

Printemps 2018 de la médecine d'urgence

Nouveautés dans la prise en charge du STEMI

Eric BONNEFOY (Lyon)

Conflicts of interest

Soutien d'enseignement par AZ
sinon je n'ai pas de conflit d'intérêt

CHANGE IN RECOMMENDATIONS

2012

2017

Radial access^a

MATRIX¹⁴³

DES over BMS

EXAMINATION^{150, 151}
COMFORTABLE-AMI¹⁴⁹, NORSTENT¹⁵²

Complete Revascularization^b

PRAMI¹⁶⁸, DANAMI-3-PRIMULTI¹⁷⁰,
CVLPRIT¹⁶⁹, Compare-Acute¹⁷¹

Thrombus Aspiration^c

TOTAL¹⁵⁹, TASTE¹⁵⁷

Bivalirudin

MATRIX²⁰⁹, HEAT-PPCI²⁰⁵

Enoxaparin

ATOLL^{200, 201}, Meta-analysis²⁰²

Early Hospital Discharge^d

Small trials & observational data^{259–262}Oxygen when
 $\text{SaO}_2 < 95\%$ AVOID⁶⁴,
DETO2X⁶⁶Oxygen when
 $\text{SaO}_2 < 90\%$ Dose i.V. TNK-tPA
same in all patientsSTREAM¹²¹Dose i.V. TNK-tPA
half in Pts ≥ 75 years

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- Additional lipid lowering therapy if LDL $> 1.8 \text{ mmol/L}$ (70 mg/dL) despite on maximum tolerated statins IMPROVE-IT³⁷⁶, FOURIER³⁸²

- Complete revascularization during index primary PCI in STEMI patients in shock
Expert opinion

- Cangrelor if P2Y₁₂ inhibitors have not been given CHAMPION¹⁹³
- Switch to potent P2Y₁₂ inhibitors 48 hours after fibrinolysis
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- Extend Ticagrelor up to 36 months in high-risk patients PEGASUS-TIMI 54³³³
- Use of polypill to increase adherence FOCUS³²³
- Routine use of deferred stenting DANAMI 3-DEFER¹⁵⁵

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- Definition of “time 0” to choose reperfusion strategy (i.e. the strategy clock starts at the time of “STEMI diagnosis”).
- Selection of PCI over fibrinolysis: when anticipated delay from “STEMI diagnosis” to wire crossing is ≤ 120 min.
- Maximum delay time from “STEMI diagnosis” to bolus of fibrinolysis agent is set in 10 min.
- “Door-to-Ballon” term eliminated from guidelines.

TIME LIMITS FOR ROUTINE OPENING OF AN IRA^e:

- 0–12h (Class I); 12–48h (Class IIa); >48h (Class III).

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- Left and right bundle branch block considered equal for recommending urgent angiography if ischemic symptoms.

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- Acute and chronic management presented.

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Early Hospital Discharge^d Small trials & observational data ^{259–262}	
Oxygen when SaO ₂ <95%	AVOID ⁶⁴ , DETO2X ⁶⁶
Dose i.V. TNK-tPA same in all patients	Oxygen when SaO ₂ <90%
STREAM ¹²¹	Dose i.V. TNK-tPA half in Pts \geq 75 years

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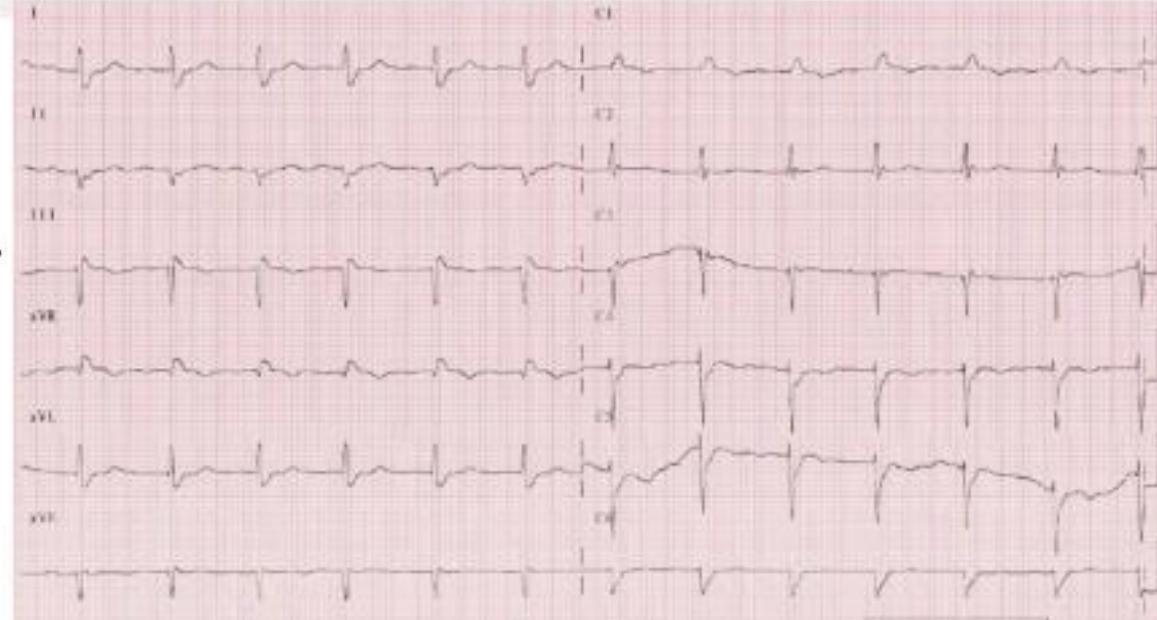
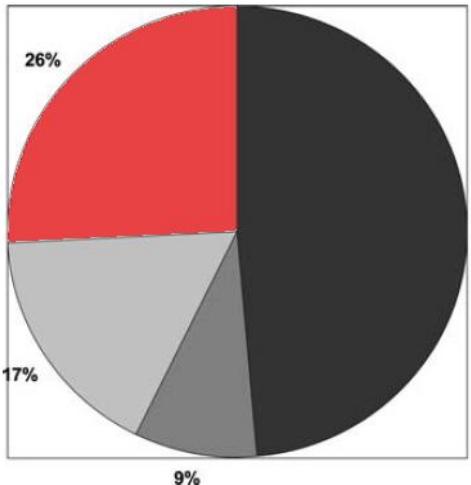
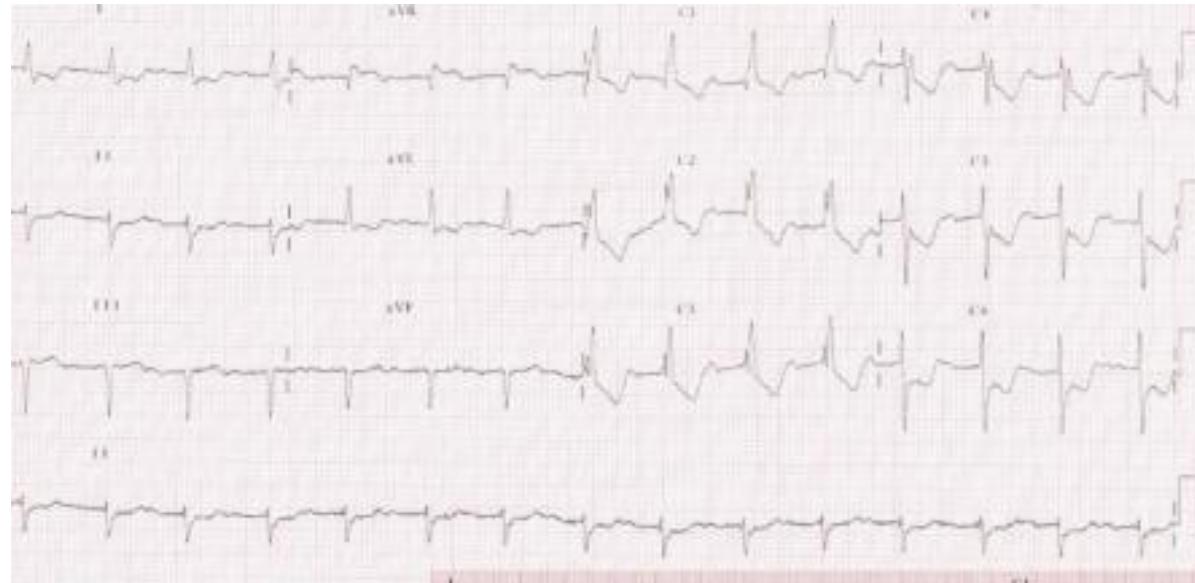
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Infarctus aigu du myocarde avec bloc de branche droit

