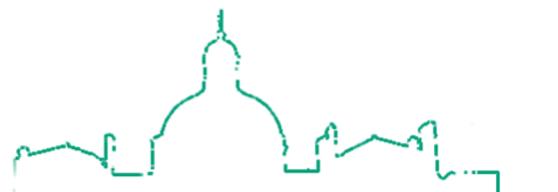


# Durée du traitement antiagrégant plaquettaire : toujours plus court



## Gilles Montalescot

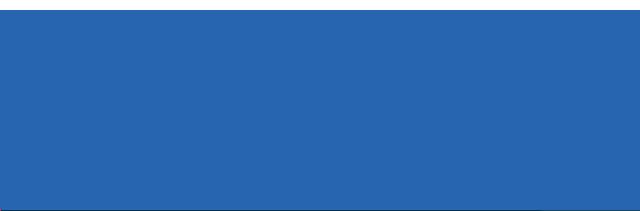
Dr. Montalescot reports research Grants to the Institution or Consulting/Lecture Fees from Abbott, AIM group, Amgen, Actelion, ACC Foundation, AstraZeneca, Axis-Santé, Bayer, Boston-Scientific, BMS, Beth Israel Deaconess Medical, Brigham Women's Hospital, Fréquence Médicale, ICOM, Idorsia, Elsevier, ICAN, Lead-Up, Menarini, MSD, Novo-Nordisk, Pfizer, Quantum Genomics, Sanofi-Aventis, SCOR global life, Servier, WebMD.



INSTITUT DE CARDIOLOGIE  
Pitié-Salpêtrière  
Paris



Paris, France



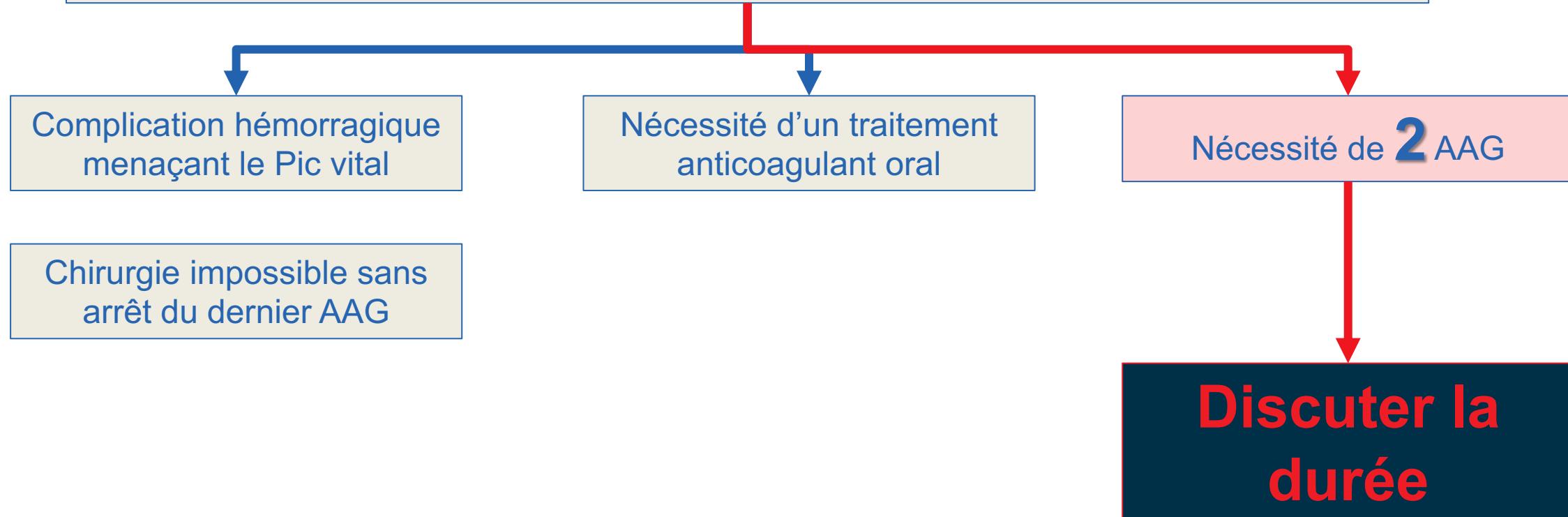
+ court?



