OCT Evaluation of Neointimal Coverage and Malapposition

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Neointimal Coverage After DES

Pathological Correlates of Late Drug-Eluting Stent Thrombosis

Strut Coverage as a Marker of Endothelialization

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Background—Late stent thrombosis (LST) after Cypher and Taxus drug-eluting stent placement has emerged as a major

The most powerful histological predictor of stent thrombosis was endothelial coverage.

The best morphometric predictor of LST was the ratio of uncovered to total stent struts.

to 22).

Conclusions—The most powerful histological predictor of stent thrombosis was endothelial coverage. The best morphometric predictor of LST was the ratio of uncovered to total stent struts. Heterogeneity of healing is a common finding in drug-eluting stents with evidence of LST and demonstrates the importance of incomplete healing of the stented segment in the pathophysiology of LST. (Circulation. 2007;115:2435-2441.)



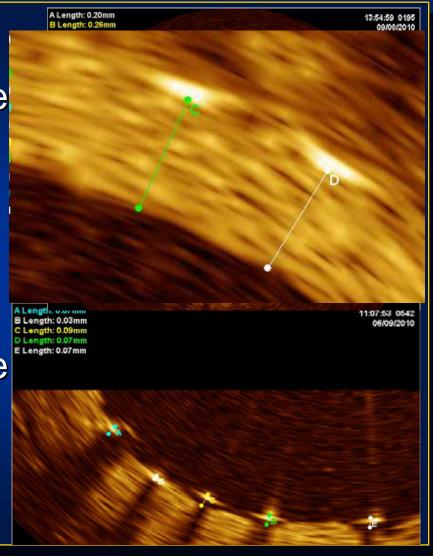
NIH Thickness & Malapposition

1. Neointimal thickness

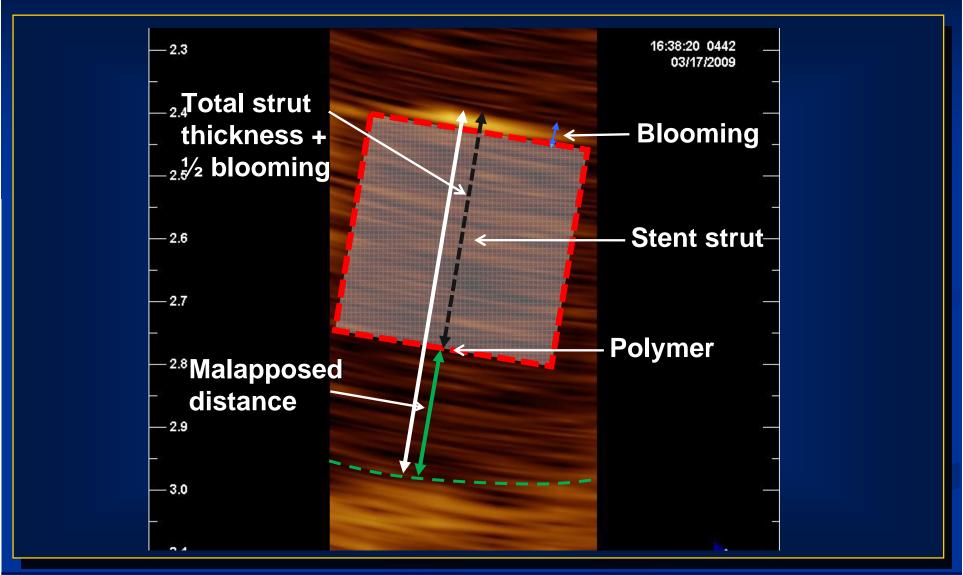
The distances between the endoluminal surface of neointimal and the strut reflection

2. Stent apposition

The distances between the endoluminal surface of the strut reflection and the vessel wall



Apposition Analysis





Apposition Analysis

1. Malapposition

Stent malappositions were defined as struts with detachment from the vessel wall \geq 160 μ m for SES, \geq 130 μ m for PES, \geq 110 μ m for ZES and \geq 100 μ m for EES

Tanigawa J, et al. EuroInterv 2007; 3:127-36.