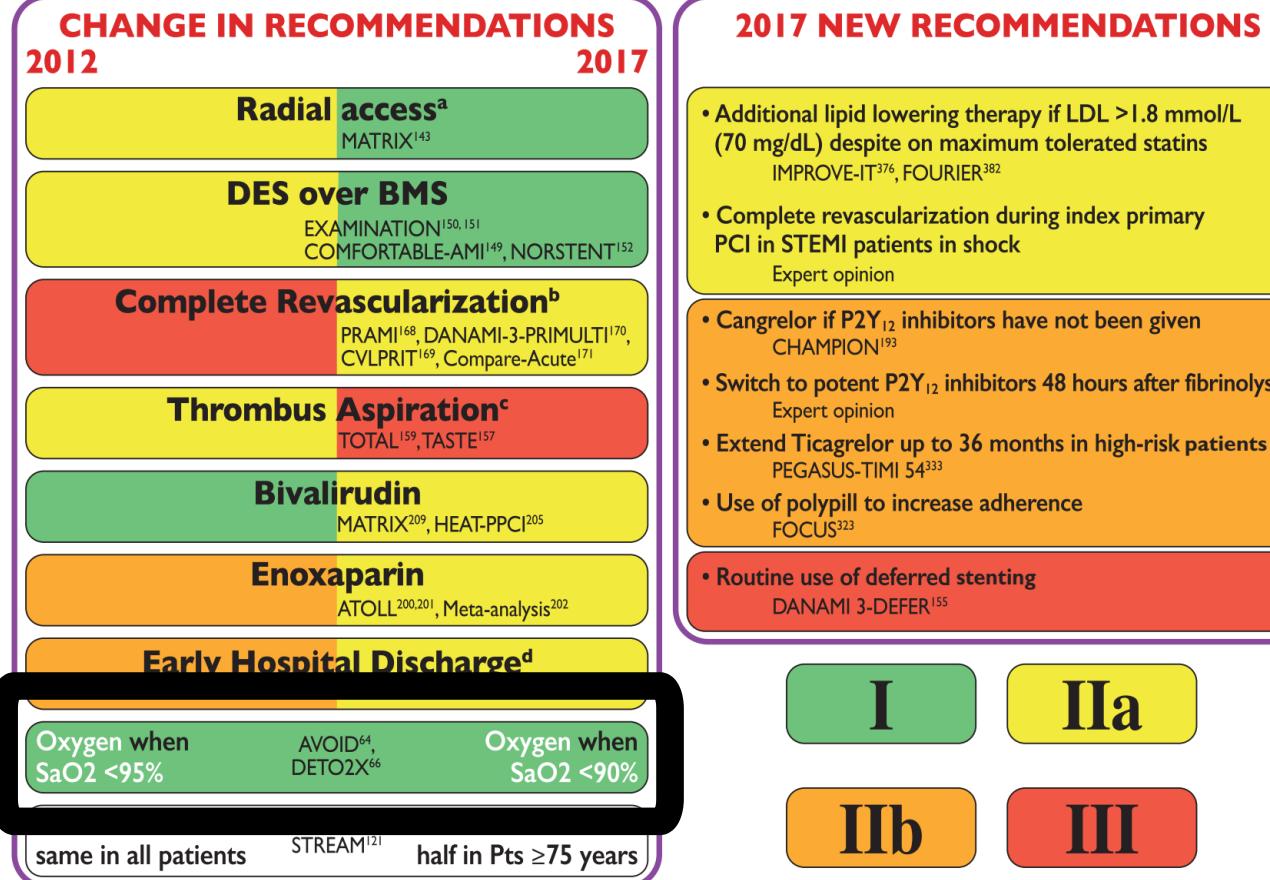


Incidence of cardiogenic shock

STE	6.7 %
LBBB	15.8 %
RBBB	15.4 %

In hospital mortality

STE	5.4 %
LBBB	13.2 %
RBBB	18.8 %

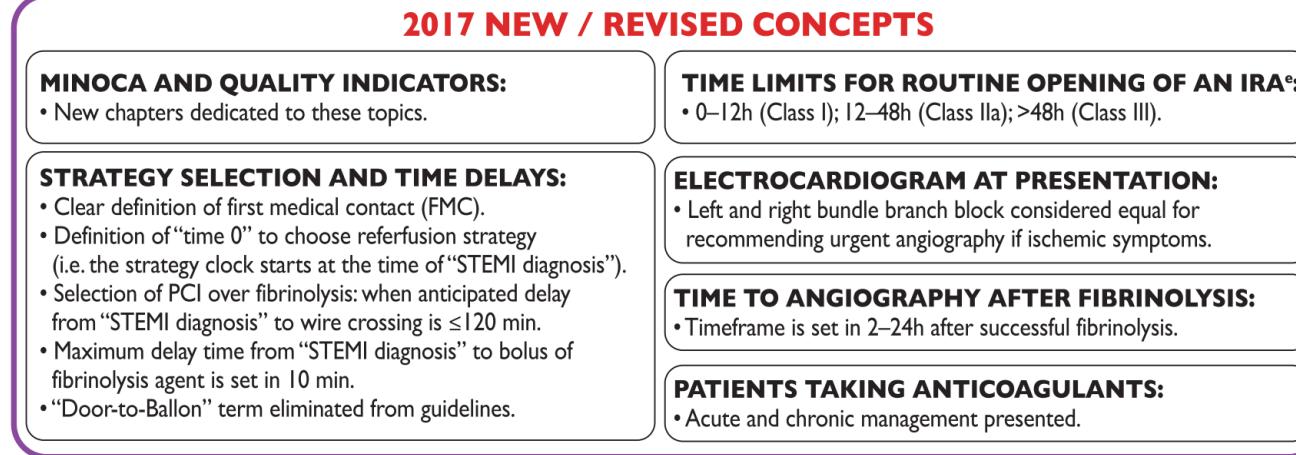


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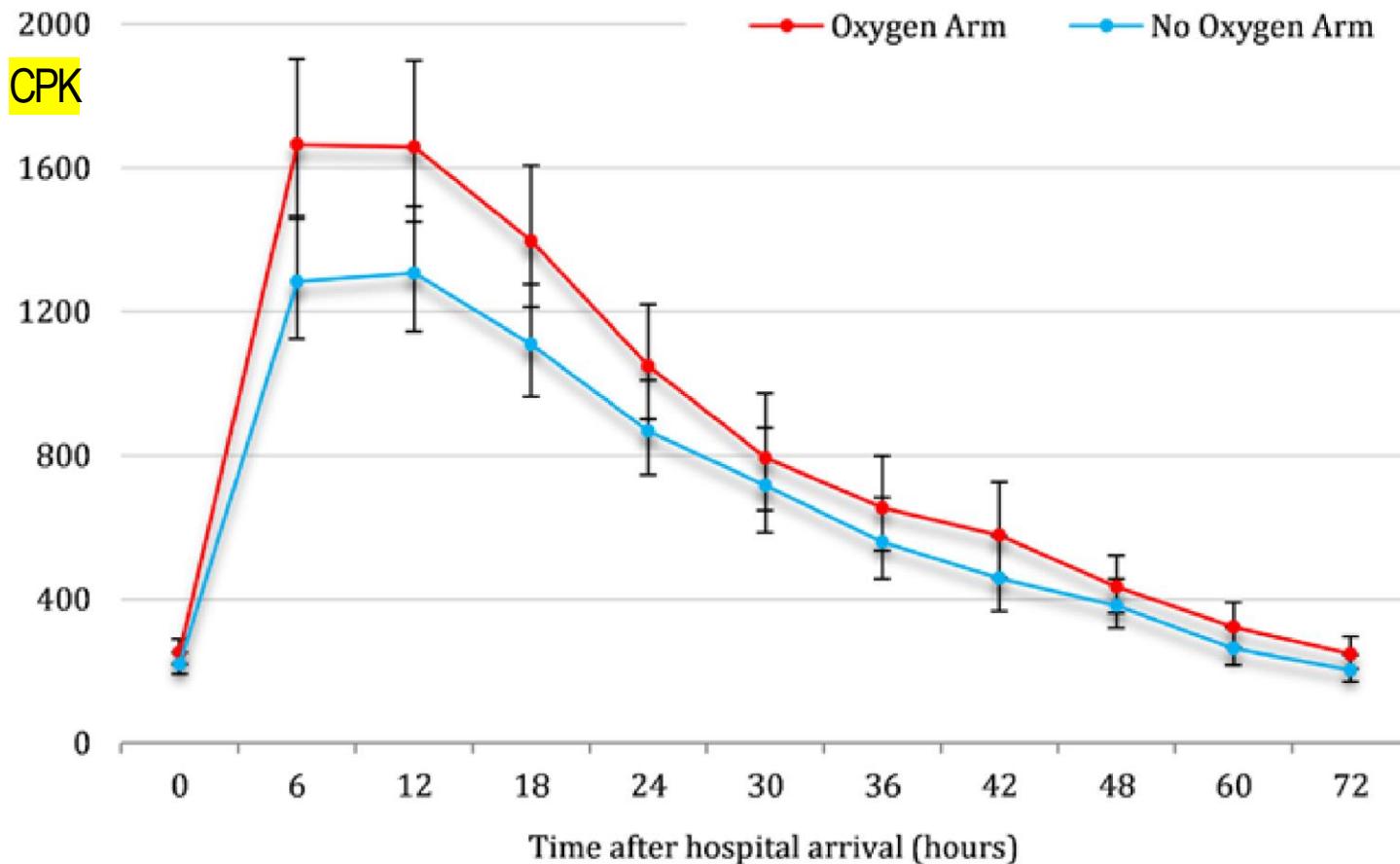


Relief of hypoxaemia and symptoms

Recommendations	Class	Level
Hypoxia		
Oxygen is indicated in patients with hypoxaemia ($\text{SaO}_2 < 90\%$ or $\text{PaO}_2 < 60 \text{ mmHg}$).	I	C
Routine oxygen is not recommended in patients with $\text{SaO}_2 \geq 90\%$.	III	B
Symptoms		
Titrated i.v. opioids should be considered to relieve pain.	IIa	C
A mild tranquilizer (usually a benzodiazepine) should be considered in very anxious patients.	IIa	C