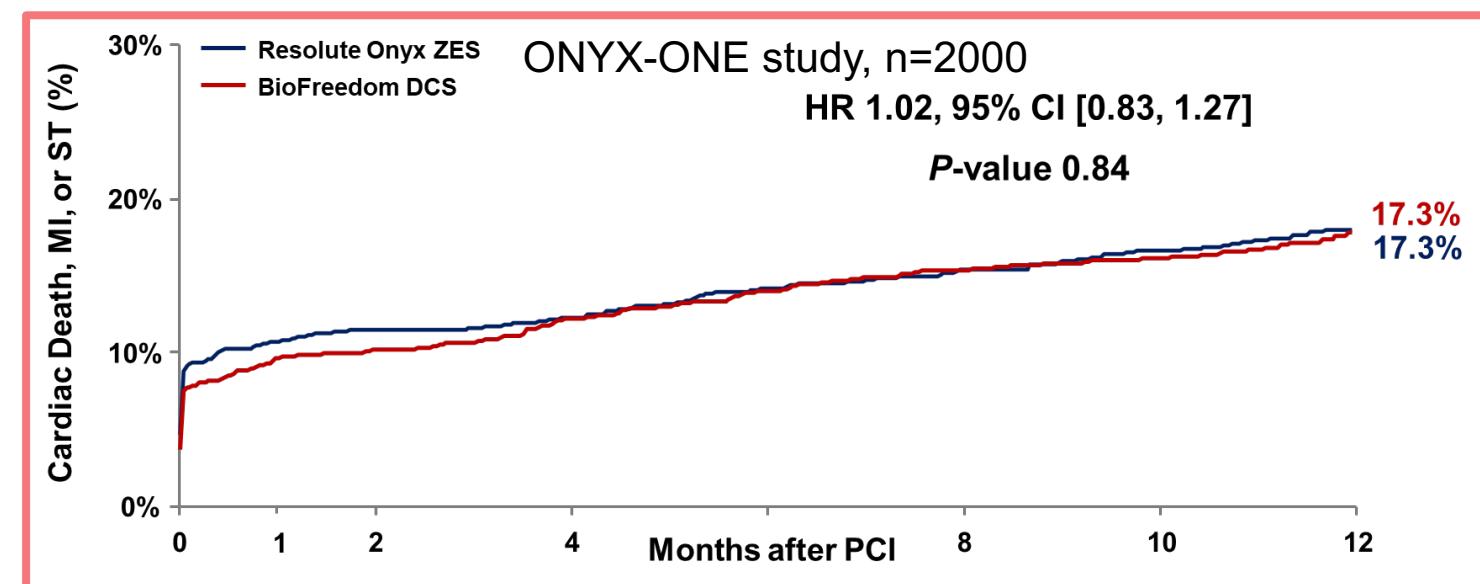
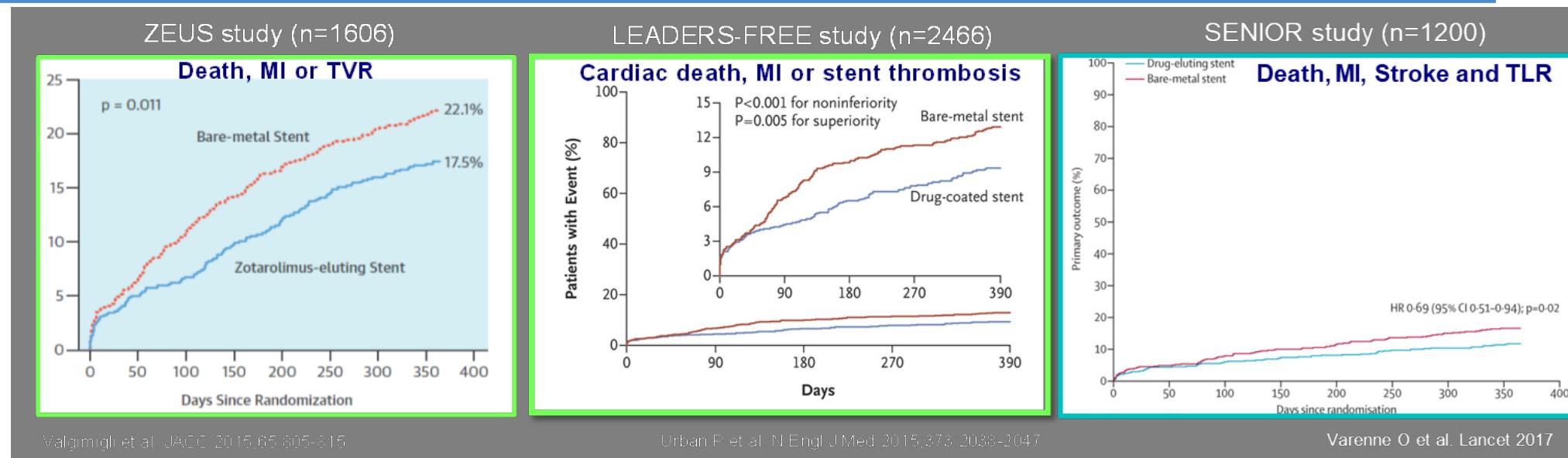
# De quoi parle-t-on?



**Le coronarien est redevable d'**1** AAG à vie!  
(sauf exception)**



# STENT + HBR = Arrêt précoce du clopi



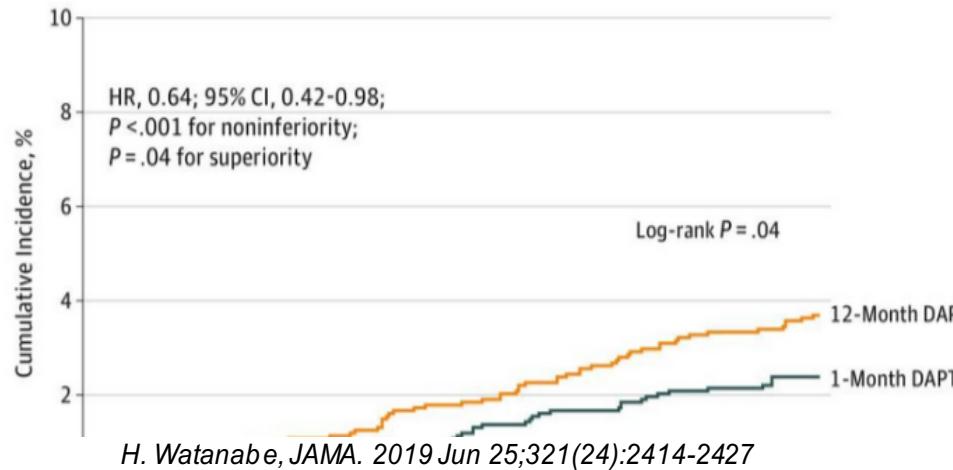
HBR: haut risque hémorragique

# STOP-DAPT-2

[op\\_pmc\\_inline.html?title=Click on image to zoom&p=PMC3&id=6593641\\_jama-321-2414-g002.jpg](#)

Drag image to reposition. Double click to magnify further.

**A** Primary end point (composite of cardiovascular death, MI, definite stent thrombosis, ischemic and hemorrhagic stroke, or TIMI major or minor bleeding)

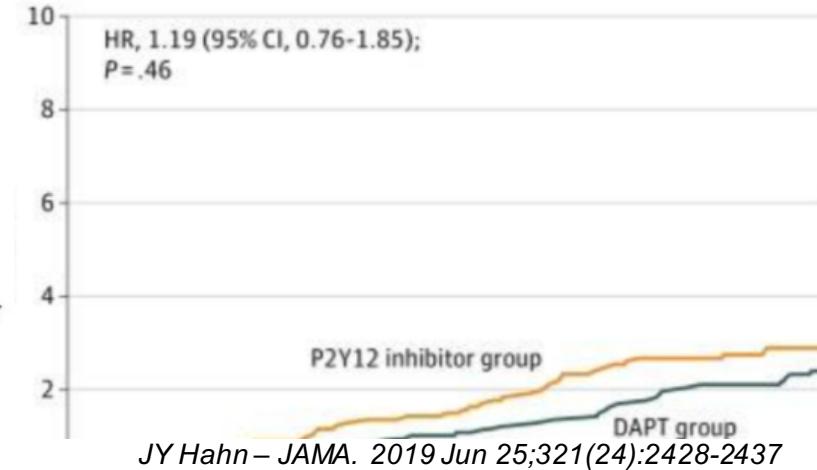


# SMART-CHOICE

[y.insermbiblio.inist.fr/core/lw/2.0/html/tileshop\\_pmc/tileshop\\_pmc\\_inline.html?title=Click on image to zoom&p=PMC3&id=6593641\\_jama-321-2414-g002.jpg](#)