2. Covered strut

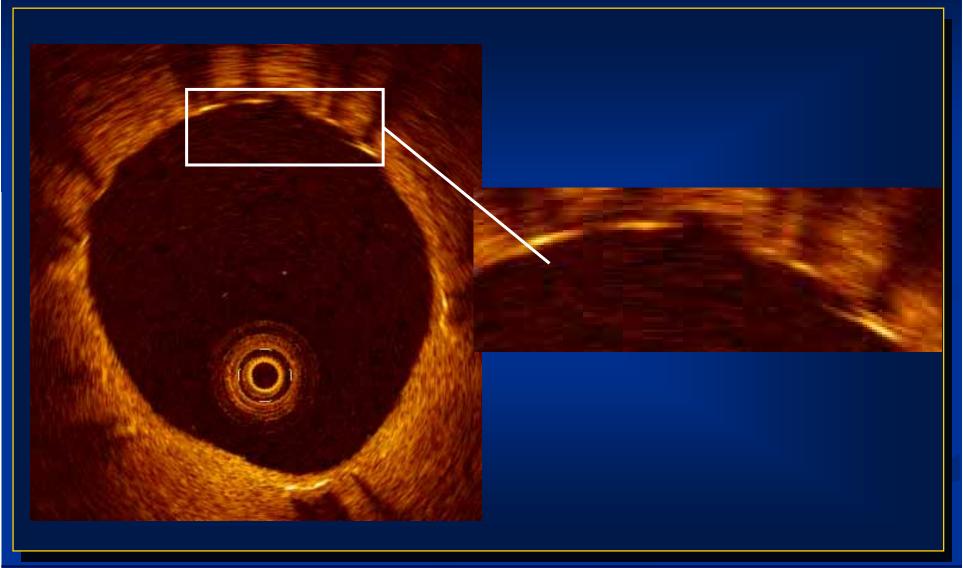
Covered strut defined as stent strut with detectable neointima by OCT

Takano M, et al. Am J Cardiol 2007;99:1033-8.