Air ambiant ou O₂ dans l'infarctus du myocarde avec élévation du segment ST

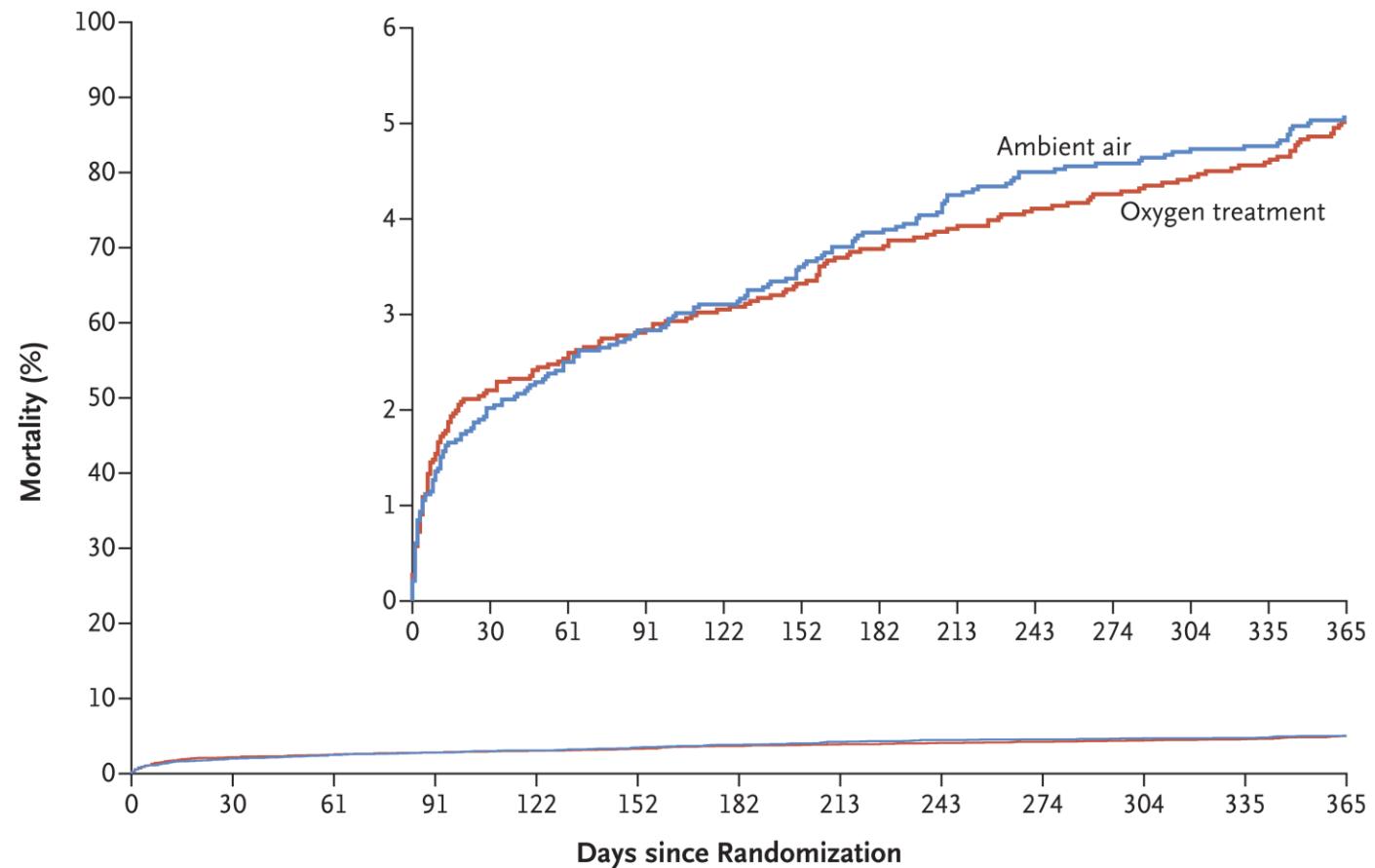
✍ AVOID study, 441 patients with acute STEMI and SaO₂ > 94% were randomized to receive 0 vs. 8 L/min O₂



CPK peak 1948 versus 1543 U/L;
means ratio, 1.27; 95% confidence interval, 1.04–1.52; $P=0.01$

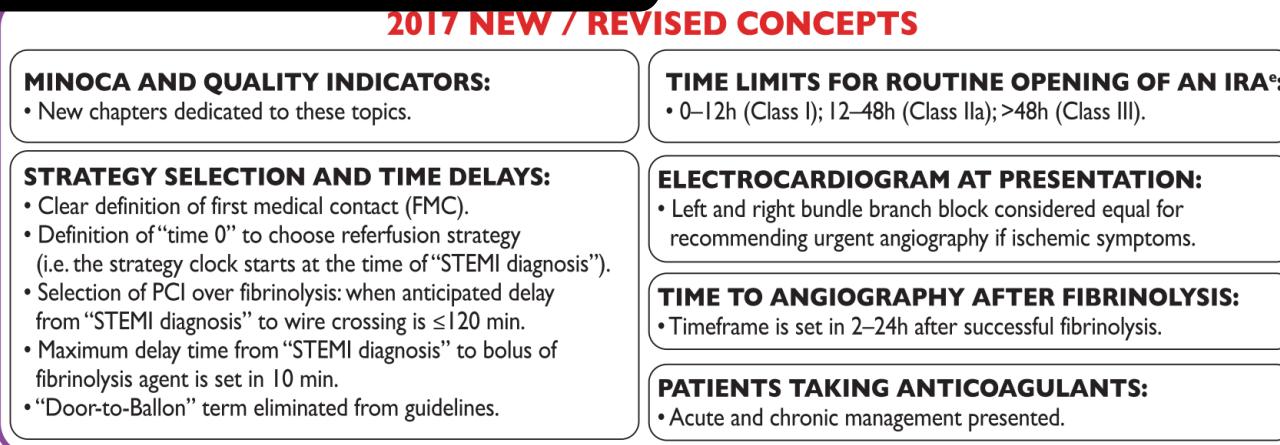
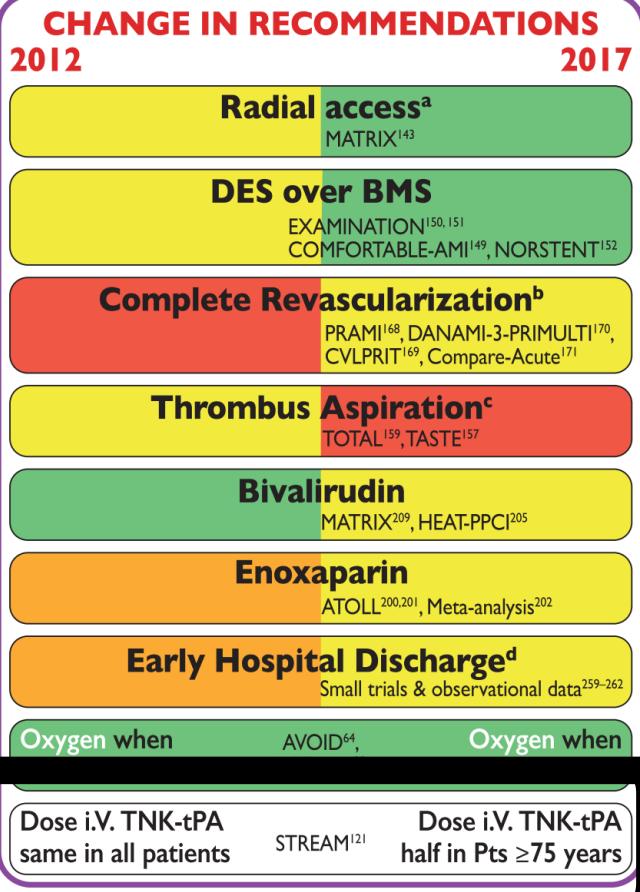
Ellims & al. Circulation. 2015;131:2143-2150

Oxygénothérapie dans l'infarctus aigu du myocarde



Oxygénothérapie dans l'infarctus aigu du myocarde

Timing and End Point	Oxygen Group (N=3311)	Ambient-Air Group (N=3318)	Hazard Ratio (95% CI)	P Value
365 Days after randomization				
Death from any cause — no. (%)	166 (5.0)	168 (5.1)	0.97 (0.79–1.21)	0.80
Rehospitalization with myocardial infarction — no. (%)	126 (3.8)	111 (3.3)	1.13 (0.88–1.46)	0.33
Composite of death from any cause or rehospitalization with myocardial infarction — no. (%)	275 (8.3)	264 (8.0)	1.03 (0.87–1.22)	0.70
30 Days after randomization				
Death from any cause — no. (%)	73 (2.2)	67 (2.0)	1.07 (0.77–1.50)	0.67
Rehospitalization with myocardial infarction — no. (%)	45 (1.4)	31 (0.9)	1.46 (0.92–2.31)	0.11
Composite of death from any cause or rehospitalization with myocardial infarction — no. (%)	114 (3.4)	95 (2.9)	1.19 (0.91–1.56)	0.21
During hospital stay				
Median highest measured level of highly sensitive troponin T (IQR) — ng/liter*	946.5 (243.0–2884.0)	983.0 (225.0–2931.0)	—	0.97