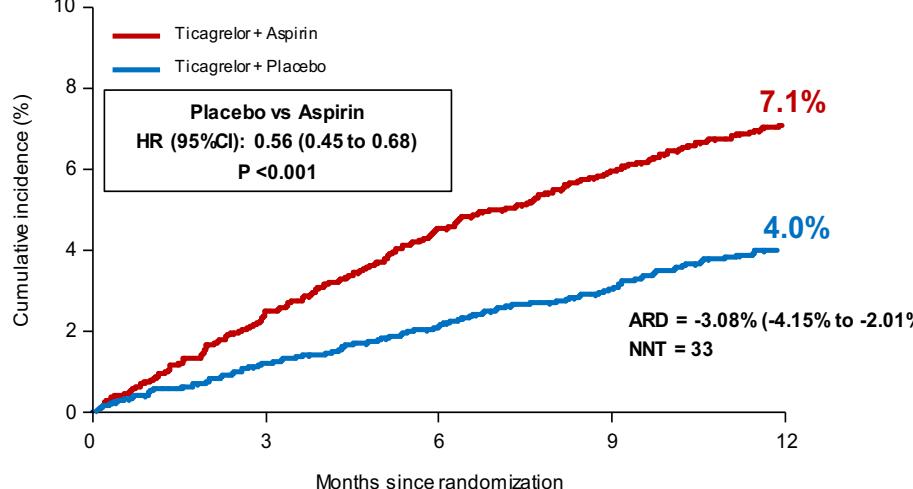
Drag image to reposition.

Composite events (primary outcome)



# TWILIGHT

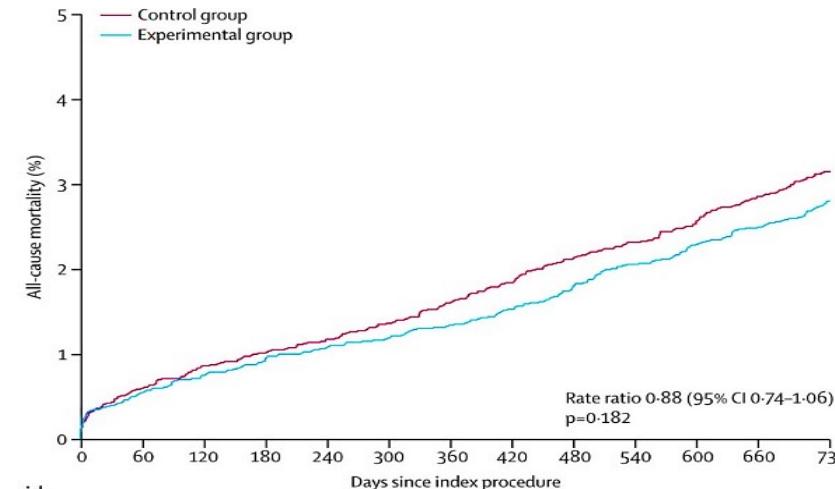
BARC 2, 3 or 5 Bleeding (primary outcome)



Mehran R et al. N Engl J Med. 2019 Nov 21;381(21):2032-2042

# GLOBAL-LEADERS

Control group  
Experimental group



Vranckx P et al. Lancet 2018

	Elderly age $\geq 75$ years		Thrombocytopenia ( $<100,000/\text{mm}^3$ )		
	OAC planned after PCI			Cancer diagnosed or treated w/i 3 years	
	Renal failure ( $\text{CrCl} <40 \text{ ml/min}$ )			Stroke within 1 year or any prior ICH	
	Planned surgery $<1$ year			Severe chronic liver disease	
	Anemia ( $\text{Hgb} <11 \text{ g/dl}$ )			Long-term NSAID or steroid use	
	Hospitalization for bleeding within 1		<input type="checkbox"/>	Expected DAPT non-compliance	

# Moyen mémotechnique



## A AGE

- Frail elderly >75 years\*
- Advanced age >85 years\*
- Life expectancy <1 year

\* Must be accompanied with an additional risk factor



## B BLEEDING

- Spontaneous intracranial haemorrhage
- Recurrent gastrointestinal bleeding
- Haemoglobin <9 g/dL



## O ORGAN DYSFUNCTION

- Liver cirrhosis
- End-stage renal failure, requiring dialysis
- Bone marrow failure, e.g. severe thrombocytopaenia, platelet count < 50,000/ $\mu$ L
- Stroke in the last 6 months



# MASTER DAPT Trial

**Screened Population:** HBR pts, treated exclusively with Ultimaster stent, with no restriction based on clinical presentation or PCI complexity

## Randomization and Regimens

### 30 (+14) Days after PCI

Free from cardiac and cerebral ischemic events  
and active bleeding  
No further revascularization planned



### Abbreviated DAPT

#### Immediate DAPT discontinuation

followed by SAPT for 11 months  
or 5 months if OAC is indicated

### Standard DAPT

DAPT for  $\geq 2$  or 5 months in pts with  
or without OAC indication, respectively

