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# Clinical Outcomes Using Coronary CT Angiography and $\text{FFR}_{\text{CT}}$ Guided Management of Stable Chest Pain Patients

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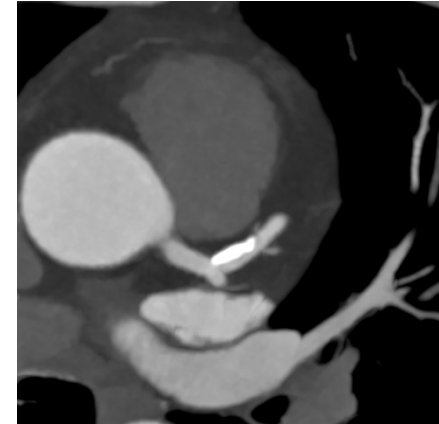


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# Background

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- **Coronary CT Angiography:**



- Can accurately exclude the presence of CAD<sup>1</sup>
- Prognostic implications<sup>2</sup>
- Cannot determine the physiologic significance of lesions<sup>3</sup>

**Non-invasive strategies are needed to identify those patients with CAD who may benefit from cardiac catheterization and those who do not require further testing**

<sup>1</sup>Abdulla J et al, EHJ 2007; <sup>2</sup>Nielsen LH et al, EHJ 2017; <sup>2</sup>Xie JX et al, iJACC 2018; <sup>3</sup>Meijboom WB et al, JACC 2008; <sup>3</sup>Norgaard BL et al, JACC 2014

# Background

- CTA derived fractional flow reserve ( $\text{FFR}_{\text{CT}}$ ):



- High and improved diagnostic performance compared to CTA<sup>1</sup>
- Have shown promise in guiding downstream management of patients with CAD<sup>2</sup>
- One-year outcomes of  $\text{FFR}_{\text{CT}}$  guided care in a clinical trial setting was favorable<sup>2</sup>

**Longer term clinical outcome data in patients undergoing CTA testing with  $\text{FFR}_{\text{CT}}$  guidance in day-to-day practice is sparse**

<sup>1</sup>Koo BK et al, JACC 2011; <sup>1</sup>Min JK et al, JAMA 2012; <sup>1</sup>Norgaard BL et al, JACC 2014; <sup>2</sup>Douglas PS et al, JACC 2016

# Overall purpose of the study

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- To assess the safety and clinical outcomes of utilizing a diagnostic strategy of first-line coronary CTA with selective  $\text{FFR}_{\text{CT}}$  testing in real world symptomatic patients with suspected stable CAD

# Study design

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- Single-center, observational all-comer study of symptomatic patients undergoing non-emergent coronary CTA for suspected CAD with selective FFR<sub>CT</sub> testing between May 2014 and December 2016
- Data sources
  - The Western Denmark Cardiac Computed Tomography Registry<sup>1</sup>
    - Patient demographics, CTA results
  - The Danish National Patient Registry<sup>2</sup>
    - Discharge diagnoses, test and procedures for all in and outpatient encounters
  - The Danish Civil Registration system<sup>3</sup>
    - Data on mortality

<sup>1</sup>Nielsen LH et al, Clin Epidemiol 2014; <sup>2</sup>Schmidt M et al, Clin Epidemiol 2015; <sup>3</sup>Schmidt M et al, Eur J Epidemiol 2014

# Patients

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- All Aarhus University Hospital patients with new onset chest pain and suspected CAD who had non-emergent coronary CTA performed from May 2014 to December 2016
  - Coronary CTA is the first-line test in such patients
  - CTA acquisition was performed according to societal guidelines<sup>1</sup>
    - Exclusion from CTA: Contrast allergy, pregnancy, scenarios where a diagnostic image quality cannot be expected (combination of e.g. obesity, arrhythmia, and severe calcification)

<sup>1</sup>Abbara S et al, JCCT 2016

# Post-test management, *Coronary CTA*

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	Test outcome	Post-test management recommendations
<b>Coronary CTA</b>		
Diagnostic conclusive	High-risk anatomy	ICA
	Intermediate-risk anatomy	<u>FFR<sub>CT</sub></u>
	Low-risk anatomy	No further testing
Diagnostic inconclusive	-	MPI, or ICA

