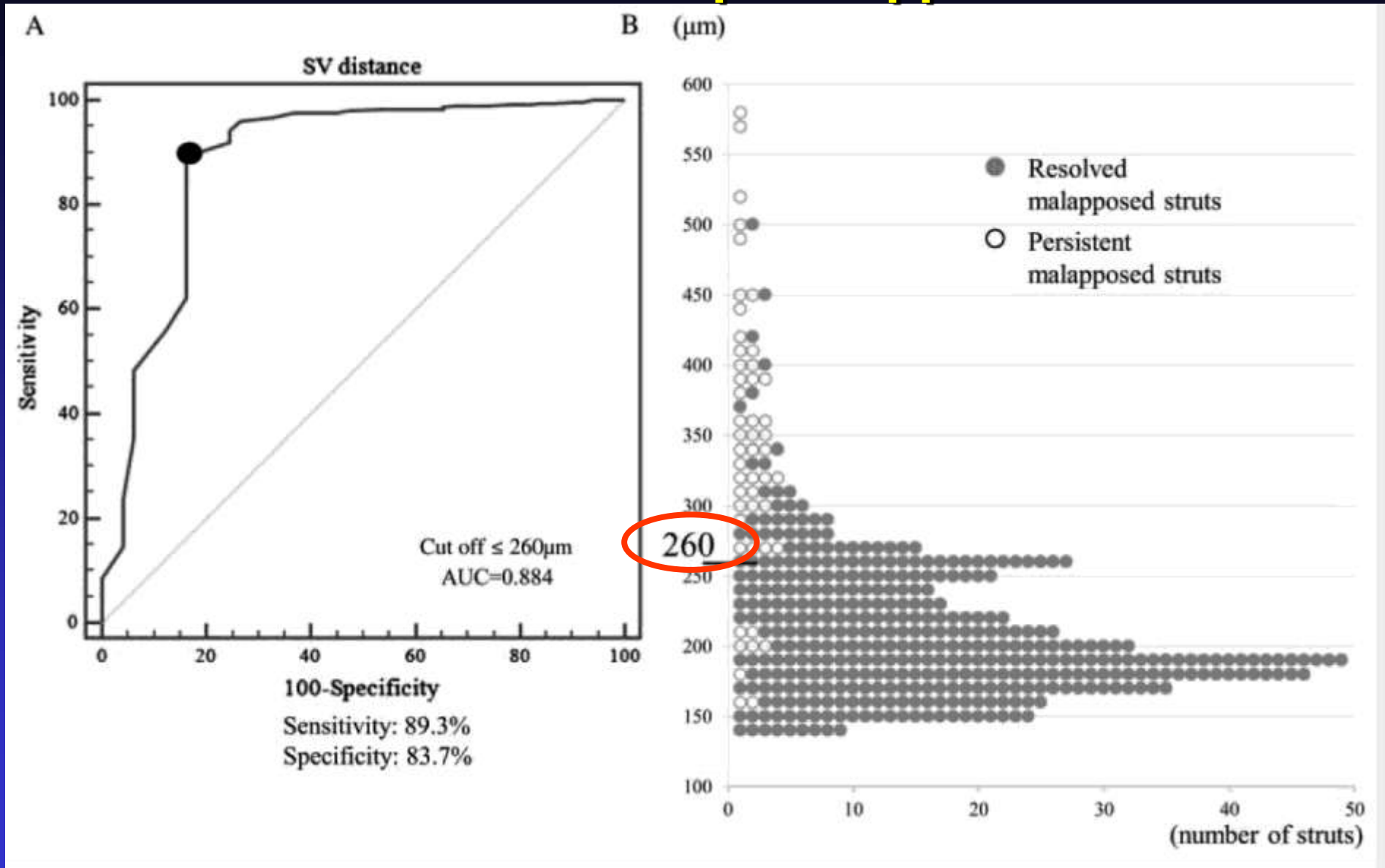
Definition of incomplete stent apposition (ISA)



ISA was defined as a ISA distance of $>100 \mu\text{m}$ in EES and $>170 \mu\text{m}$ in SES.