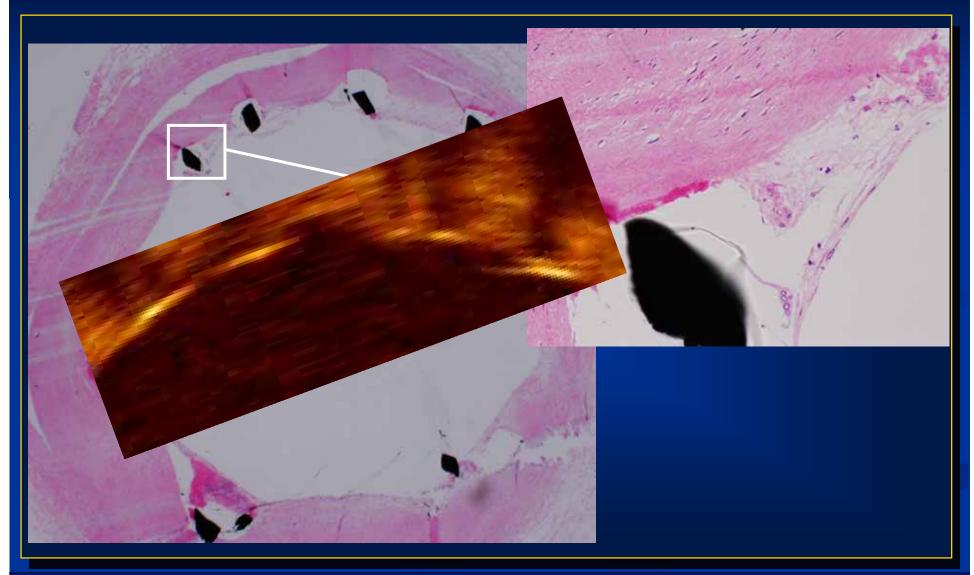
Case 1: Sirolimus stent @2M



Case 1: Sirolimus stent uncovered



Case 1: Sirolimus stent uncovered



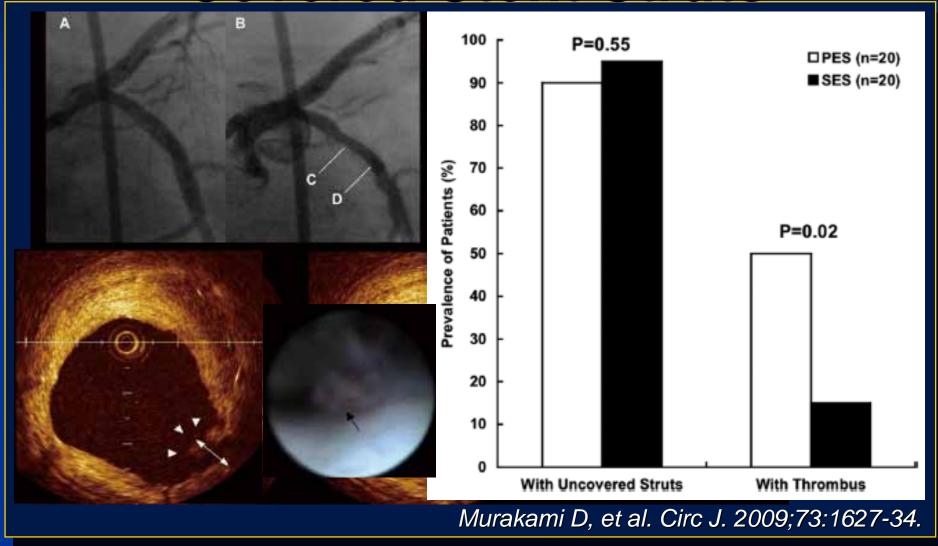
OCT Assessment of Strut Coverage

Uncovered stent strut



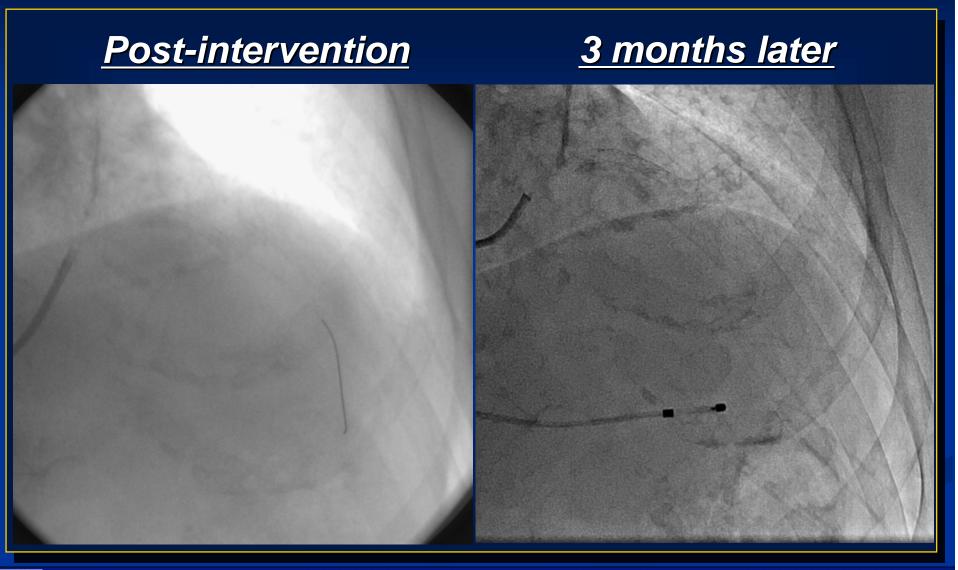
Lack of re-endothelialization