Optimal medical treatment was recommended in all patients with CAD  
ICA =invasive coronary angiography, MPI =myocardial perfusion imaging

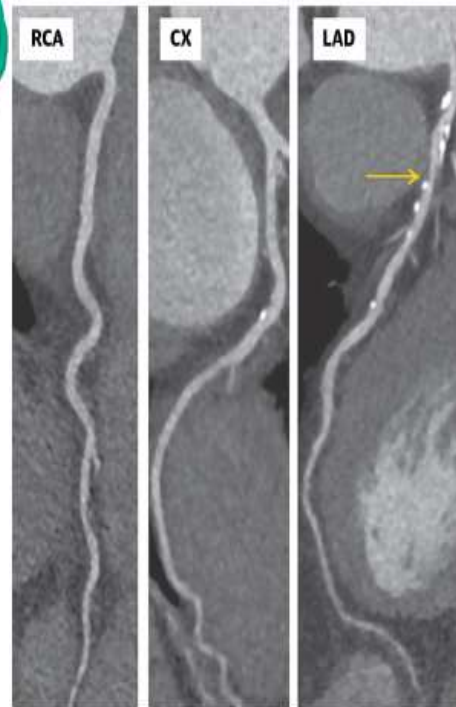
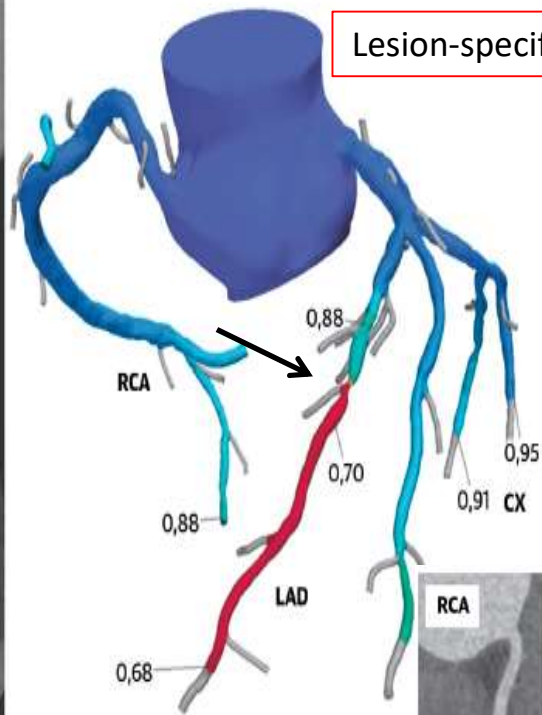
## Post-test management, $FFR_{CT}$

	Test outcome	Post-test management recommendations
<b><math>FFR_{CT}</math></b>		
Diagnostic conclusive	<b>Negative</b> , all values $>0.80$	OMT, no additional testing
	<b>Positive</b> , one or more values $\leq 0.80$	
	-Lesion-specific ischemia	OMT or ICA
	-Distal vessel positivity	OMT
Diagnostic inconclusive	-	MPI, or ICA

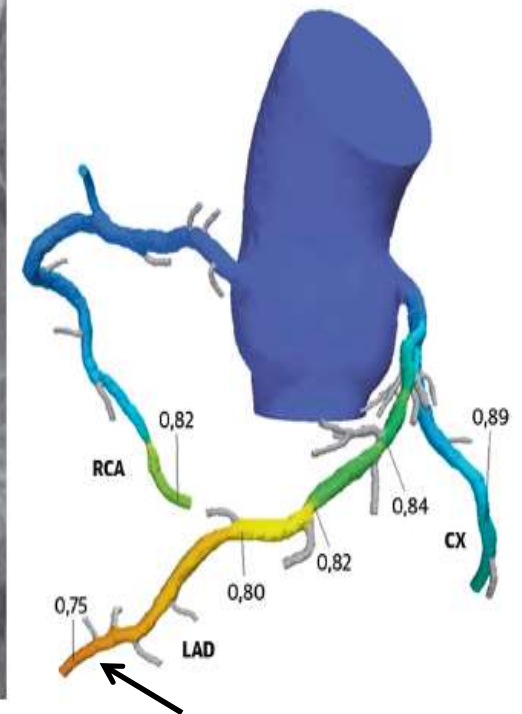
ICA =invasive coronary angiography, MPI =myocardial perfusion imaging, OMT =optimal medical treatment