followed by SAPT up to 11 months

# High Bleeding Risk Definition

**Patients are at high bleeding risk if at least one of the following criteria applies:**

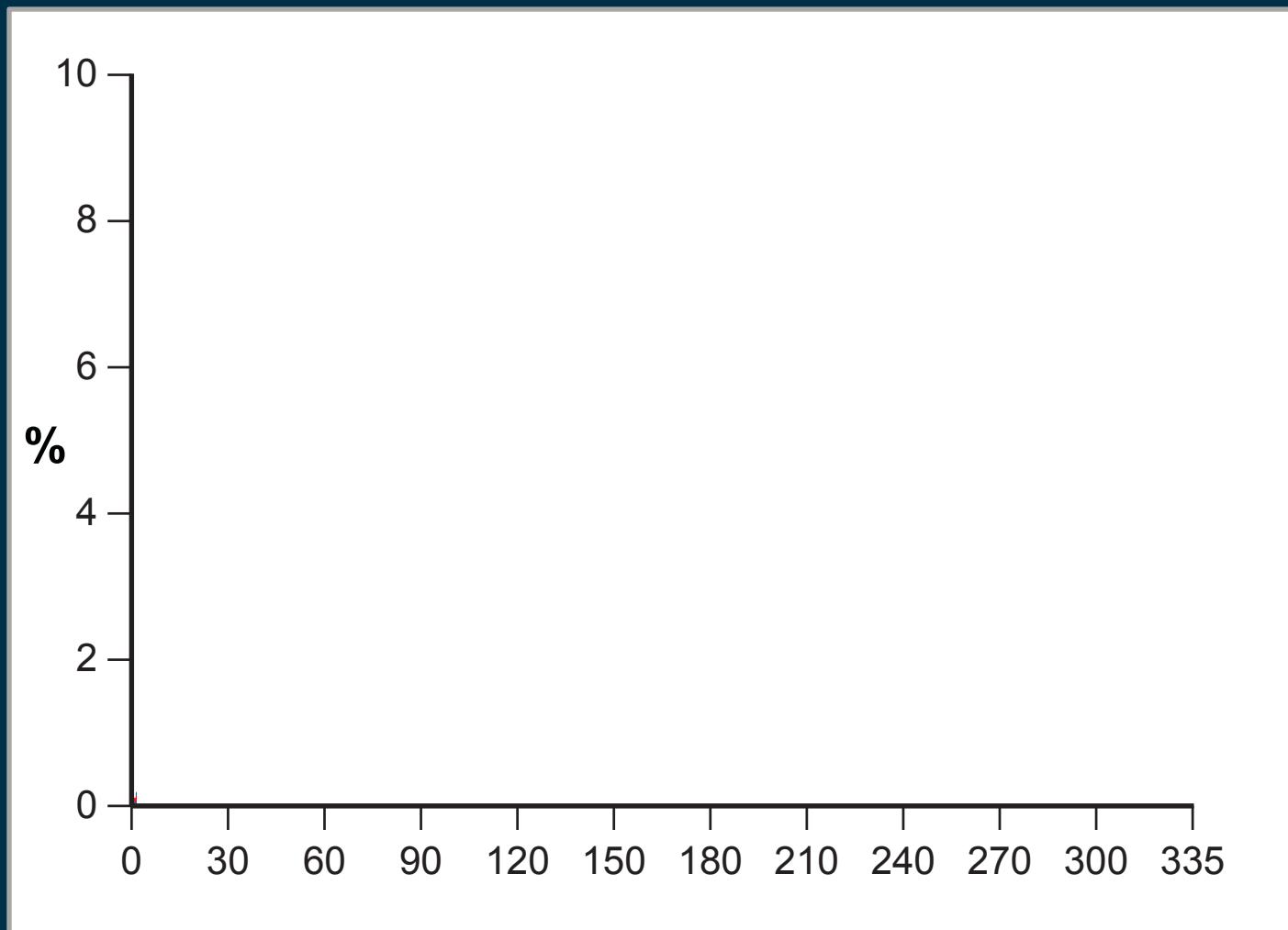
1. **Clinical indication to oral anticoagulants (OAC) for at least 12 months**
2. **Recent (<12 months) non-access site bleeding episode(s), which required medical attention**
3. **Previous bleeding episode(s) which required hospitalization if the underlying cause has not been definitively treated (i.e. surgical removal of the bleeding source)**
4. **Age  $\geq 75$  years**
5. **Systemic conditions associated with an increased bleeding risk**
6. **Documented anemia ( $Hb < 11$  g/dL) or transfusion within 4 weeks before randomization**
7. **Need for chronic treatment with steroids or non-steroidal anti-inflammatory drugs**
8. **Diagnosed malignancy (other than skin) considered at high bleeding risk**
9. **Stroke at any time or transient ischemic attack (TIA) in the previous 6 months**
10. **PRECISE DAPT score  $\geq 25$**

# Baseline Characteristics and Clinical Presentation

Mean±SD 2.1±1.1 HBR criteria	Abbreviated DAPT (N=2295)	Standard DAPT (N=2284)
Age — yr	76.1±8.7	76.0±8.8
Male sex — no. (%)	1590 (69.3)	1581 (69.2)
Diabetes mellitus — no. (%)	754 (32.9)	784 (34.3)
Prior MI— no. (%)	434 (18.9)	430 (18.8)
Prior PCI— no. (%)	594 (25.9)	594 (26.0)
Prior CVA— no. (%)	268 (11.7)	302 (13.2)
Chronic kidney disease— no. (%)	418 (18.2)	458 (20.1)
Atrial fibrillation — no. (%)	770 (33.6)	720 (31.5)
Oral anticoagulant — no. (%)	849 (37.0)	820 (35.9)
CCS— no. (%)	1167 (50.8)	1201 (52.6)
Non-ST-elevation ACS— no. (%)	855 (37.2)	818 (35.8)
ST-elevation MI— no. (%)	273 (11.9)	265 (11.6)
Killip II, III or IV	252 (11.0)	254 (11.1)