Thrombus Appearance on Covered Stent Struts

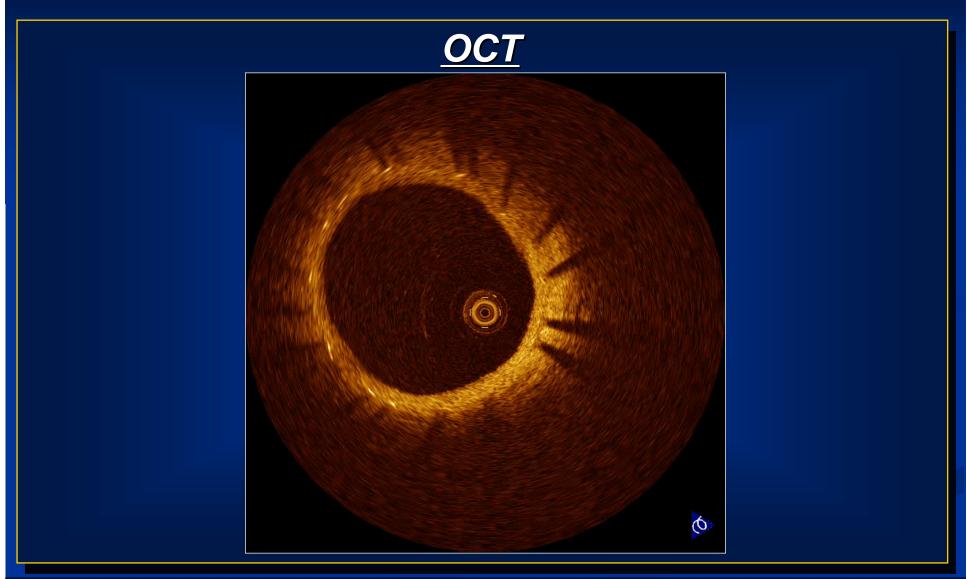




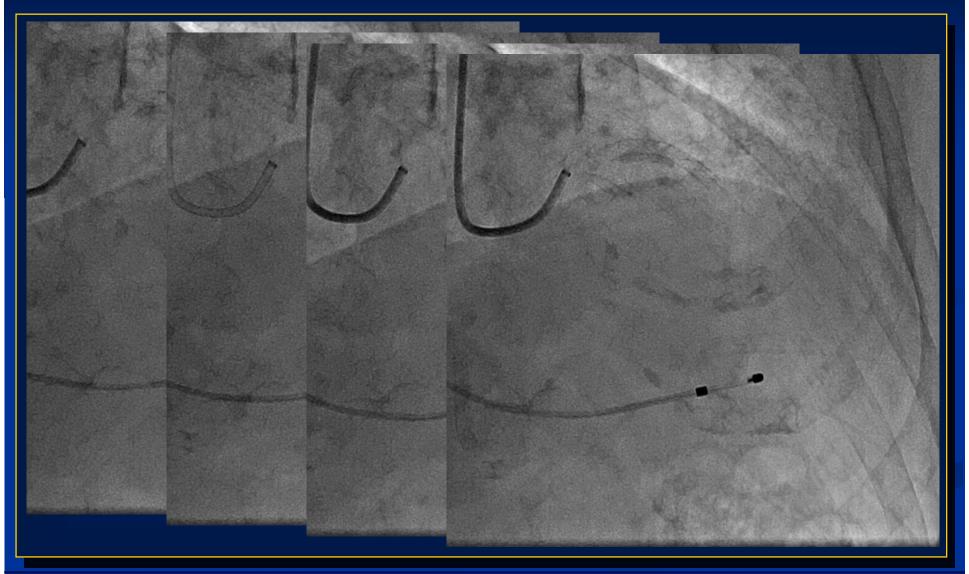
Case 2: Zotarolimus stent



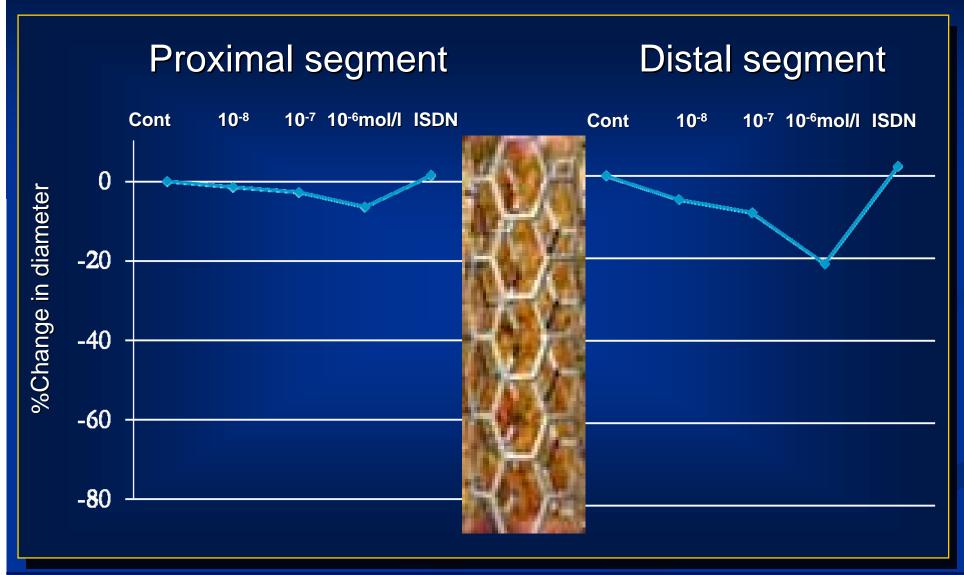
Case 2: Zotarolimus stent @3M



Case 2: Zotarolimus stent @3M

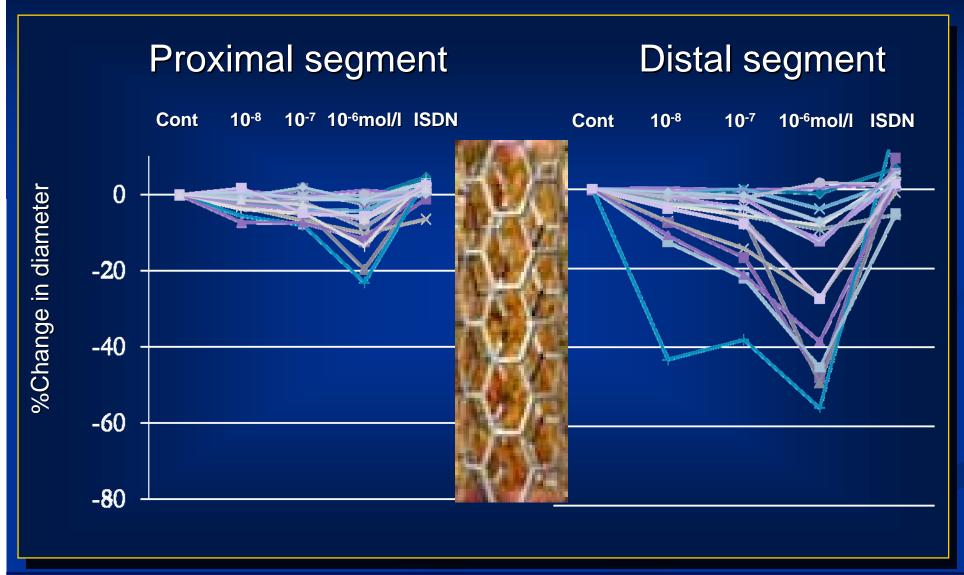


Vasomotor Response: ZES@3M





Vasomotor response: ZES@3M