### Lesion-specific ischemia



### Distal vessel FFR<sub>CT</sub> positivity



# Endpoint, Follow-up, and Study aims

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- **Endpoint:** Composite of all-cause death, non-fatal myocardial infarction, hospitalization for unstable angina, and unplanned revascularization
- **Follow-up:** Median 24 (interquartile range, 16-32; range, 8-41) months. No patients were lost to follow-up

● **Primary aim:** The cumulative incidence of the combined endpoint in patients with  $\text{FFR}_{\text{CT}} > 0.80$ , and no additional testing compared to patients with no or minimal (stenosis severity  $< 30\%$ ) CAD

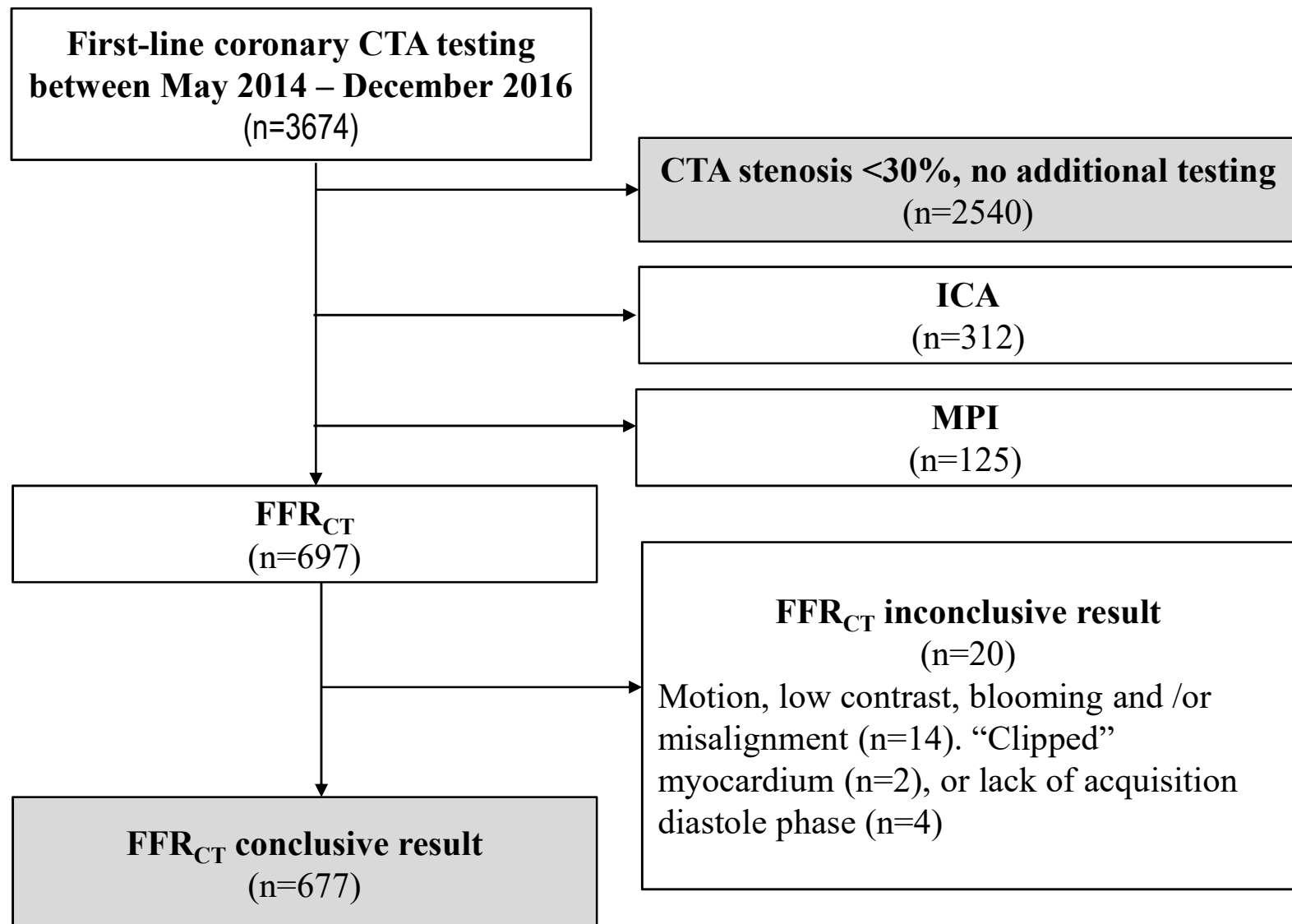
# Endpoint, Follow-up, and Study aims

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- **Endpoint:** Composite of all-cause death, non-fatal myocardial infarction, hospitalization for unstable angina, and unplanned revascularization
- **Follow-up:** Median 24 (interquartile range, 16-32; range, 8-41) months. No patients were lost to follow-up

- **Primary aim:** The cumulative incidence of the combined endpoint in patients with  $\text{FFR}_{\text{CT}} > 0.80$ , and no additional testing compared to patients with no or minimal (stenosis severity  $< 30\%$ ) CAD
- **Secondary aim:** The cumulative incidence of the combined endpoint in patients with  $\text{FFR}_{\text{CT}} \leq 0.80$  (OMT or ICA), compared to patients with CTA stenosis  $< 30\%$

# Results: Patients Flow-chart



# Results: Baseline characteristics

	CTA stenosis <30%	CTA stenosis ≥30%		P-value (FFR <sub>CT</sub> >0.80 versus FFR <sub>CT</sub> ≤0.80 group)
	OMT (n =2540)	FFR <sub>CT</sub> >0.80, OMT (n=410)	FFR <sub>CT</sub> ≤0.80, OMT or ICA (n=267)	
Age, yrs, mean	56	60	62	0.006
Male, %	43	55	65	0.02
Diabetes mellitus,%	6	9	14	0.16
Hypertension,%	30	40	50	0.005
Updated D-F score, mean %	31	43	47	0.01