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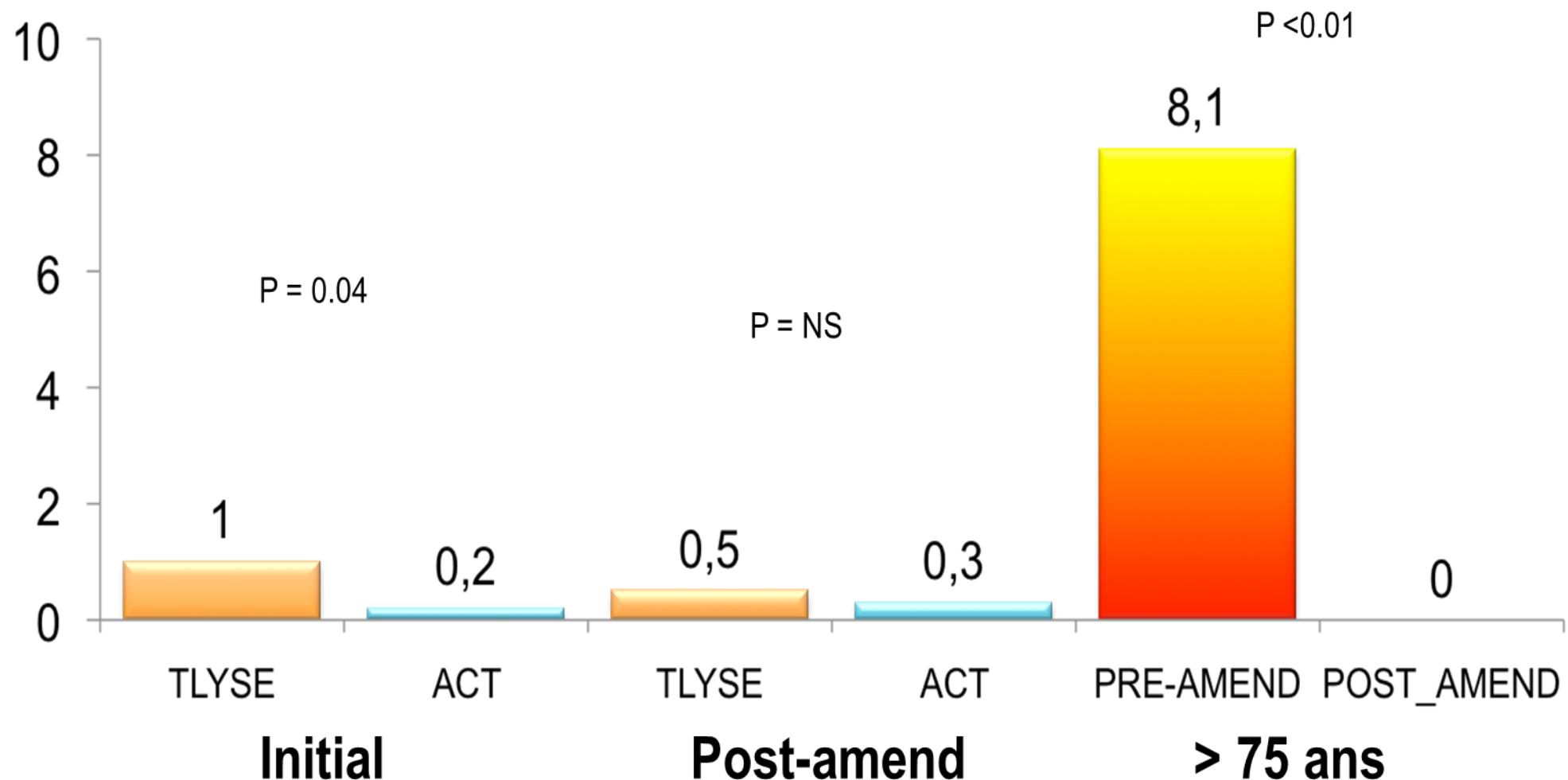
III

Ce que dit l'étude STREAM

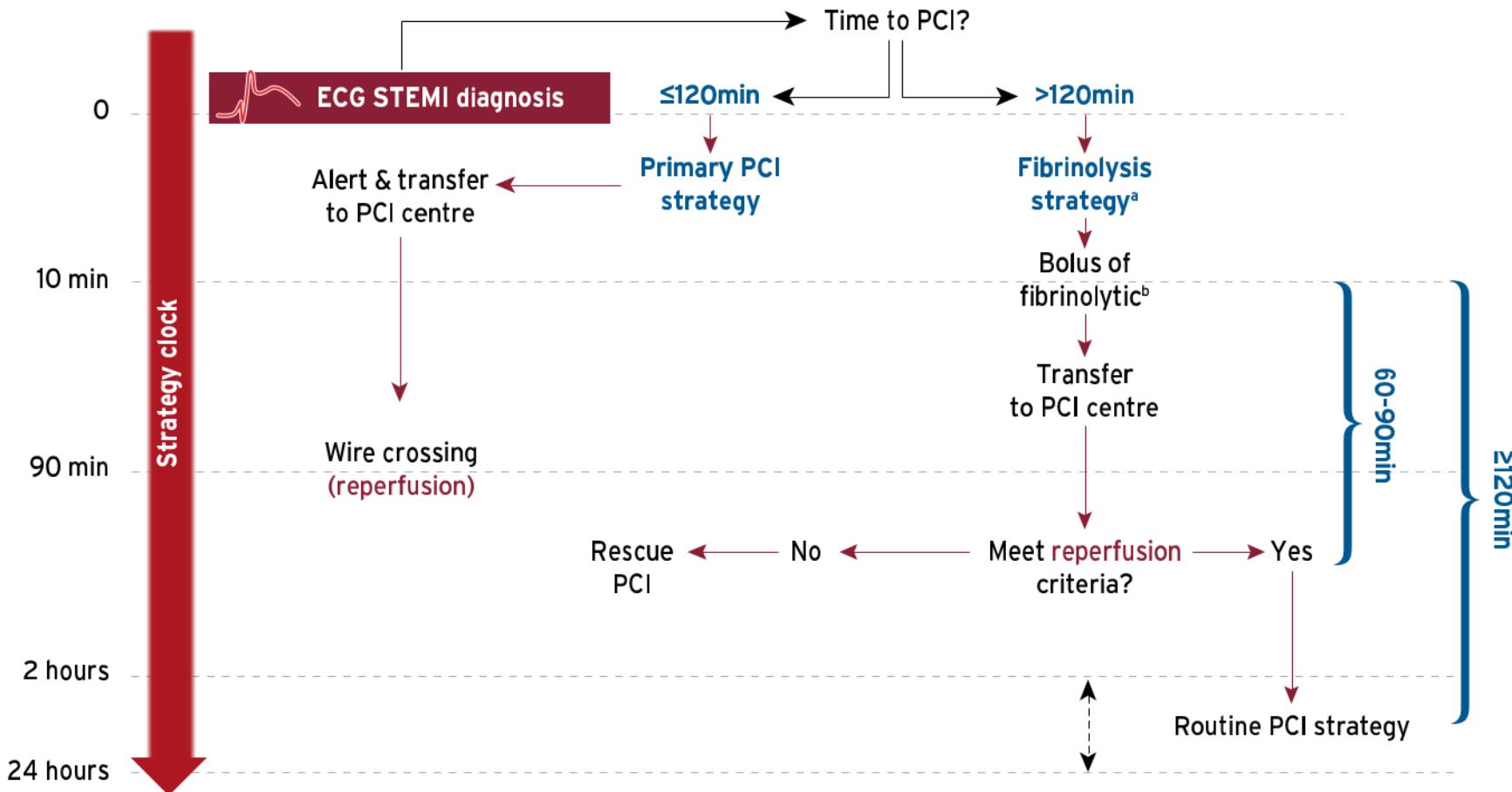
1892 patients. <3H00 début des symptômes et PCI possible seulement >1H00
Fibrinolyse + rescue ou angioplastie dans les 24 heures
Angioplastie primaire

Variable	Fibrinolysis (N = 944)	Primary PCI (N = 948)	P Value
End Point	no./total no. (%)		
Primary composite end point: death, shock, congestive heart failure, or reinfarction at 30 days	116/939 (12.4)	135/943 (14.3)	0.21
Death from any cause	43/939 (4.6)	42/946 (4.4)	0.88
Cardiogenic shock	41/939 (4.4)	56/944 (5.9)	0.13
Congestive heart failure	57/939 (6.1)	72/943 (7.6)	0.18
Reinfarction	23/938 (2.5)	21/944 (2.2)	0.74
Death from cardiovascular causes	31/939 (3.3)	32/946 (3.4)	0.92
Rehospitalization for cardiac causes	45/939 (4.8)	41/943 (4.3)	0.64
Intracranial hemorrhage			
Any	9/939 (1.0)	2/946 (0.2)	0.04

TNK ½ dose >75 ans – Hémorragies cérébrales



Stratégies de reperfusion



^aIf fibrinolysis is contra-indicated, direct for primary PCI strategy regardless of time to PCI.

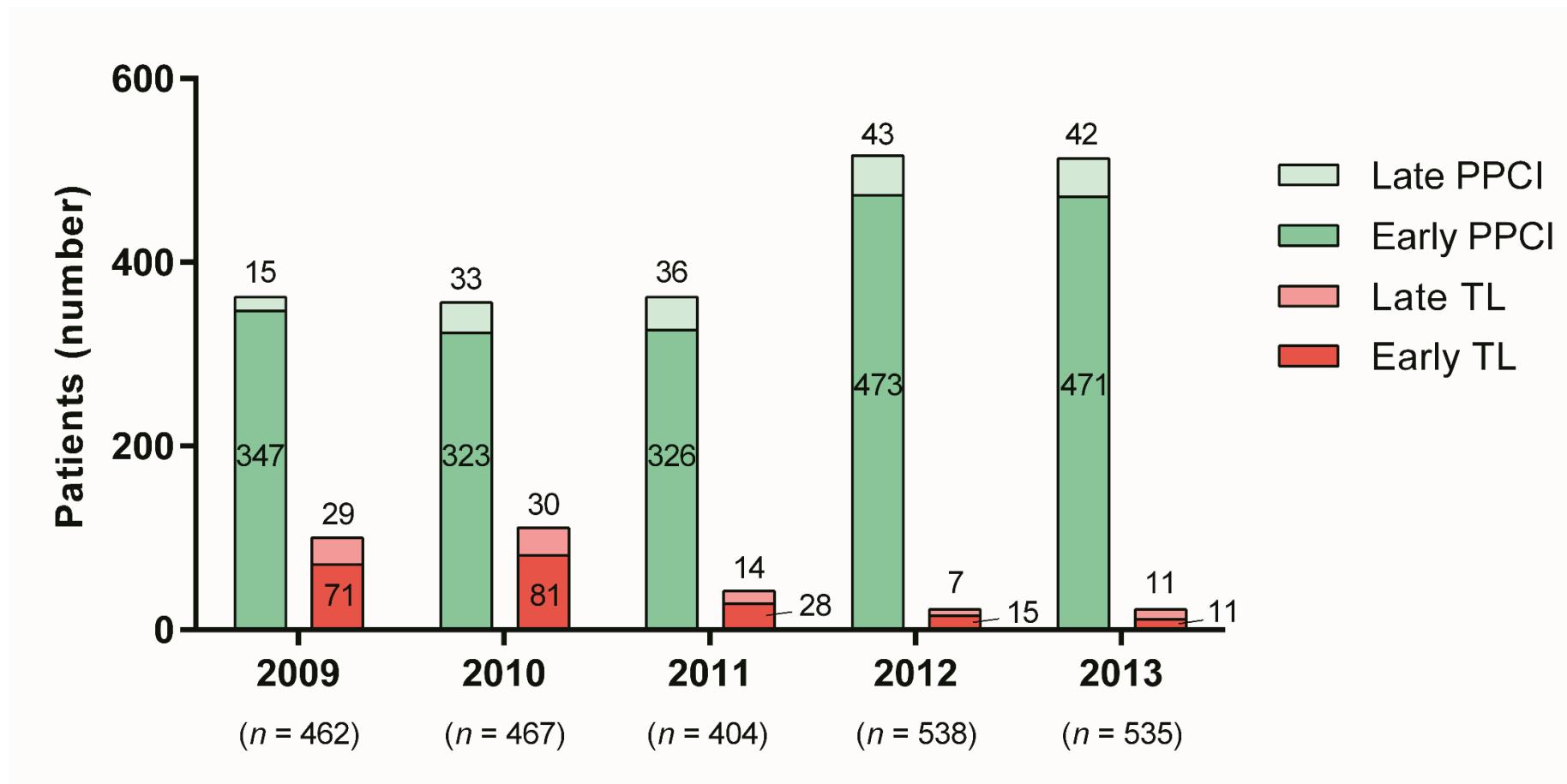
^b10 min is the maximum target delay time from STEMI diagnosis to fibrinolytic bolus administration, however, it should be given as soon as possible after STEMI diagnosis (after ruling out contra-indications).