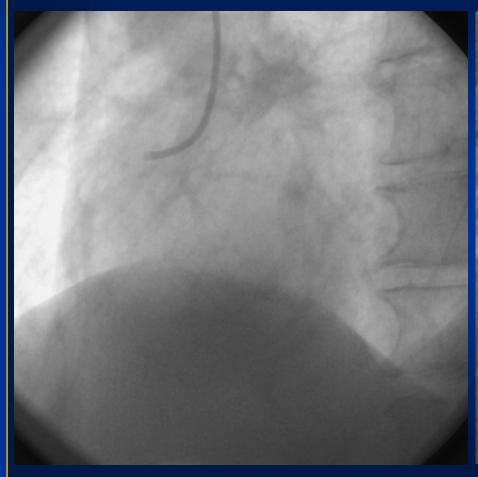




Case 3: Zotarolimus stent



3-month later

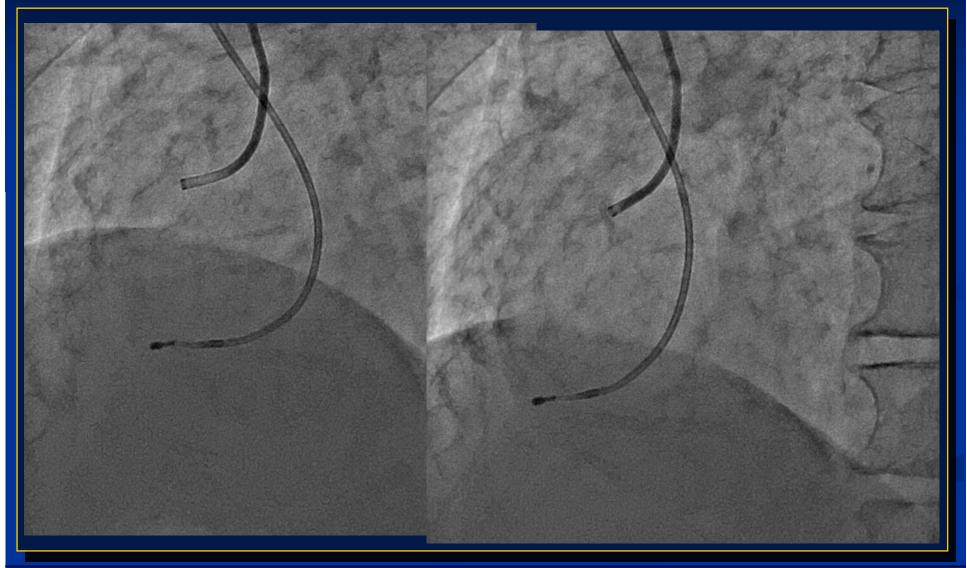




Case 3: Zotarolimus stent @3M

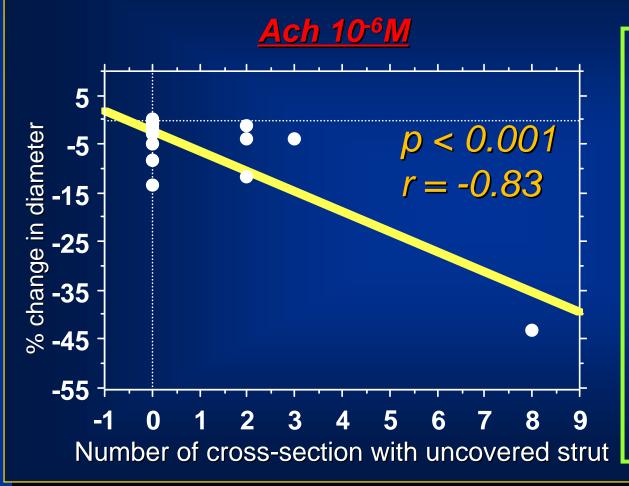


Case 3: Zotarolimus stent @3M



Vasomotor response: ZES @3M

OCT NIH & vasomotor response(distal to stent)



grades;