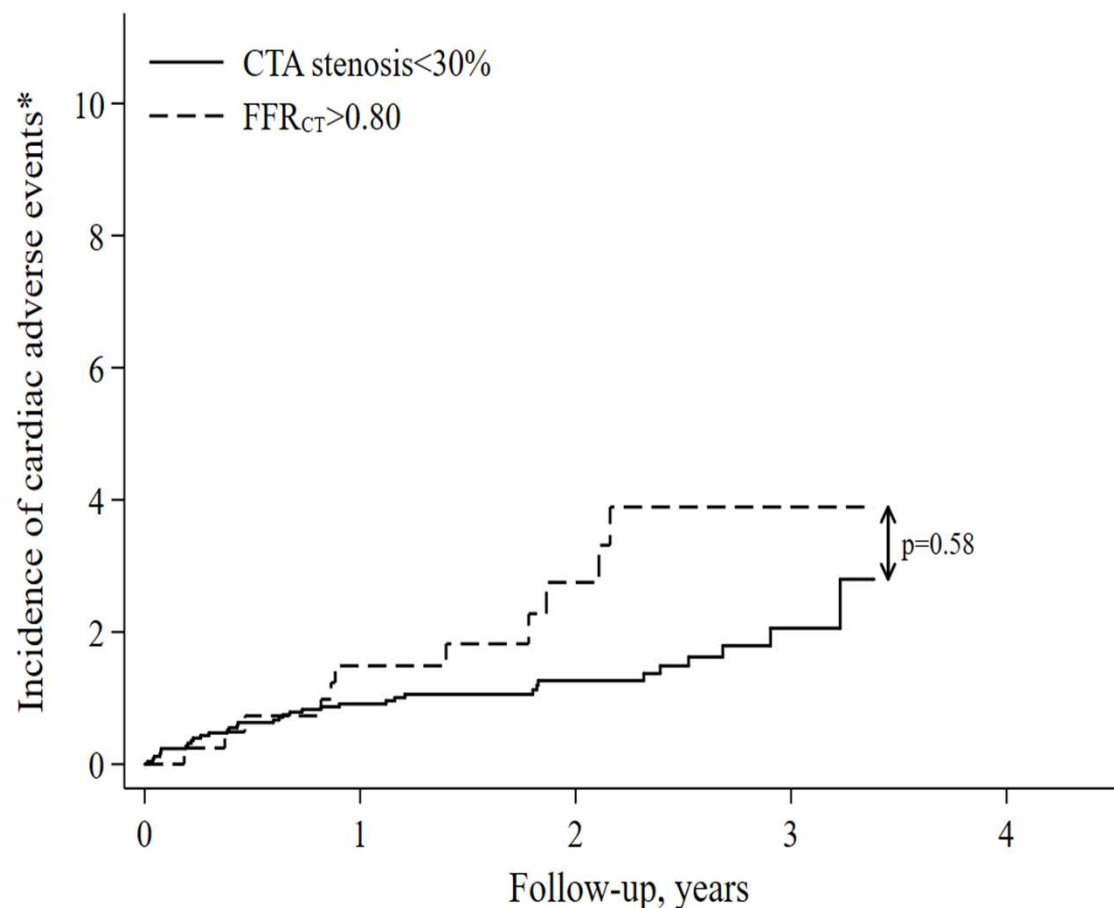
DF = Diamond-Forrester

# Results: Anatomical characteristics

	FFR <sub>CT</sub> >0.80, OMT (n=410)	FFR <sub>CT</sub> ≤0.80, OMT or ICA (n=267)	P-value
Maximum CTA stenosis			
30-49%	25%	9%	<0.001
50-69%	65%	59%	0.10
≥70%	10%	32%	<0.001
Vessels with stenosis ≥50%			<0.001
1	63%	56%	
2	10%	27%	
3	1%	7%	
Mean Agatston score	164	456	<0.001

# Results: Clinical outcomes

\*All-cause death,  
non-fatal myocardial MI,  
hospitalization for unstable angina,  
unplanned revascularization