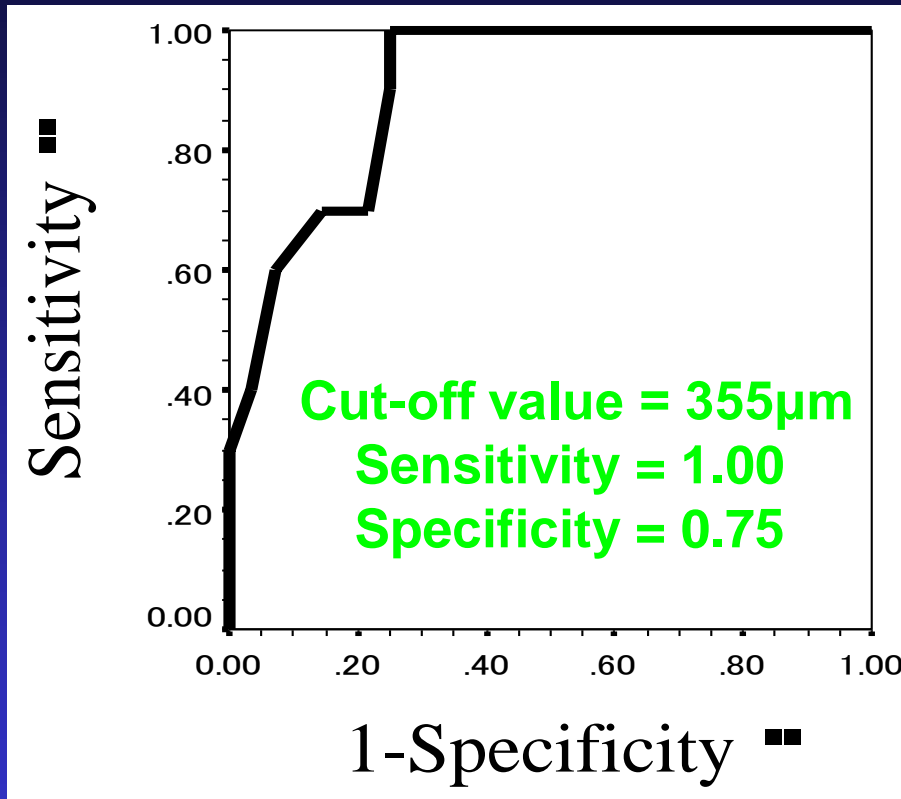


Persistent incomplete apposition



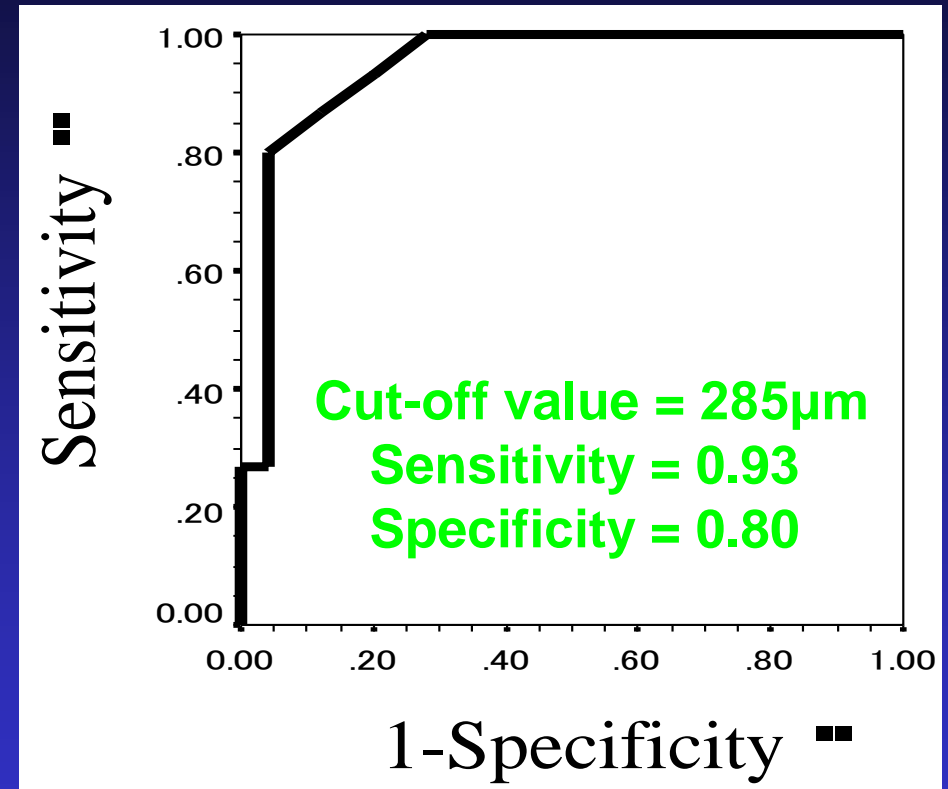
ROC curve analysis of maximum ISA distance for predicting persistent ISA

EES



ROC curve analysis identified a maximum ISA distance of **EES > 355 μ m** with as separating persistent from resolved ISA (sensitivity 100%, specificity 75%, area under the curve = 0.905; 95%CI, 0.812 to 0.999).