Grade 0: exposed strut

Grade 1: NIH thickness

 $< 100 \mu m$

Grade 2: NIH thickness

bet 100 and 200

 μ m

Grade 3: NIH thickness

 $> 200 \mu m.$

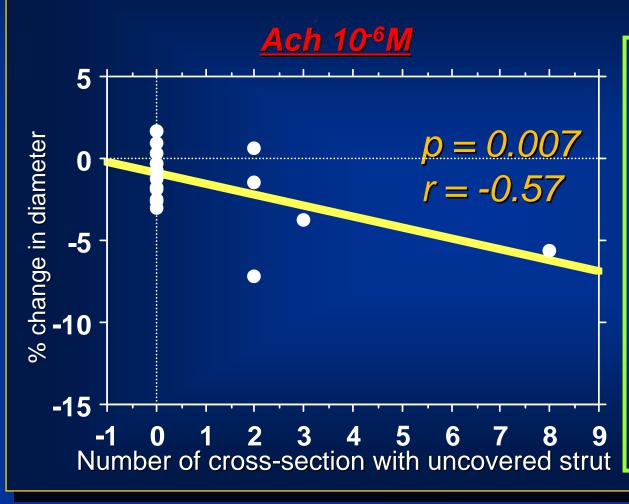
The grade was determined as minimal grade including ≥ 10 % of stent struts at each

cross section



Vasomotor response: ZES @3M

OCT NIH & vasomotor response(proximal to stent)



grades;

Grade 0: exposed strut

Grade 1: NIH thickness

 $< 100 \mu m$

Grade 2: NIH thickness

bet 100 and 200

 μ m

Grade 3: NIH thickness

 $> 200 \mu m.$

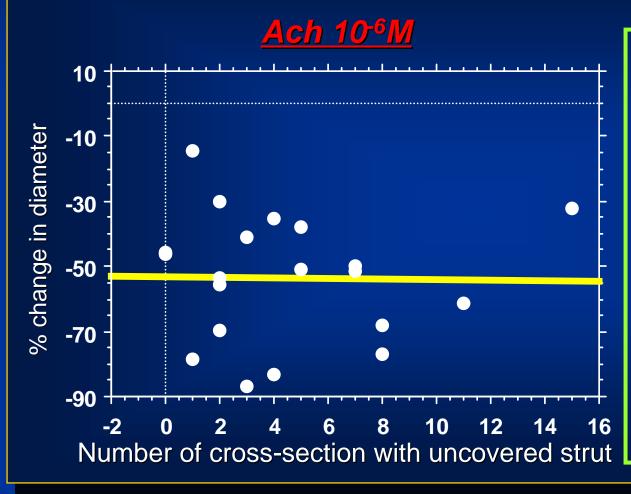
The grade was determined as minimal grade including ≥ 10 % of stent struts at each

cross section



Vasomotor response: SES @3M

OCT NIH & vasomotor response(distal to stent)



grades;

Grade 0: exposed strut

Grade 1: NIH thickness

 $< 100 \mu m$

Grade 2: NIH thickness

bet 100 and 200

 μ m

Grade 3: NIH thickness

 $> 200 \mu m.$

The grade was determined as minimal grade including ≥ 10 % of stent struts at each

cross section



OCT Qualitative Assessment of Neointimal Tissue

Restenotic tissue structure



Homogeneous: restenotic tissue has uniform optical properties and does not show focal variations in backscattering pattern.



Heterogeneous: restenotic tissue has focally changing optical properties and shows various backscattering patterns



Layered: restenotic tissue consists of concentric layers with different optical properties: an adluminal high scattering layer and an abluminal low scattering layer

Restenotic tissue backscatter



High: the majority of the tissue shows high backscatter and appears



Low: the majority of the tissue shows low backscatter and appears dark or black

Microvessels visible



Yes: microvessels appear as well delineated low backscattering structures less than 200 micron in diameter that show a trajectory within the



No

Lumen shape



Regular: lumen border is sharpy delineated, smooth and circular



Irregular: lumen border irregular with tissue protrusions from the vessel wall into the lumen

Presence of intraluminal material



Yes: there is visible material inside the vessel lumen.