Reference: Ibañez B. Eur Heart J 2018; 39:119-177.

Web-based pdf file downloadable at: www.escardio.org/ACCA

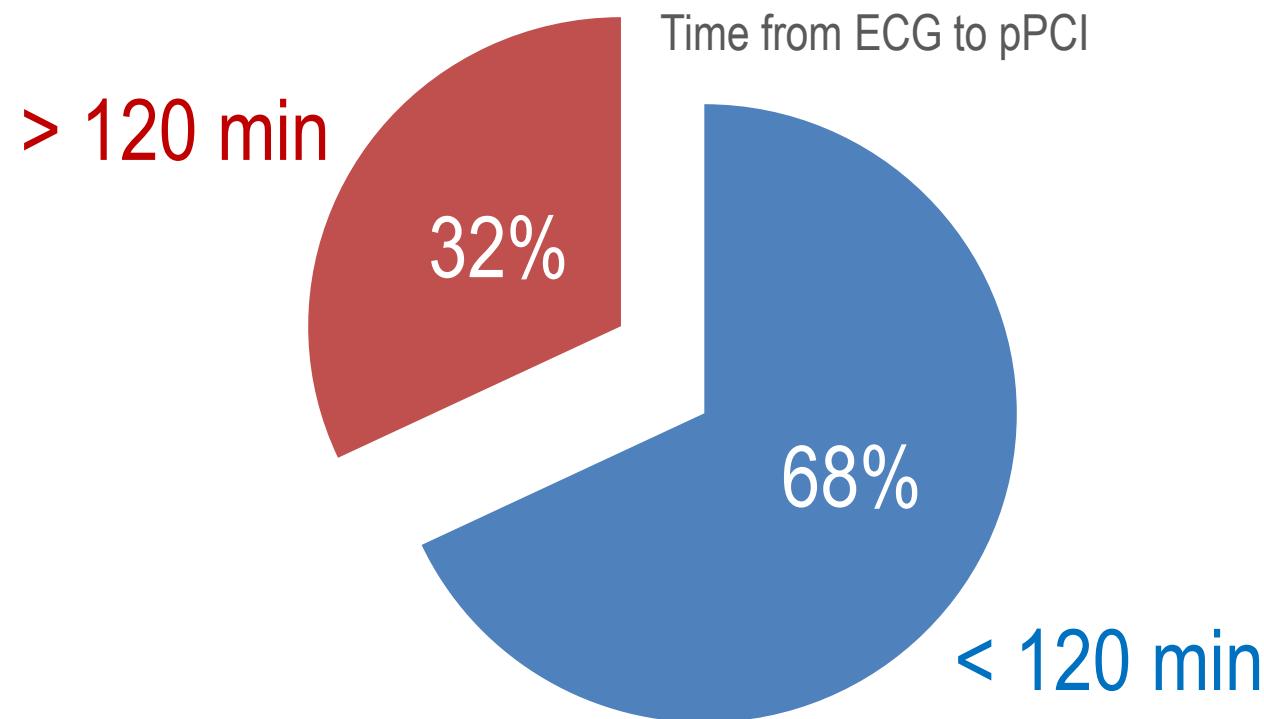
ACCA CLINICAL DECISION MAKING TOOLKIT 2018

Thrombolyse en cours de disparition ?

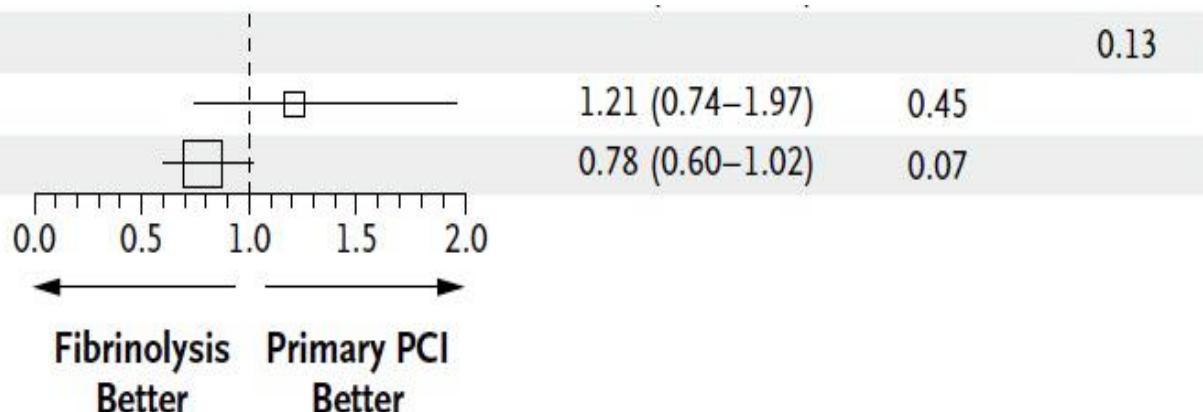
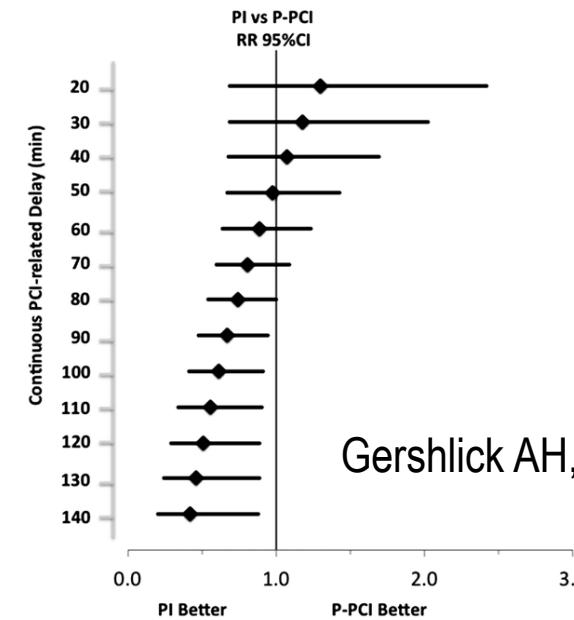
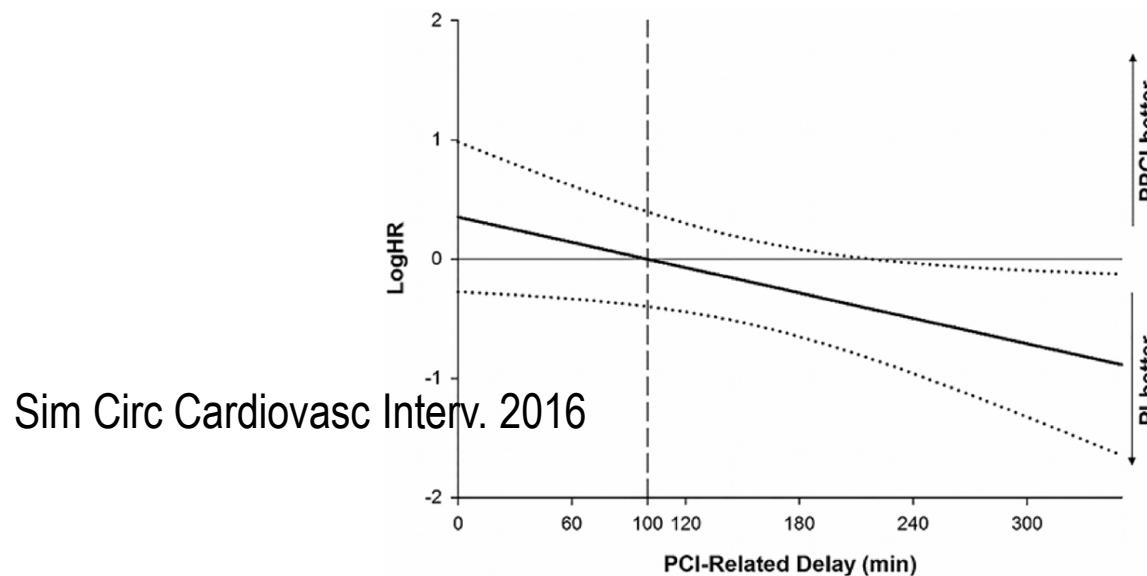