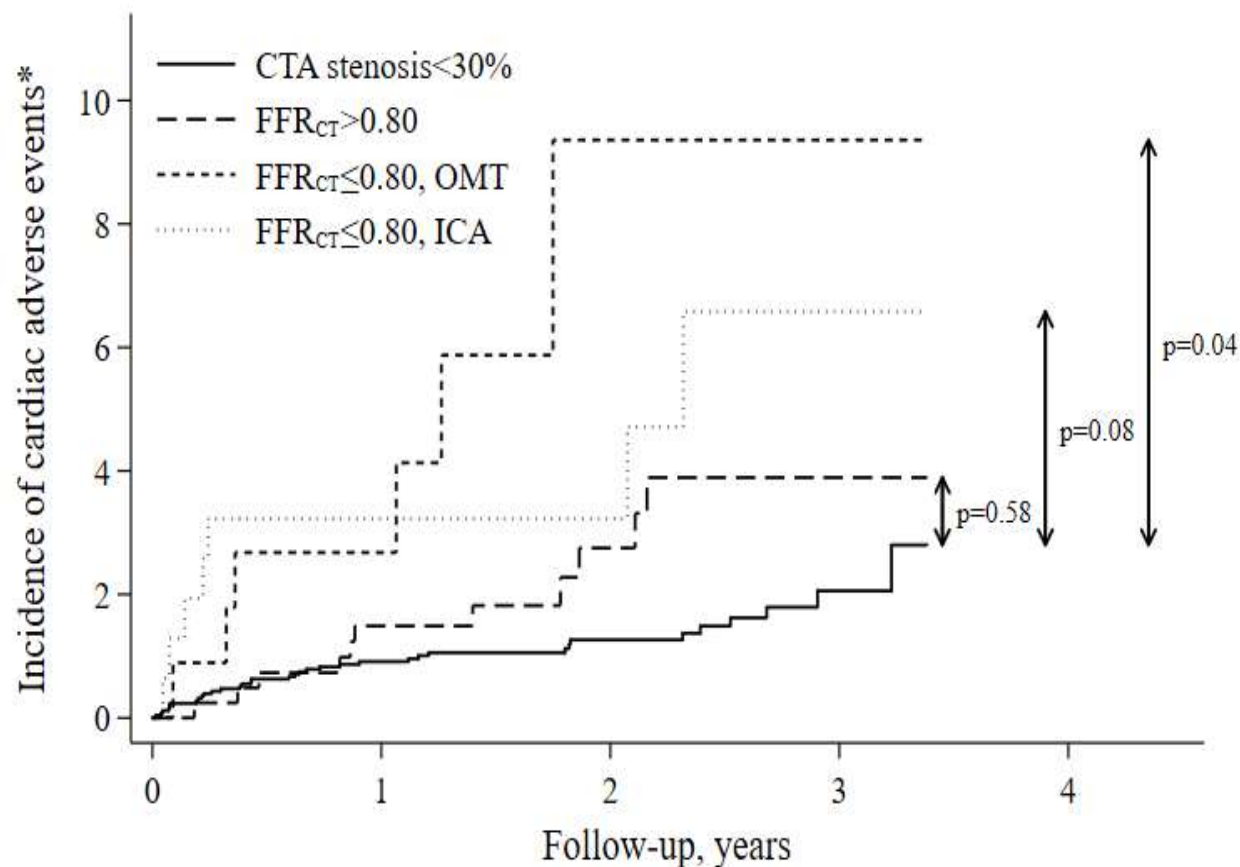


Number at risk

CTA stenosis<30%	2540	2187	1226	285
FFR <sub>CT</sub> >0.80	410	366	187	49

# Results: Clinical outcomes

\*All-cause death,  
non-fatal myocardial MI,  
hospitalization for unstable angina,  
unplanned revascularization



Number at risk

CTA stenosis<30%	2540	2187	1226	285
FFR <sub>CT</sub> >0.80	410	366	187	49
FFR <sub>CT</sub> ≤0.80, OMT	112	75	20	3
FFR <sub>CT</sub> ≤0.80, ICA	155	132	71	19