# Net adverse clinical events (NACE)

Per protocol population



Non-inferiority Analysis

Difference in cumulative incidence, -0.23

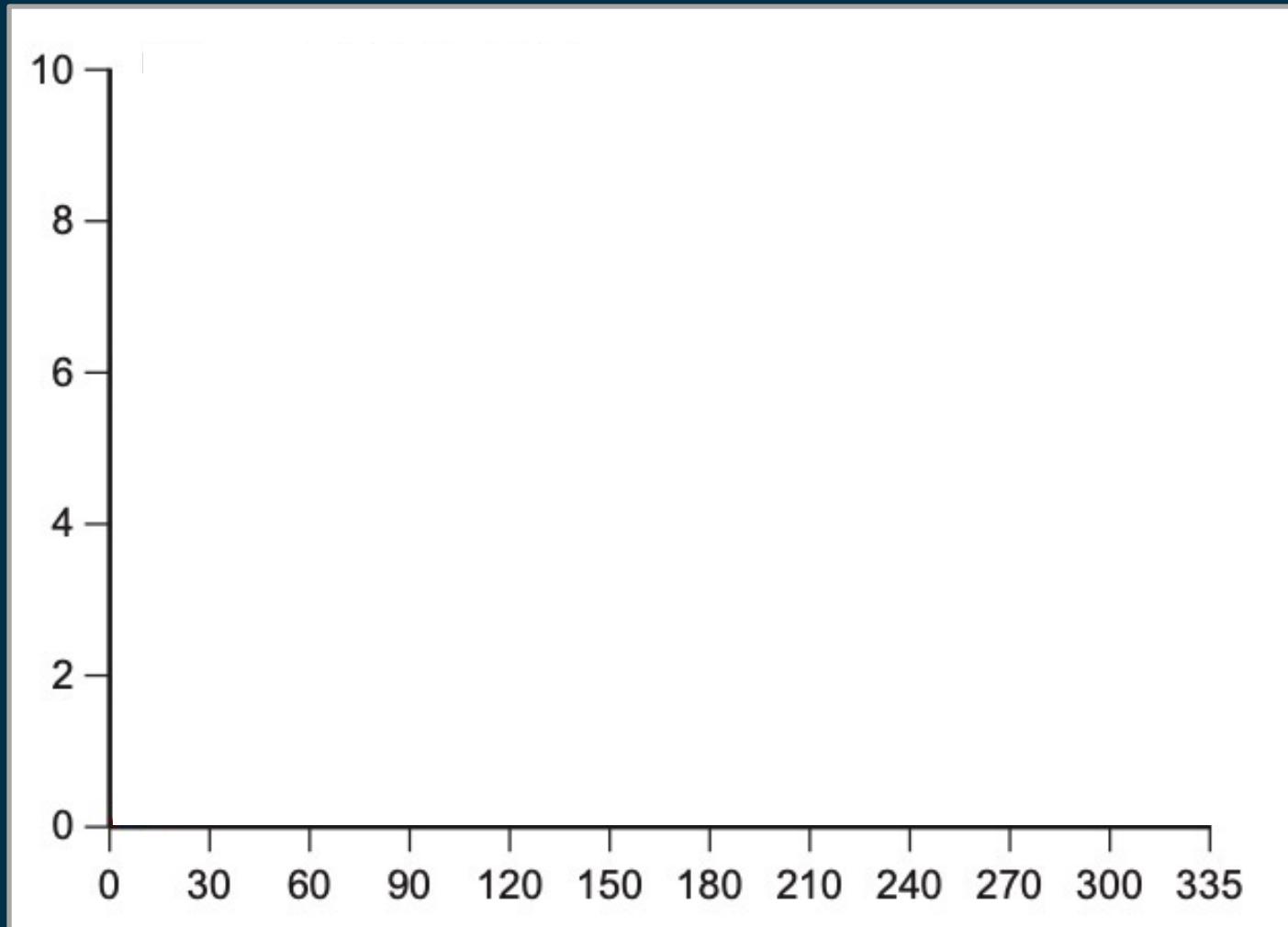
Non-inferiority margin: 3.6%

Two-sided 95%  
UCL: 1.33%  
 $P_{non-inferiority} < 0.001$

NACE: All-cause death, MI, stroke, and major bleeding events defined as BARC 3 or 5