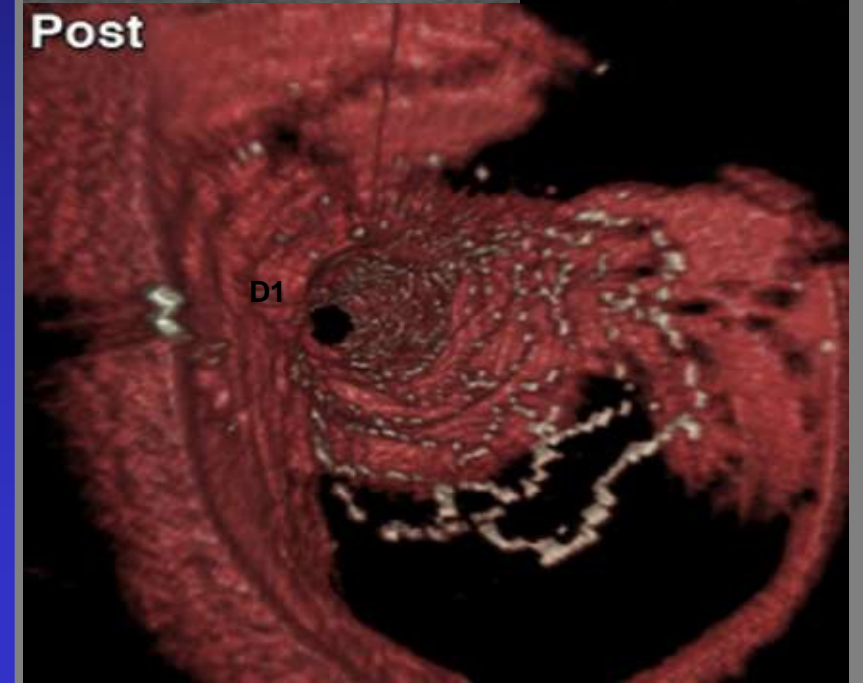
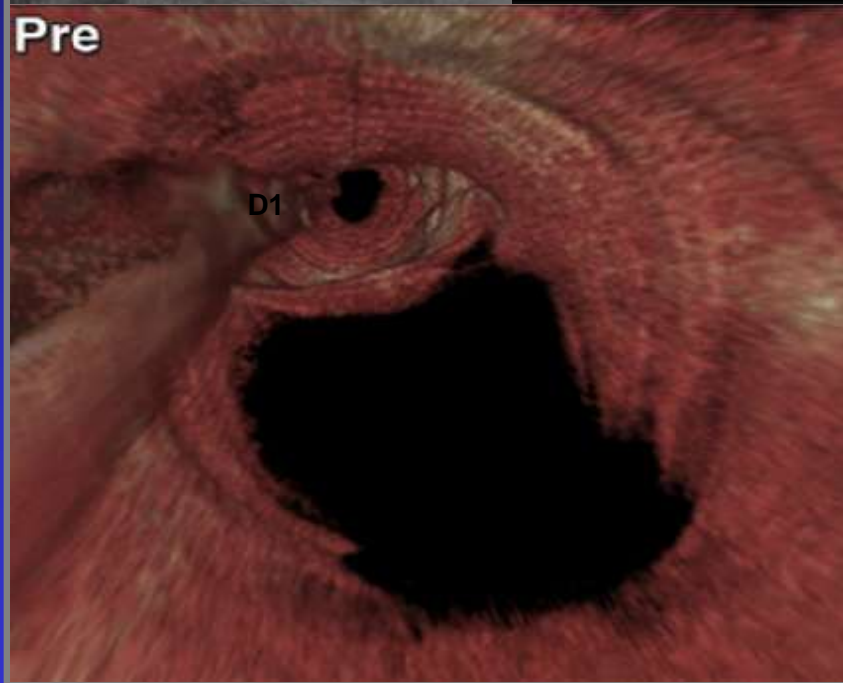
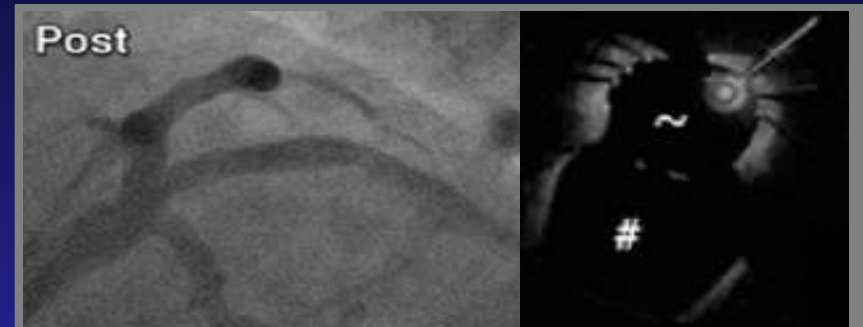
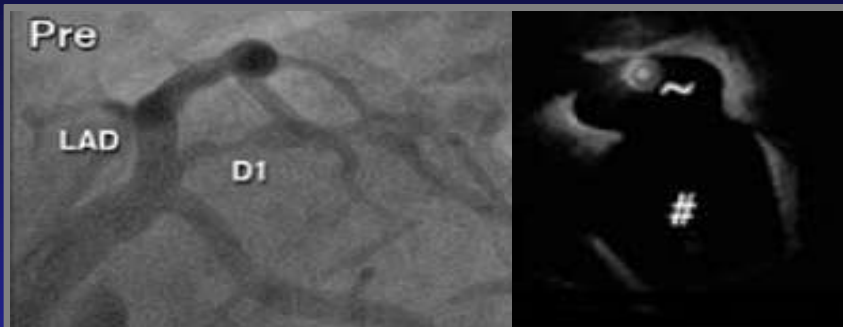
SES