No

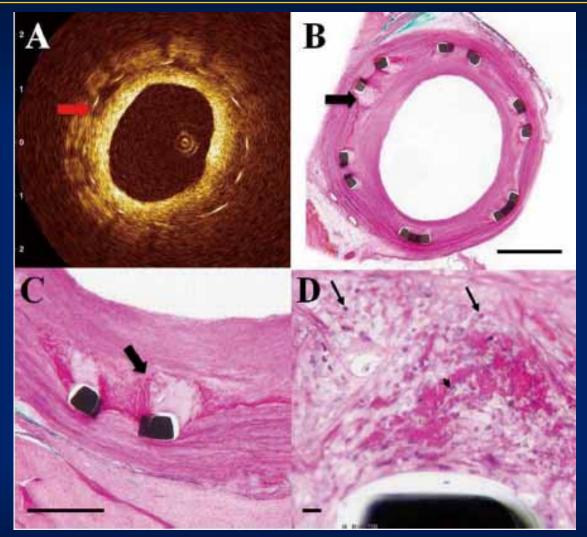
Gonzalo N, et al. Am Heart J. 2009;158:284-93.



Case 4: Paclitaxcel stent @6M

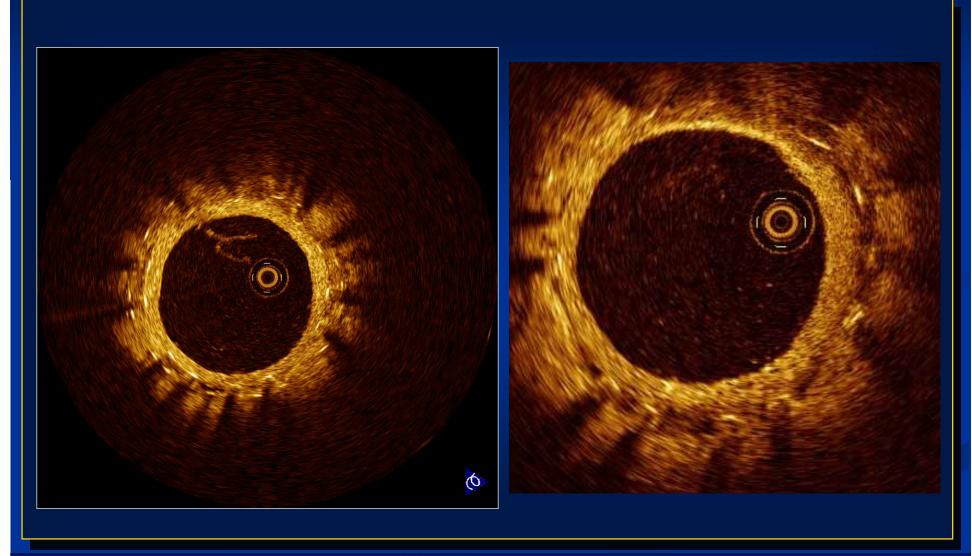


Peri-strut low-intensity area: DES



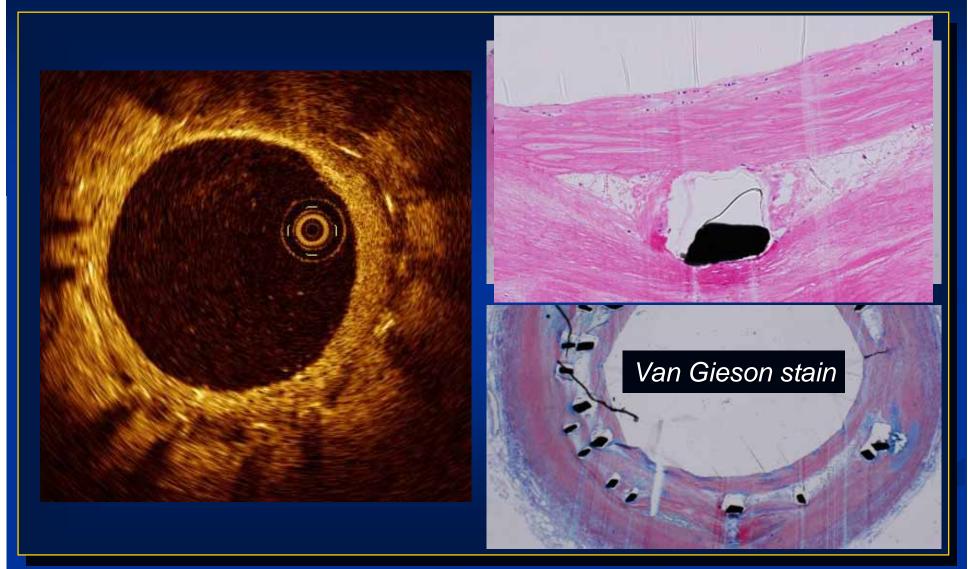
Teramoto T, et al. Circ J. 2010;74:1257-9.