# Major adverse cardiac and cerebral events (MACCE)

Per protocol population



Non-inferiority Analysis

Difference in cumulative incidence, 0.11

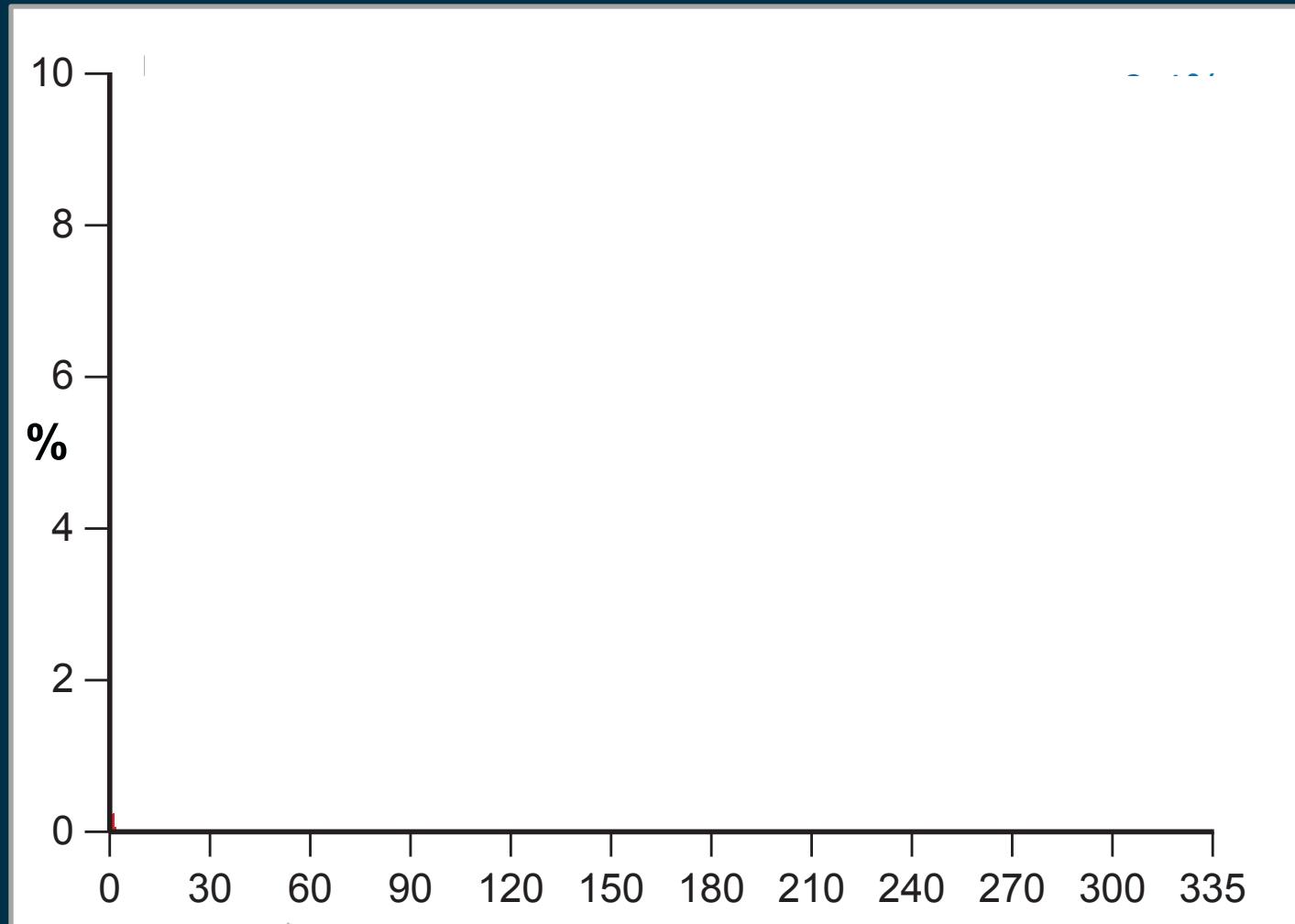
Non-inferiority margin: 2.5%

Two-sided 95%  
UCL: 1.51  
 $P_{non-inferiority} = 0.001$

MACCE: All-cause death, MI, stroke

# Major or clinically relevant nonmajor bleeding

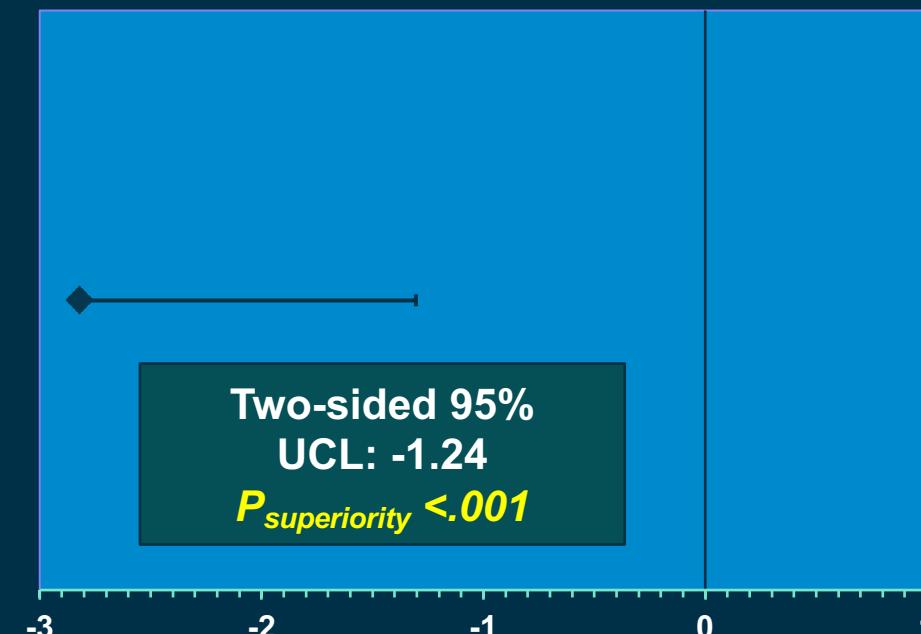
Intention to treat population



Superiority Analysis

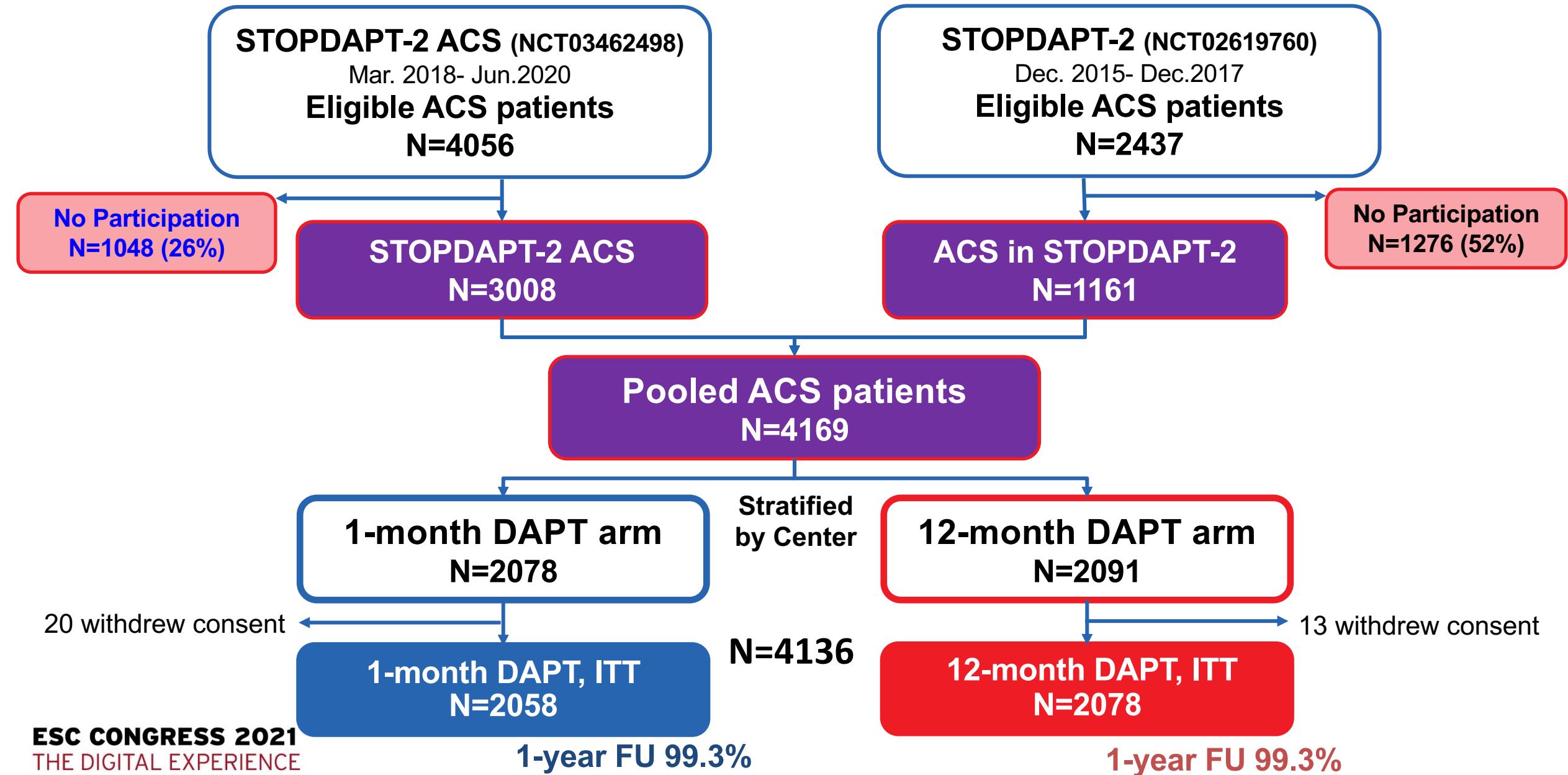
Difference in cumulative incidence, -2.82