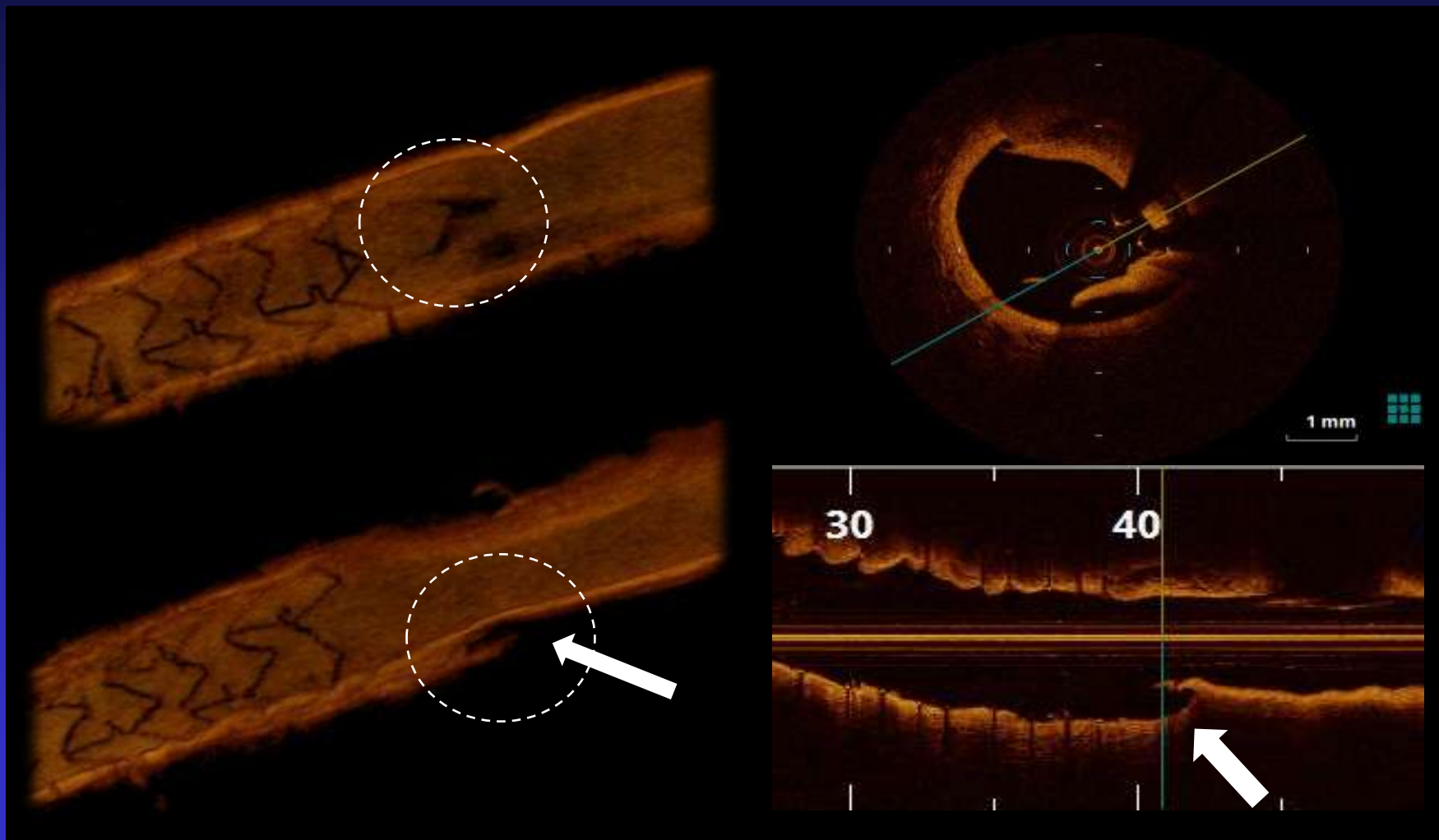
ROC curve analysis identified a maximum ISA distance of **SES > 285 μ m** with as separating persistent from resolved ISA (sensitivity 93%, specificity 80%, area under the curve = 0.947; 95%CI, 0.878 to 1.015).