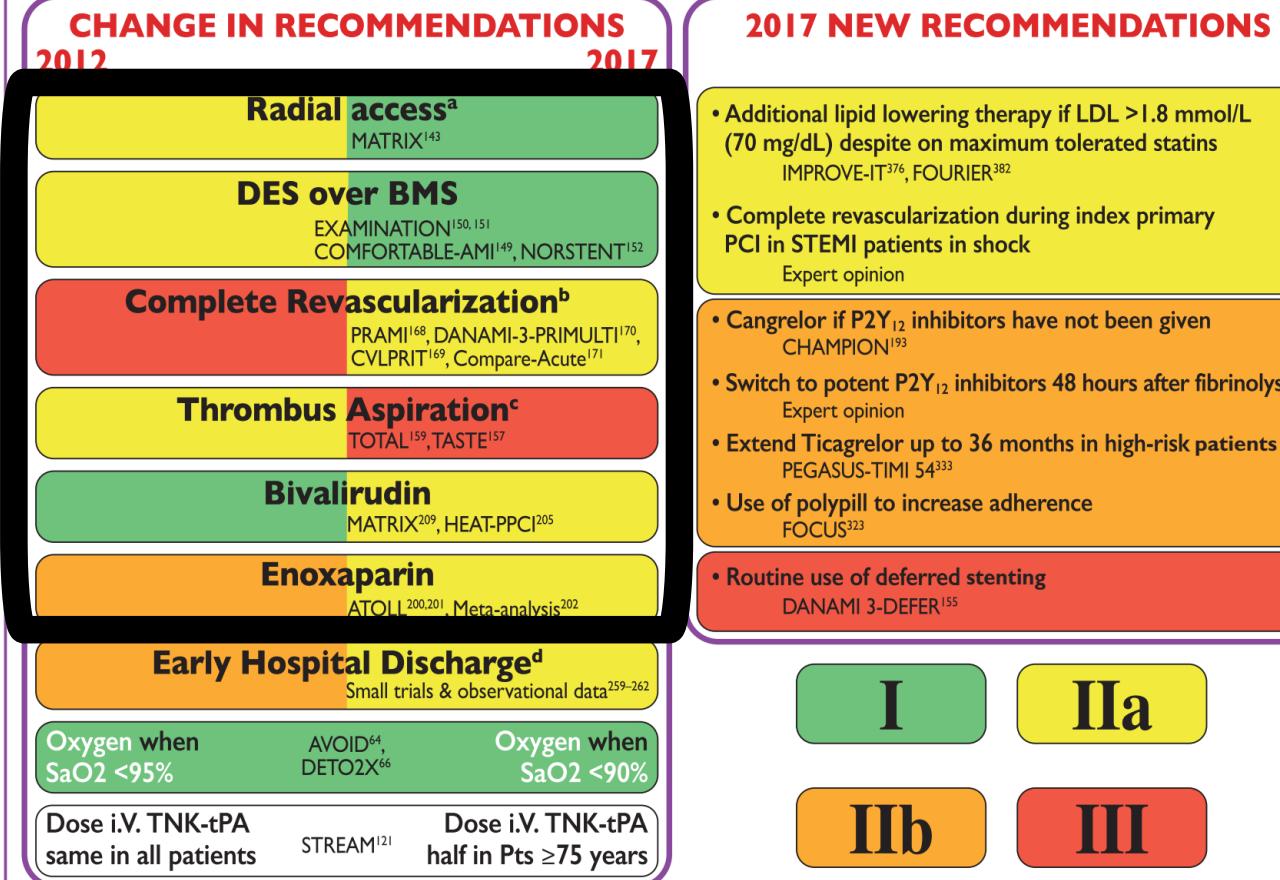


En 2015 en France, un tiers des patients ont été traités par angioplastie primaire au-delà recommandé délais



Bénéfice de la stratégie pharmacoinvasive dans les STEMI vs PPCI





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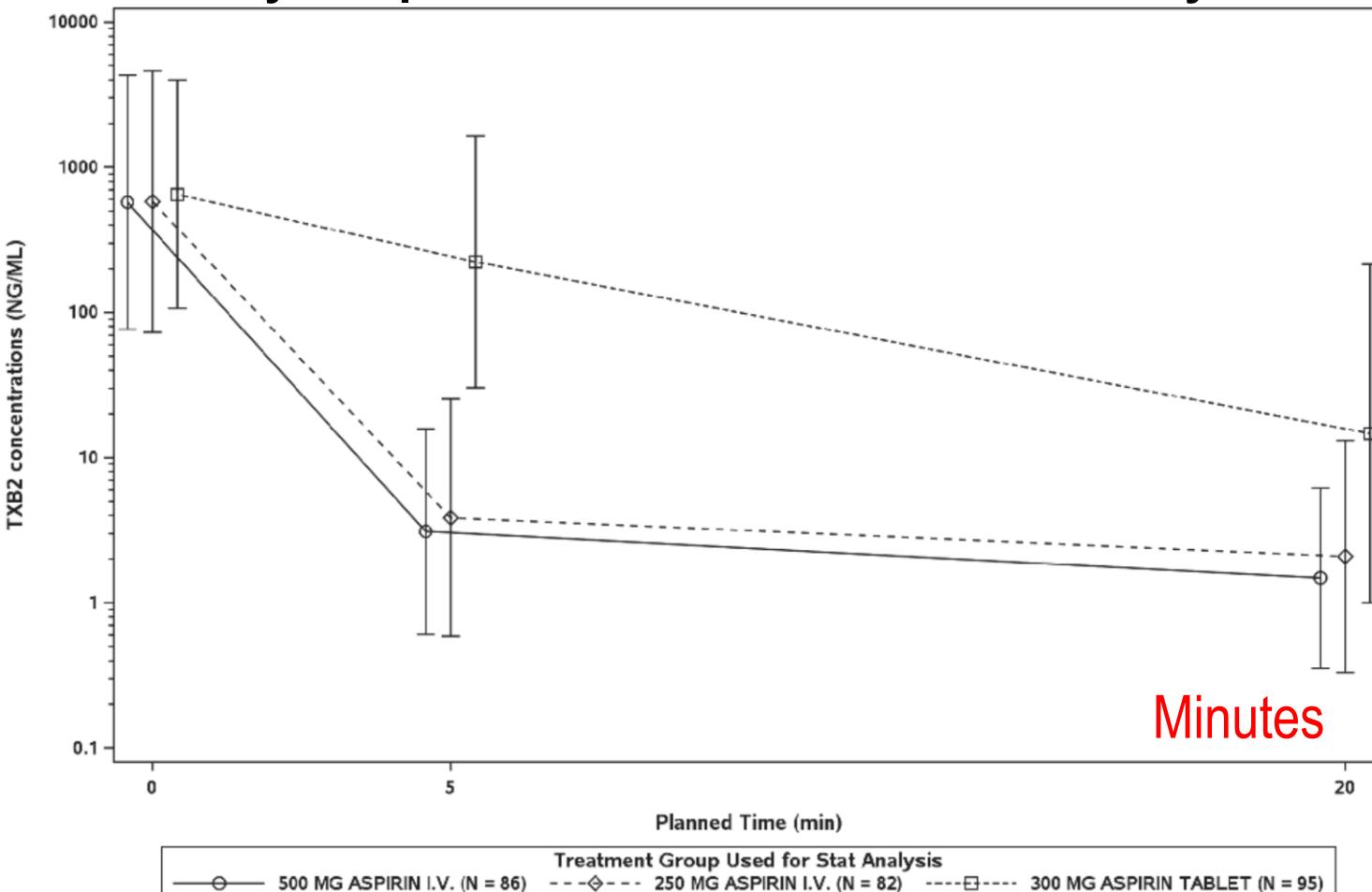
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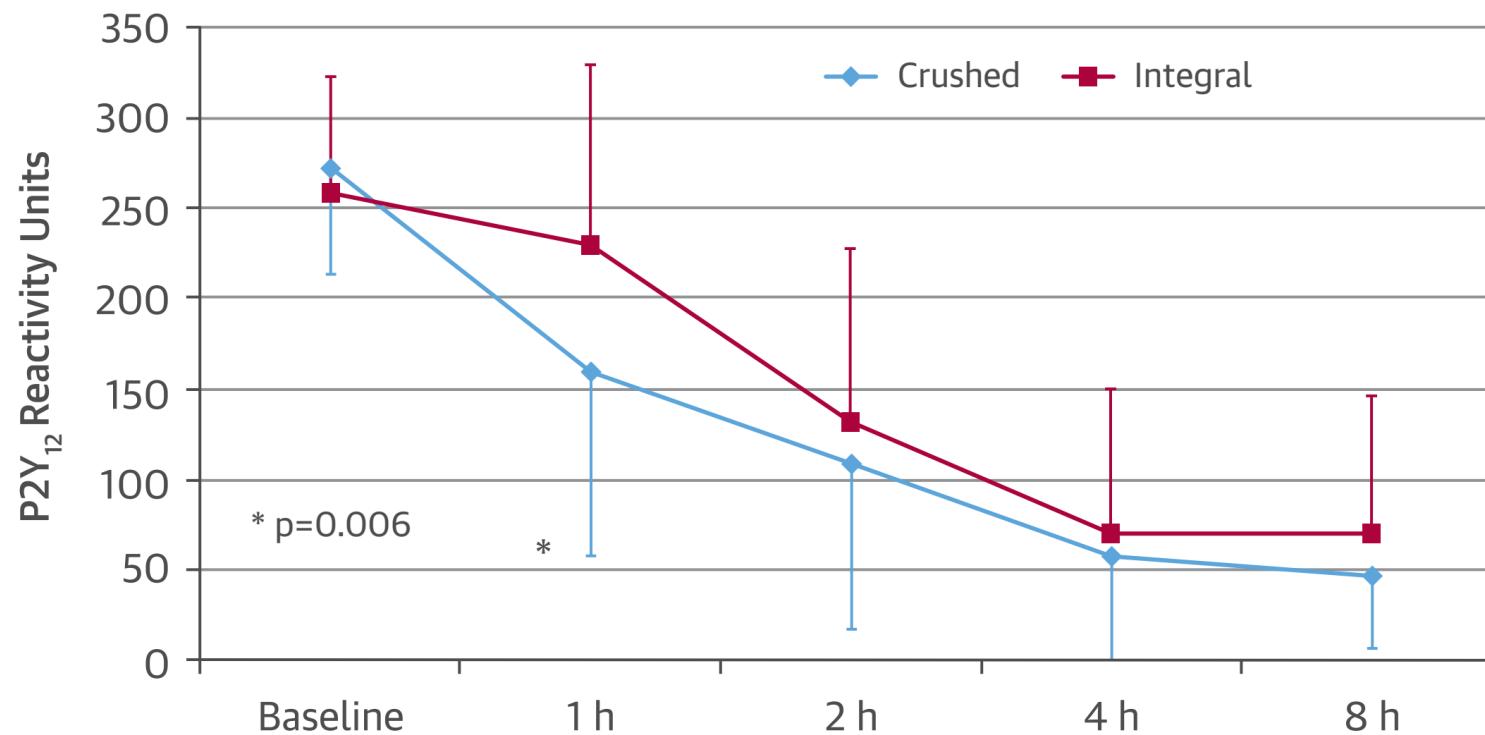
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Inhibition de la synthèse du thromboxane par 500 mg et 250 mg d'acide acétylsalicylique i. v. et 300 mg p. o.