Two-sided 95%  
UCL: -1.24  
 $P_{superiority} < .001$



NNTB: 35

# STOP-DAPT2-ACS

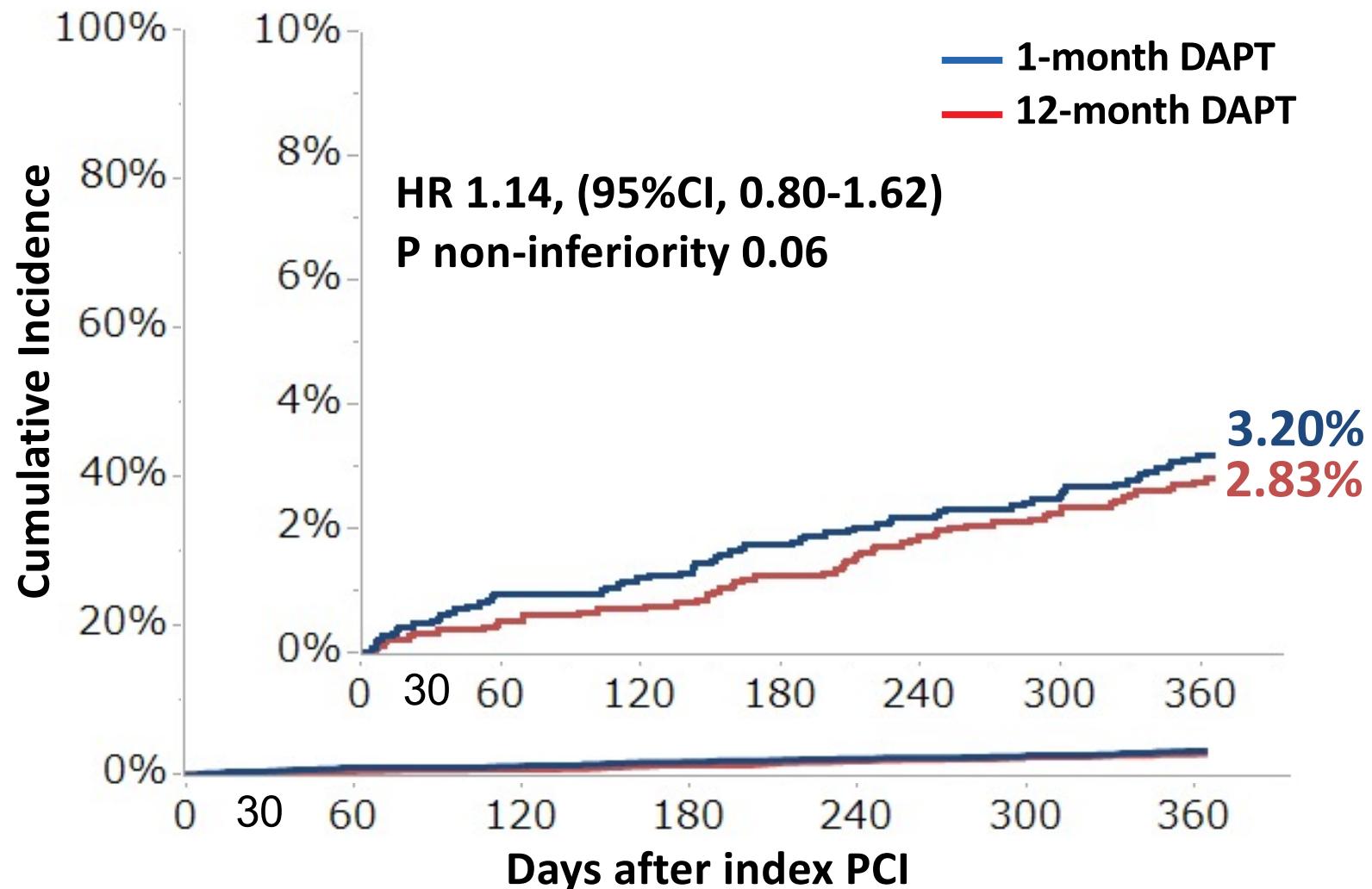


# Baseline Clinical Characteristics

	1-month DAPT N=2058	12-month DAPT N=2078
Age, years	67.0±11.9	66.6±11.9
Men	79%	79%
STEMI; NSTEMI; UA	57%; 19%; 23%	55%; 21%; 24%
Killip 4	3.4%	3.1%
ECMO; Impella; IABP	0.3%; 0.1%; 4.1%	0.3%; 0.1%; 3.1%
Diabetes	30%	30%
Severe CKD (eGFR<30ml/min/m <sup>2</sup> )	3%	3%
Prior MI	6%	5%
Prior PCI	11%	10%
CREDO-Kyoto thrombotic risk score		
High; Intermediate; Low	4%; 17%; 79%	5%; 16%; 79%
CREDO-Kyoto bleeding risk score		
High; Intermediate; Low	4%; 20%; 77%	3%; 19%; 78%