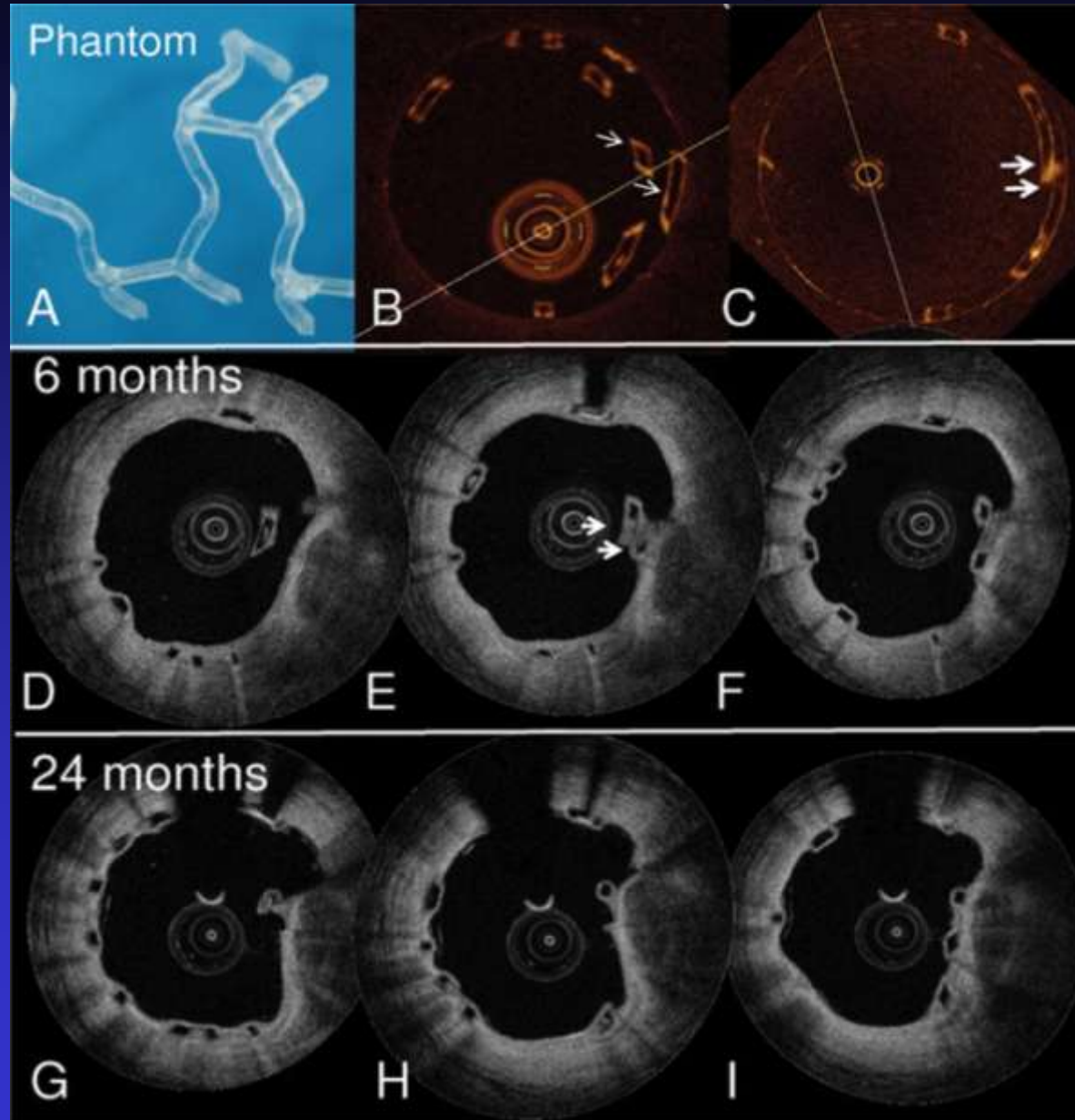
“Overhanging” struts of the D1 stent into the LAD orifice