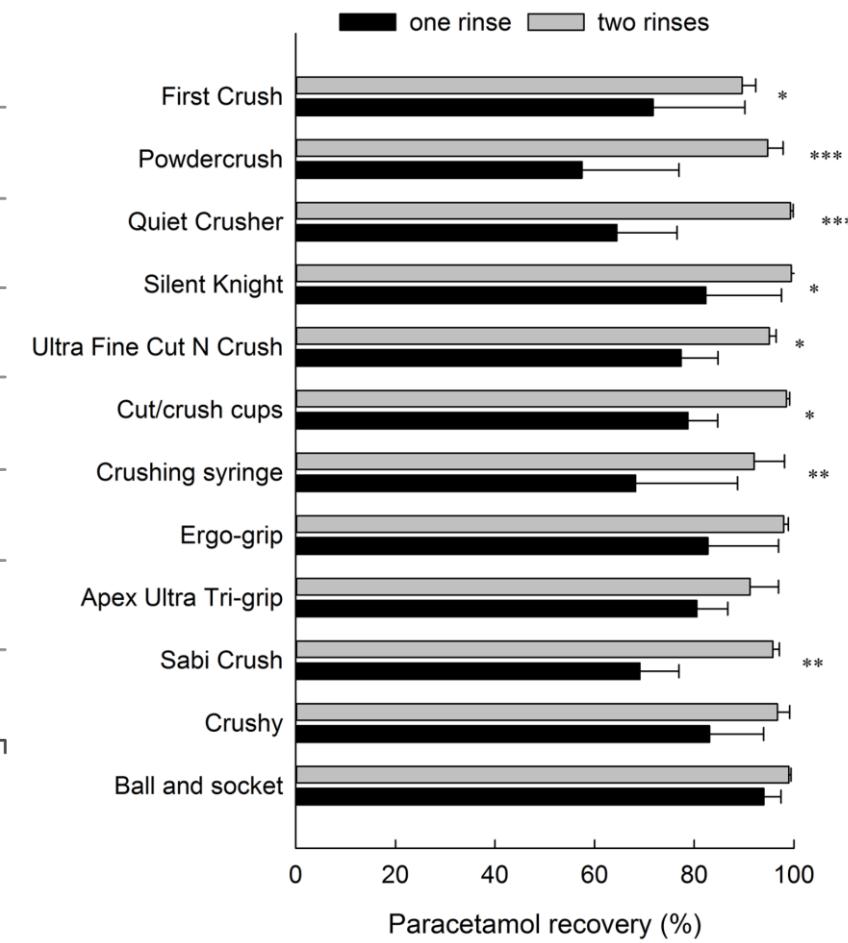
Primary Endpoint: inhibition of thromboxan-synthesis



Administration de comprimés écrasés de ticagrélor chez les patients STEMI - Etude MOJITO