# Primary Endpoint

## CV death/MI/ST/Stroke/TIMI major/minor bleeding



No. at risk

12-month DAPT

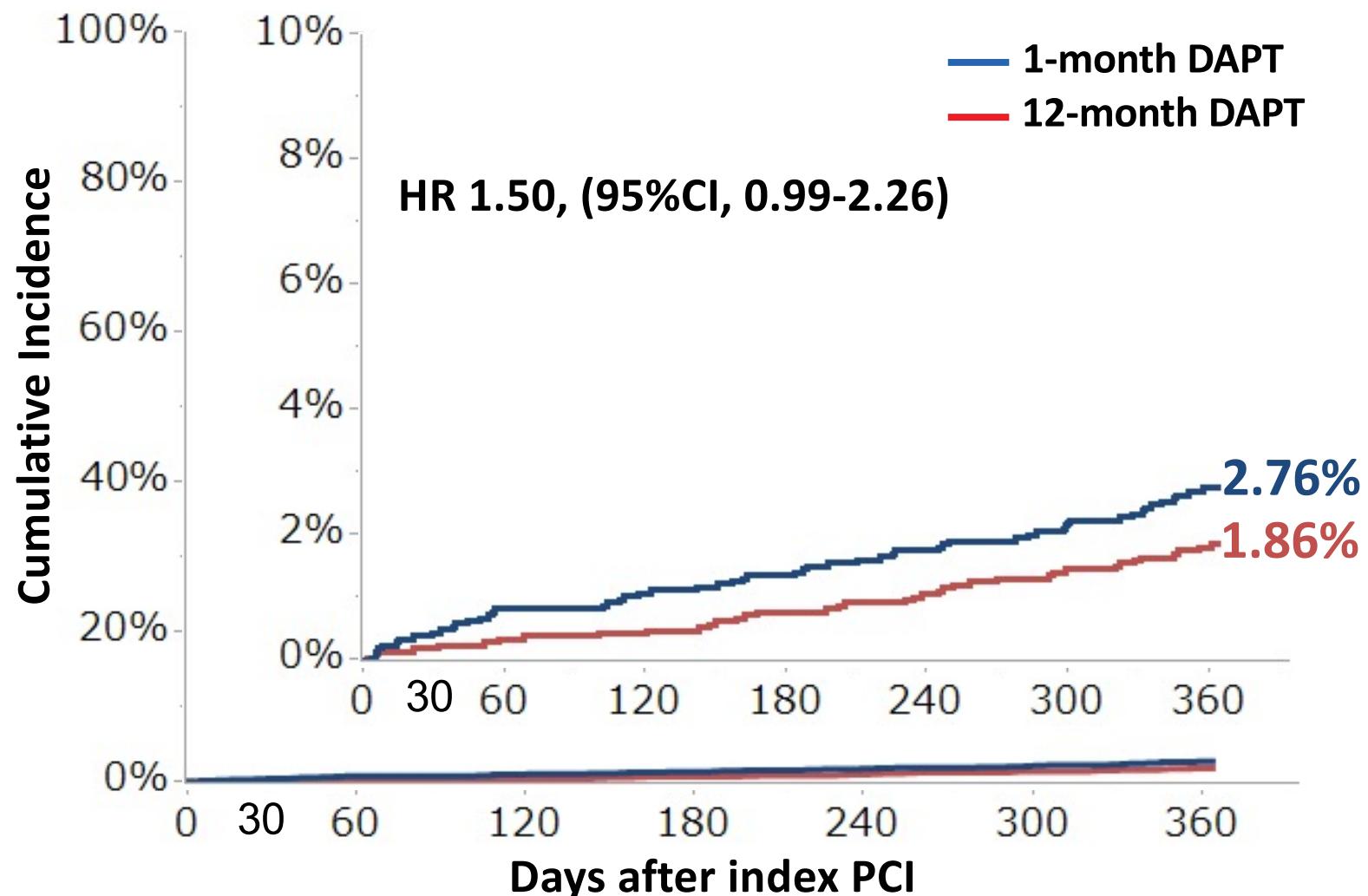
2078 2070 2055 2048 2036 2021 2010 1581

1-month DAPT

2058 2047 2028 2021 2007 1993 1982 1606

# Major Secondary CV Endpoint

## CV death/MI/ST/Stroke



No. at risk

12-month DAPT

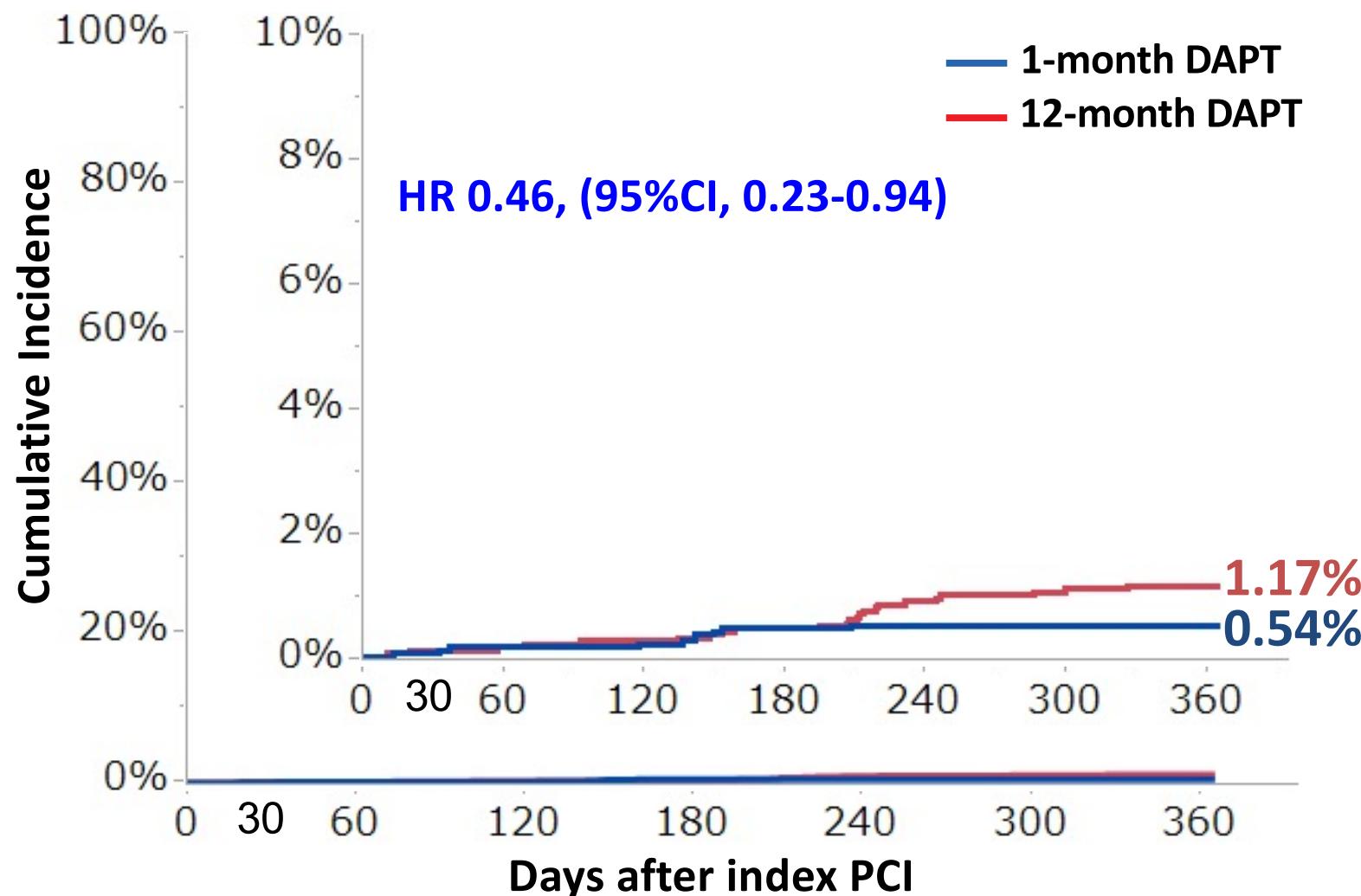
2078 2073 2059 2054 2046 2038 2028 1597

1-month DAPT

2058 2049 2031 2024 2015 2002 1991 1614

# Major Secondary Bleeding Endpoint

## TIMI major/minor bleeding



No. at risk

12-month DAPT

2078 2073 2060 2054 2045 2033 2027 1603

1-month DAPT

2058 2052 2041 2038 2030 2023 2015 1642

# Conclusion



## DAPT + courte après stent?

Oui, mais pas pour tout le monde!



STEMI  
SCA  
Ht risque ischemique  
Ht risque thrombotique