SAP, #6 99% Xience 2.75x18, Dissection



Assessment of BVS by OFDI

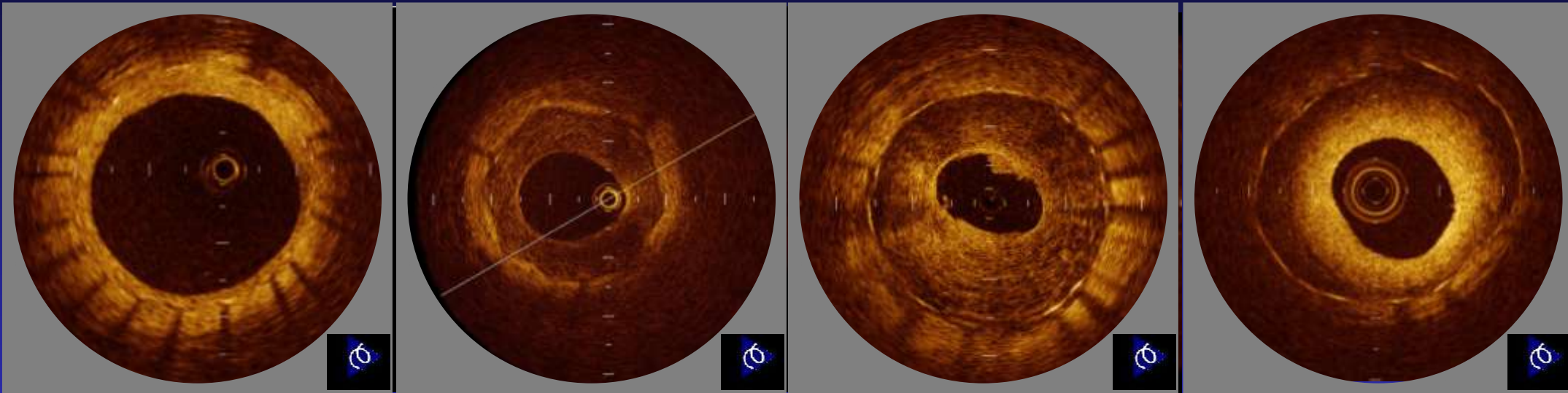


Neointimal tissue characterization by OCT

Homogeneous

Heterogeneous

Layered



Restenotic tissue has uniform optical properties and does not show focal variations in backscattering pattern.

Restenotic tissue has focally changing optical properties and shows various backscattering patterns.

Restenotic tissue consists of concentric layers with different optical properties: an adluminal high scattering layer and adluminal low scattering layer.

No data showing the relation between OCT-findings & histology in detail, and furthermore, there is no recommendation of treatment according to OCT finding.



Association between tissue characteristics evaluated with optical coherence tomography and mid-term results after paclitaxel-coated balloon dilatation for in-stent restenosis lesions: a comparison with plain old balloon angioplasty

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Aims

Morphological assessment of neointimal tissue using optical coherence tomography (OCT) is important for clarifying the pathophysiology of in-stent restenosis (ISR) lesions. The aim of this study was to determine the impact of OCT findings on recurrence of ISR after paclitaxel-coated balloon (PCB) dilatation compared with plain old balloon angioplasty (POBA).

Methods and results

Between July 2008 and May 2012, we performed percutaneous coronary intervention for 214 ISR lesions using POBA + PCB (146 lesions, PCB group) or POBA only (68 lesions, POBA group). Morphological assessment of neointimal tissue using OCT, including assessment of restenotic tissue structure and restenotic tissue backscatter, was performed. We examined the association between lesion morphologies and mid-term (6–8 months) results including ISR and target lesion revascularization (TLR) rates. Both ISR and TLR rates of lesions with a homogeneous structure were significantly lower in the PCB group than those in the POBA group (ISR: 20.0 vs. 55.6%, $P = 0.002$, TLR: 12.7 vs. 37.0%, $P = 0.019$), but there was no difference between the two groups in ISR and TLR rates of lesions with a heterogeneous or layered structure. Both ISR and TLR rates of lesions with high backscatter were significantly lower in the PCB group than those in the POBA group (ISR: 19.8 vs. 52.5%, $P < 0.001$, TLR: 13.6 vs. 42.5%, $P = 0.001$), but there was no difference between the two groups in ISR and TLR rates of lesions with low backscatter.

Conclusion

Morphological assessment of ISR tissue using OCT might be useful for identifying ISR lesions favourable for PCB dilatation.

Keywords

optical coherence tomography • in-stent restenosis lesion • paclitaxel-coated balloon

Association between restenotic tissue morphology and acute/mid-term results

	Tissue structure								
	Homogenous type			Heterogenous type			Layered type		
	PCB (n=55)	POBA (n=27)	P value	PCB (n=20)	POBA (n=8)	P value	PCB (n=71)	POBA (n=33)	P value
Acute gain mm	1.14±0.53	0.90±0.56	0.060	1.25±0.58	1.21±0.38	0.885	1.20±0.58	1.14±0.60	0.597
Late loss mm	0.25±0.50	0.70±0.58	0.000	0.45±0.72	0.84±0.85	0.234	0.23±0.60	0.61±0.69	0.005
Net gain mm	0.90±0.61	0.20±0.67	0.000	0.80±0.69	0.38±0.98	0.208	0.98±0.73	0.53±0.63	0.003
ISR n (%)	11 (20.0)	15 (55.6)	0.002	7 (35.0)	3 (37.5)	1.000	16 (22.5)	13 (39.4)	0.100
TLR n (%)	7 (12.7)	10 (37.0)	0.019	5 (25.0)	3 (37.5)	0.651	14 (19.7)	12 (36.4)	0.089