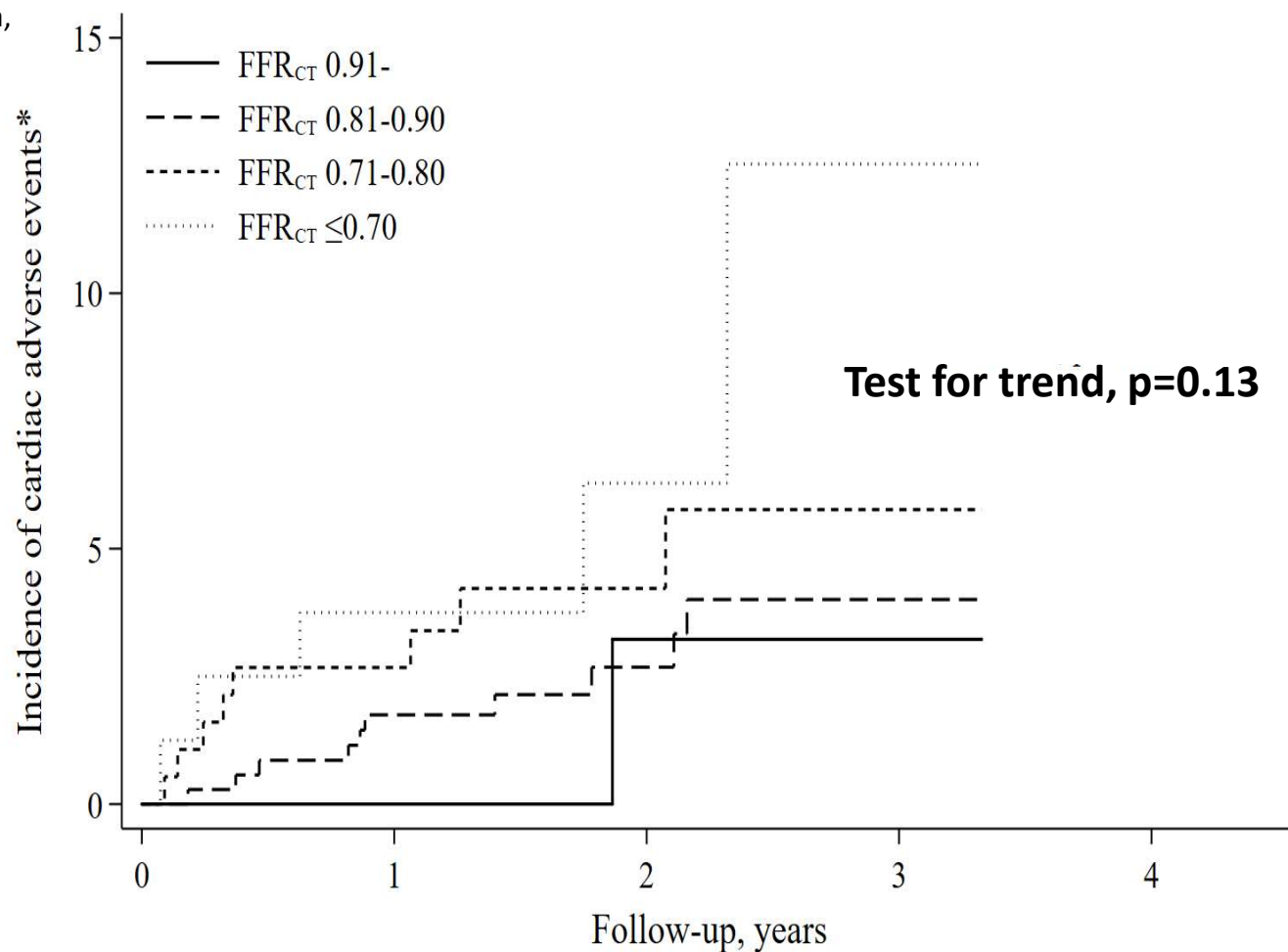


# Results: Clinical outcomes

	CTA stenosis <30%	CTA stenosis ≥30%			P-value
	OMT (n=2540)	FFR <sub>CT</sub> >0.80, OMT (n=410)	FFR <sub>CT</sub> ≤0.80, OMT (n=112)	FFR <sub>CT</sub> ≤0.80, ICA (n=155)	
Composite end-point	2.8 (1.4-4.9)	3.9 (2.0-6.9)	9.4 (3.0-20.0)	6.6. (2.5-13.4)	0.07
All-cause death	2.3 (1.0-4.4)	1.4	1.5	2.8	0.97
Non-fatal MI	0.3 (0.1-0.6)	0.3	8.0 (2.2-18.6)	1.3	<0.001
Hospitalization for UA	0.1	1.7	0.9	2.5	0.01
Unplanned revascularization	0.4 (0.2-0.8)	1.0	8.8 (2.2-18.6)	0	<0.01

# Results: Clinical outcomes

\*All-cause death,  
non-fatal myocardial MI,  
hospitalization for unstable angina,  
unplanned revascularization



# Summary

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- In a real-world setting of symptomatic patients without known CAD, the presence of intermediate range CTA stenosis and  $\text{FFR}_{\text{CT}} > 0.80$  was associated with favorable clinical outcomes similar to patients with no or minimal evidence of CAD
- Risk of an unfavorable outcome was increased (driven by a higher incidence of non-fatal MI) in patients with  $\text{FFR}_{\text{CT}} \leq 0.80$ , who were not referred to ICA

# Conclusion

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- In a real-world clinical practice, a diagnostic strategy of first-line coronary CTA in symptomatic patients suspected of CAD, and  $\text{FFR}_{\text{CT}}$  testing in those with intermediate range lesions is effective in differentiating patients who do not require further diagnostic testing or intervention ( $\text{FFR}_{\text{CT}} > 0.80$ ) from higher risk patients ( $\text{FFR}_{\text{CT}} \leq 0.80$ ) in whom further testing with ICA and possibly intervention may be needed

# Thank you!