Peri-strut low-intensity area: BMS @10Y

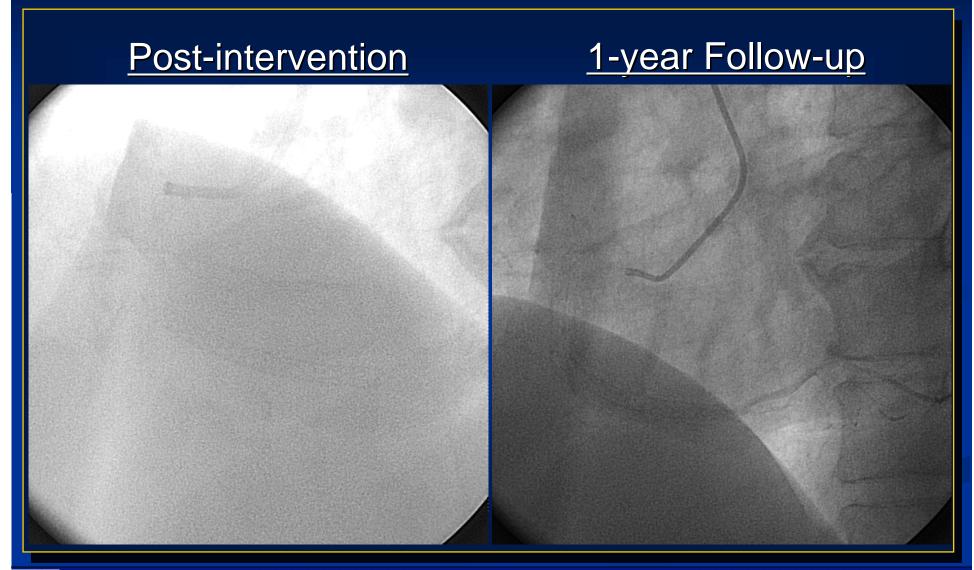




Peri-strut low-intensity area: BMS @10Y



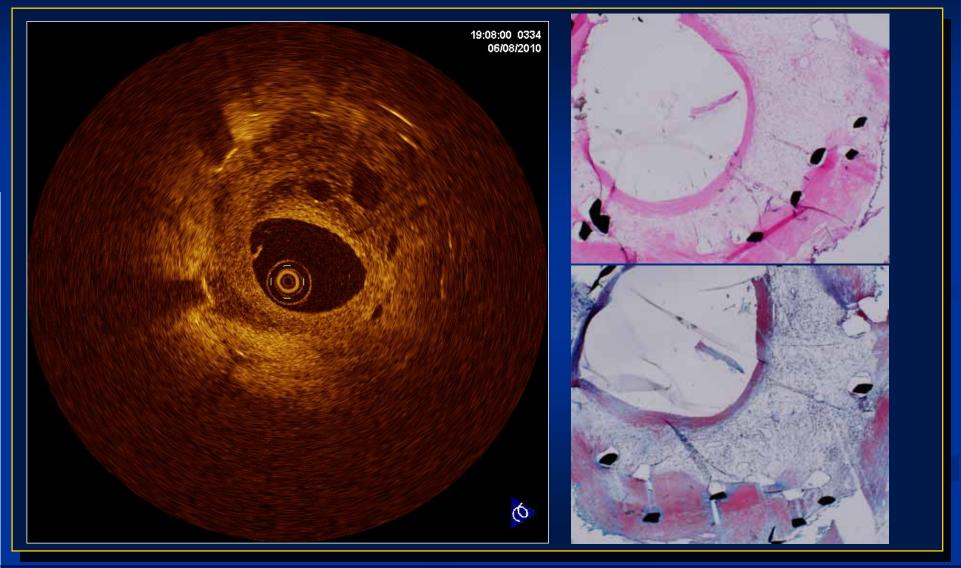
Case 5: Sirolimus stent @12M



Case 5: Sirolimus stent @12M



Microvessel Appearance: BMS @9Y



Summary

- Although OCT image did not detect strut coverage, one layer endothelium (about 600 nm) was observed with histopathological finding.
- ✓ Some case reports suggested that the degree of neointimal growth in DES was not associated with coronary endothelial function.
- ✓ Linear relationship was observed between neointimal coverage of stent struts assessed by OCT and vasomotor response to acetylcholine in ZESs, but not in SESs.

Summary

- ✓ Although abnormal morphological findings (i.e. uncovered and malapposed struts, peri-strut low-intensity area, microvessel appearance) were common in DESs, it unclear whether these stents lead to acute thrombotic event (the incidence of stent thrombosis is low).
- There were no data on clinical implications of abnormal intimal tissue morphology detected by OCT.

Thank you for your attention