Acute gain = (post-procedural – pre-procedural) MLD

Late loss = (post-procedural - follow-up) MLD

Net gain = (follow-up - pre-procedural) MLD



Association between restenotic tissue morphology and acute/mid-term results

Tissue backscatter	High backscatter			Low backscatter		
	PCB (n=81)	POBA (n=40)	P value	PCB (n=65)	POBA (n=28)	P value
Acute gain mm	1.12±0.50	0.97±0.58	0.139	1.26±0.62	1.17±0.54	0.476
Late loss mm	0.23±0.51	0.73±0.70	0.000	0.31±0.66	0.59±0.62	0.059
Net gain mm	0.90±0.61	0.25±0.67	0.000	0.96±0.76	0.58±0.70	0.027
ISR n (%)	16 (19.8)	21 (52.5)	0.000	18 (27.7)	10 (35.7)	0.467
TLR n (%)	11 (13.6)	17 (42.5)	0.001	15 (23.1)	8 (28.6)	0.606

Acute gain = (post-procedural – pre-procedural) MLD

Late loss = (post-procedural - follow-up) MLD

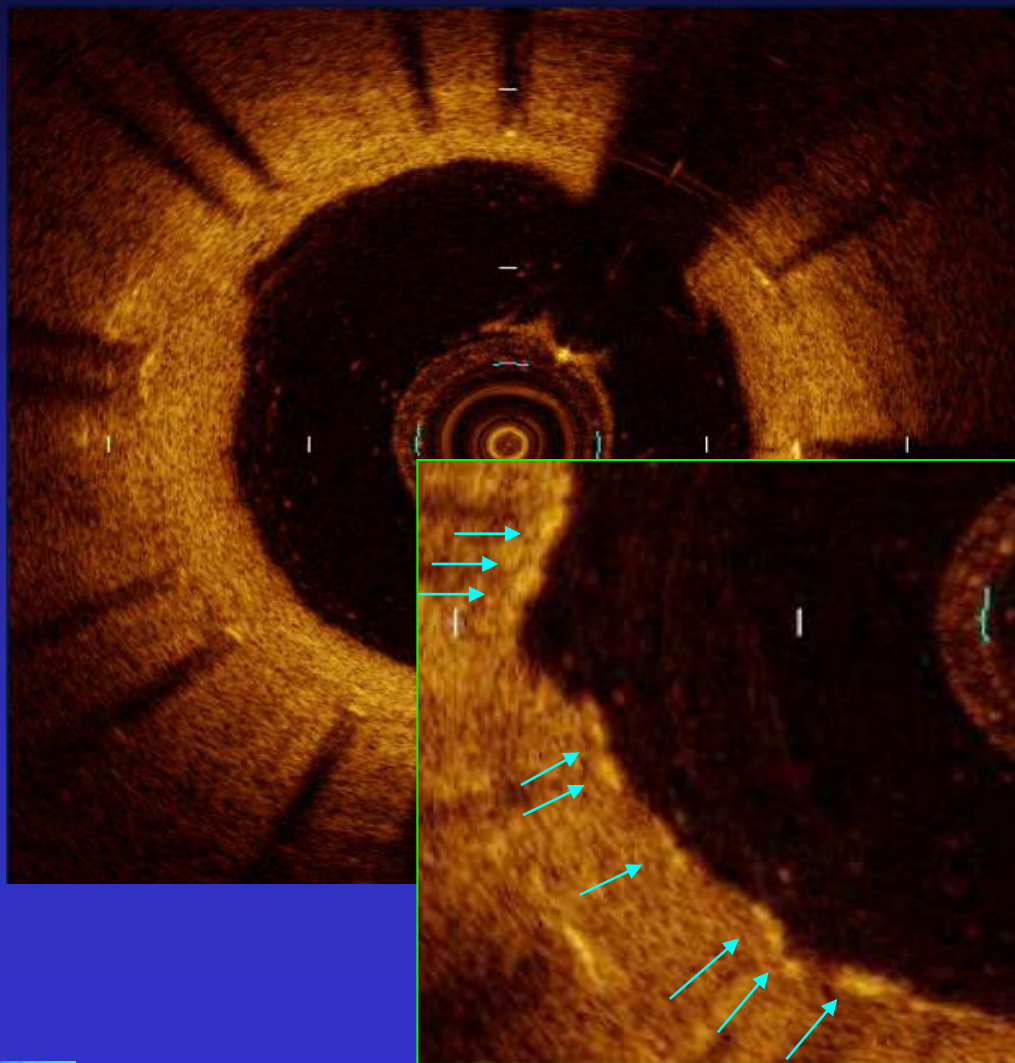
Net gain = (follow-up - pre-procedural) MLD



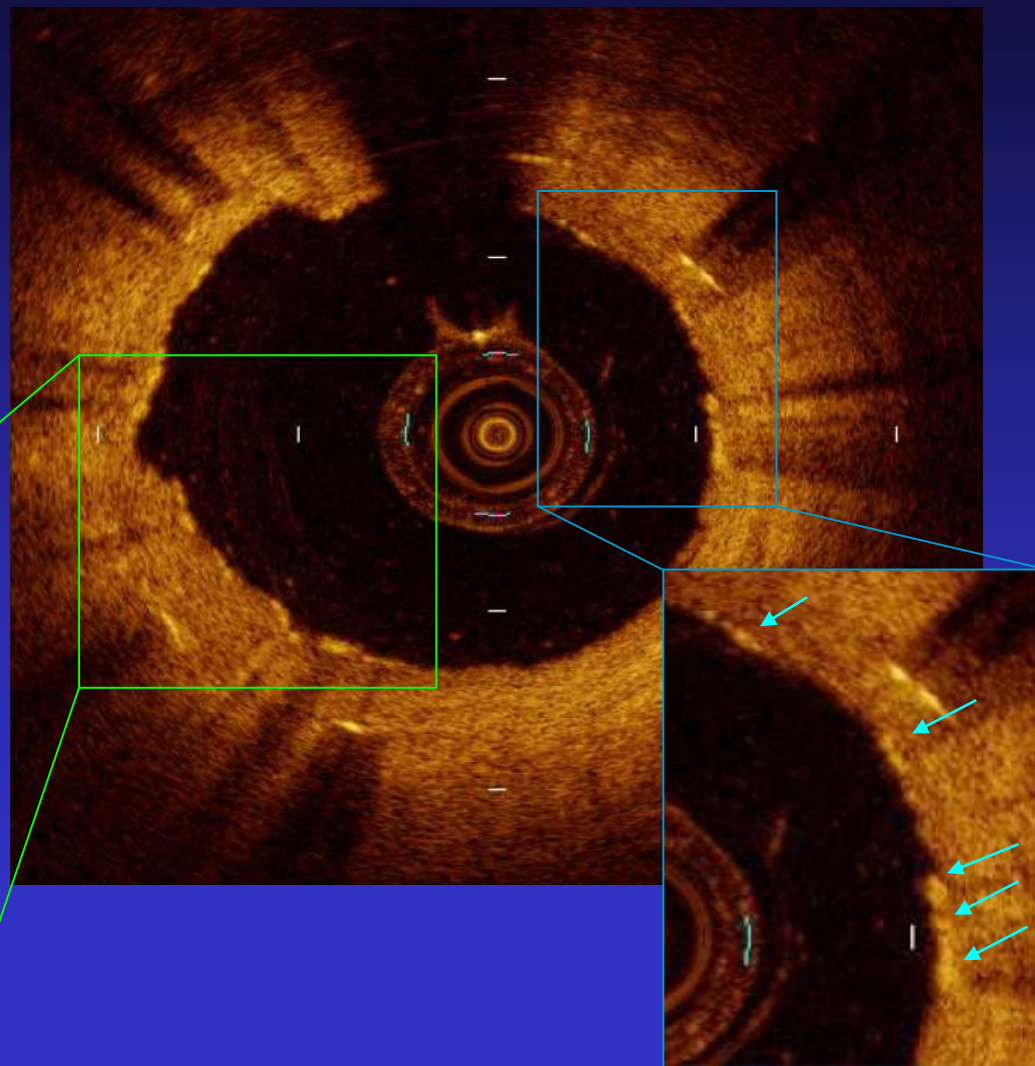
OCT findings after PCB for BMS restenosis

Courtesy by Dr. Habara S (Kurashiki Central Hosp.)

Post POBA



Post PCB



How to use OCT-guided PCI

● Pre PCI Assessment

- Image acquisition is very fast and easy.
- Precise measurements might be possible automatically.
- Lesion morphology can be assessed in detail.
Easy to plan PCI strategies, easy to decide stent landing zone, easy to identify unexpected lesions, etc.

● During and after PCI.

- Results of PCI such as tissue protrusion, incomplete apposition, mal-apposition, small dissection, etc. can be assessed precisely.
- Much more delicate treatment may be expected to bifurcation lesion stenting by 3D-OCT.
- Pathophysiology of LST & VLST could be demonstrated in detail and ideal treatment could be expected by OCT-guided PCI using PCB.
- OCT-guided PCI should be essential for BVS.



OCT: Pre- & Post Intervention (conclusions)

- **Pre- & post-PCI lesion morphology can be assessed easily & precisely by OCT because of higher resolution with high frame rate, auto-pullback & auto-measurement systems, etc.**
- **Improvement of clinical outcomes can be expected in PCI by the guidance of OCT, although there are not enough data to support the reduction of the adverse clinical events by OCT guided PCI.**
- **Randomized prospective studies should be planned to demonstrate the improvement of clinical outcome by OCT-guided PCI in the near future.**