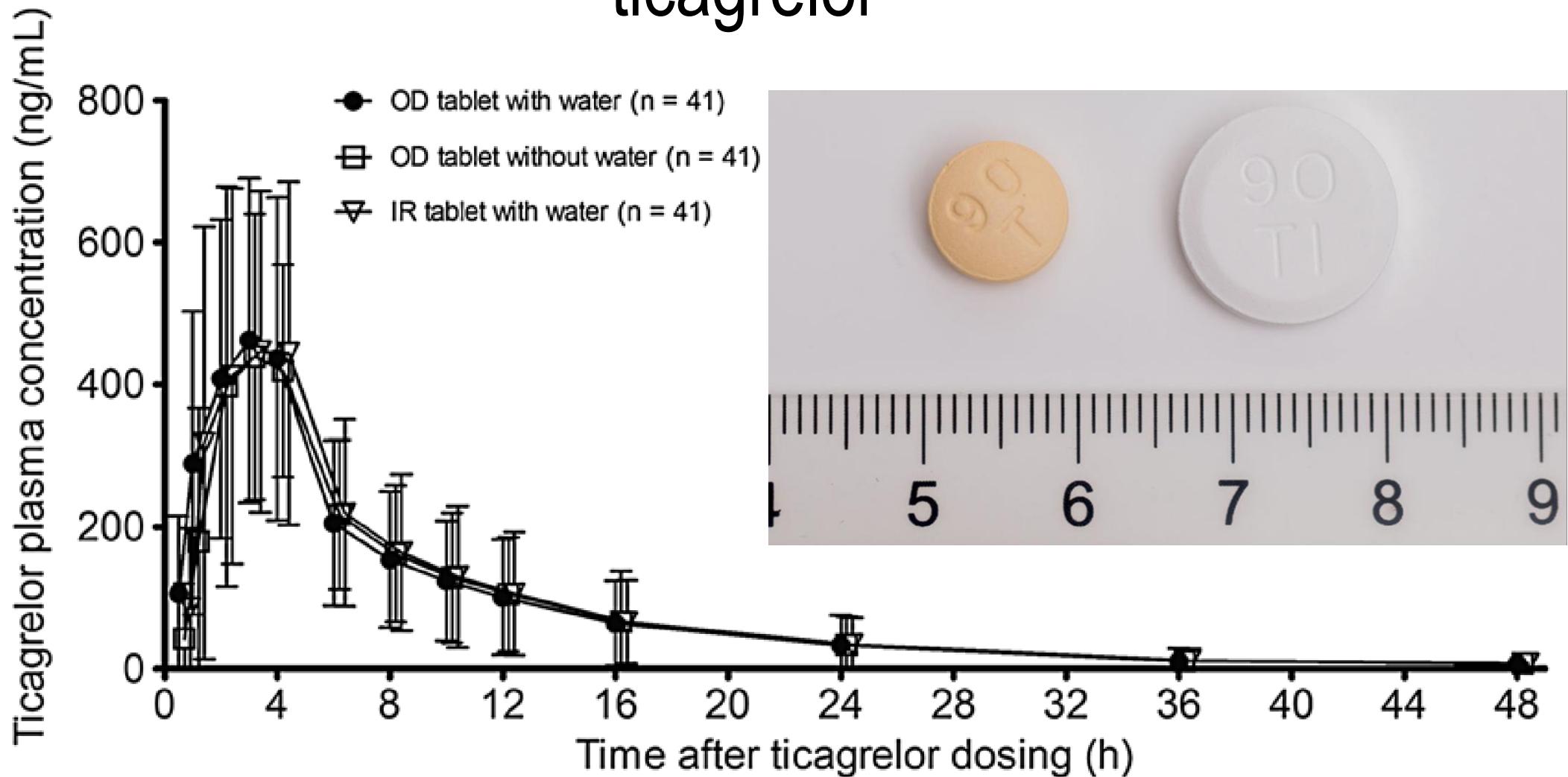


Parodi JACC 2015

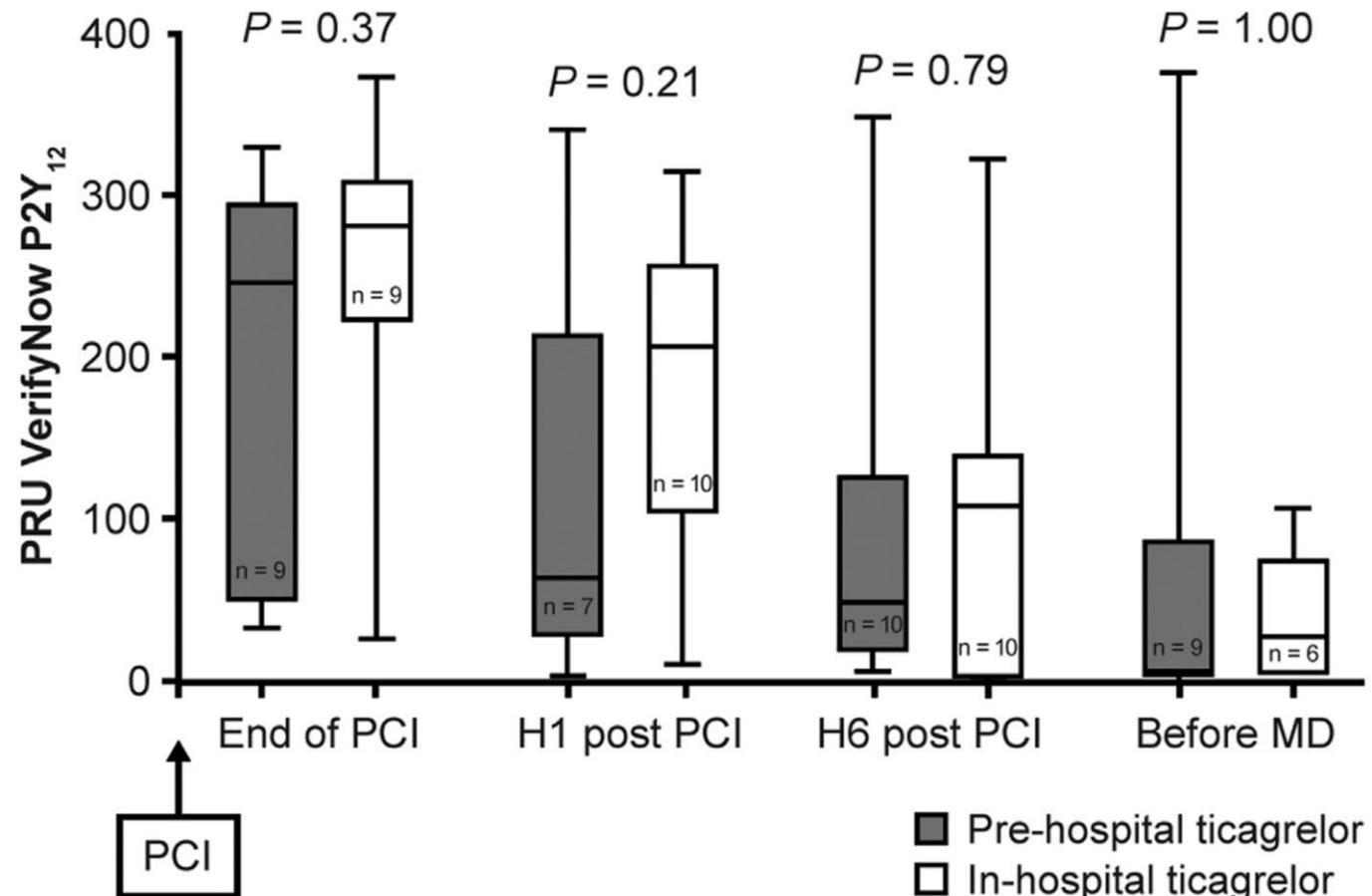


Thong PLOS ONE 2018

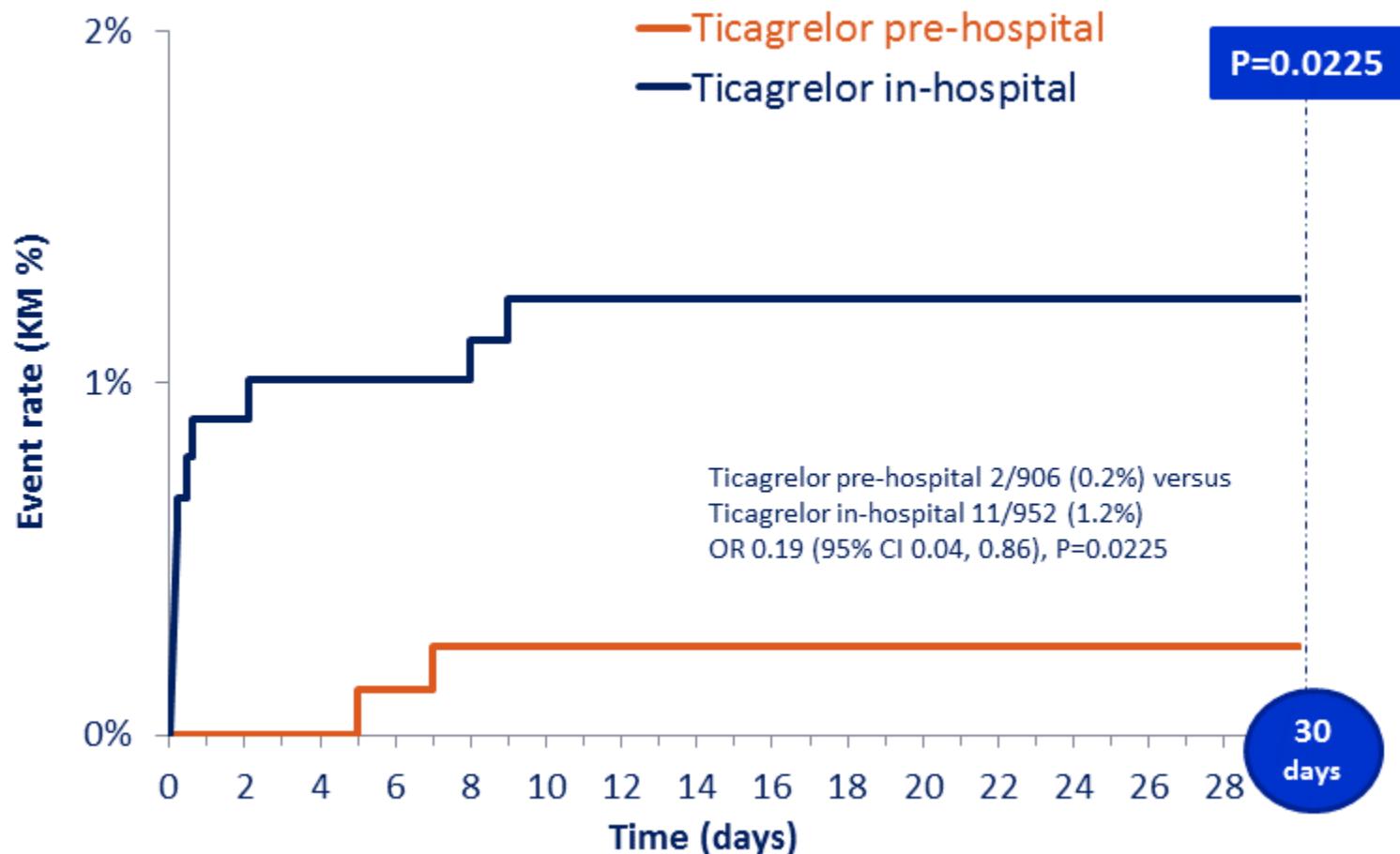
Profils pharmacocinétiques des comprimés de ticagrelor



Inhibition plaquetttaire dans l'étude ATLANTIC



Thrombose de stent – Etude ATLANTIC



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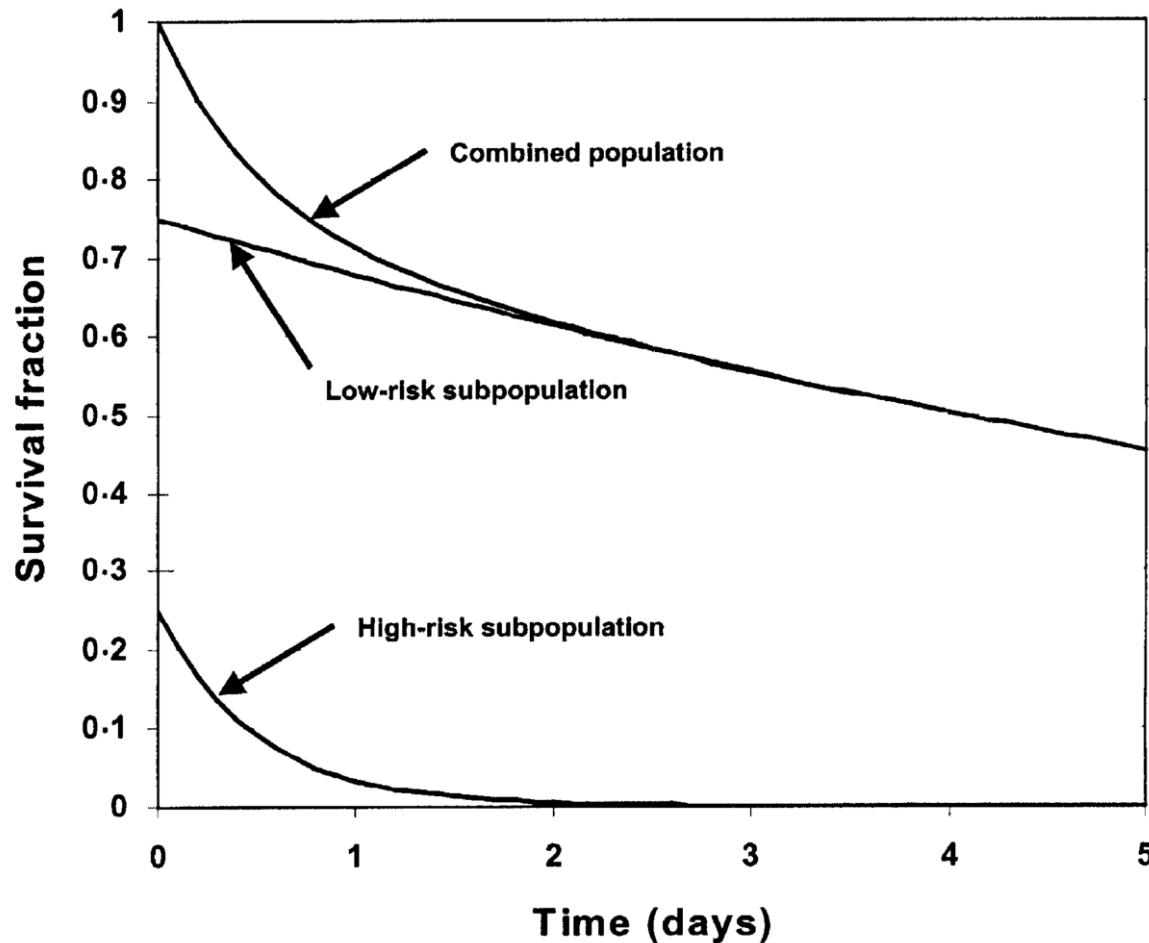
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Courbe de survie dans les STEMI

Mixed-exponential hazards modelling



Stratification du risque des patients STEMI traités par une angioplastie primaire

ZRS – Zwolle Risk Score

<u>Killip Class</u>	<u>Points</u>
1	0
2	4
3-4	9
<u>TIMI flow post</u>	
3	0
2	1
0-1	2
<u>Age</u>	
< 60	0
≥ 60	2
<u>3-vessel disease</u>	
No	0
Yes	1
<u>Anterior infarction</u>	
No	0
Yes	1
<u>Ischemia time (> 4 hours)</u>	
No	0
Yes	1
<u>Total score</u>	<u>16</u>

Front

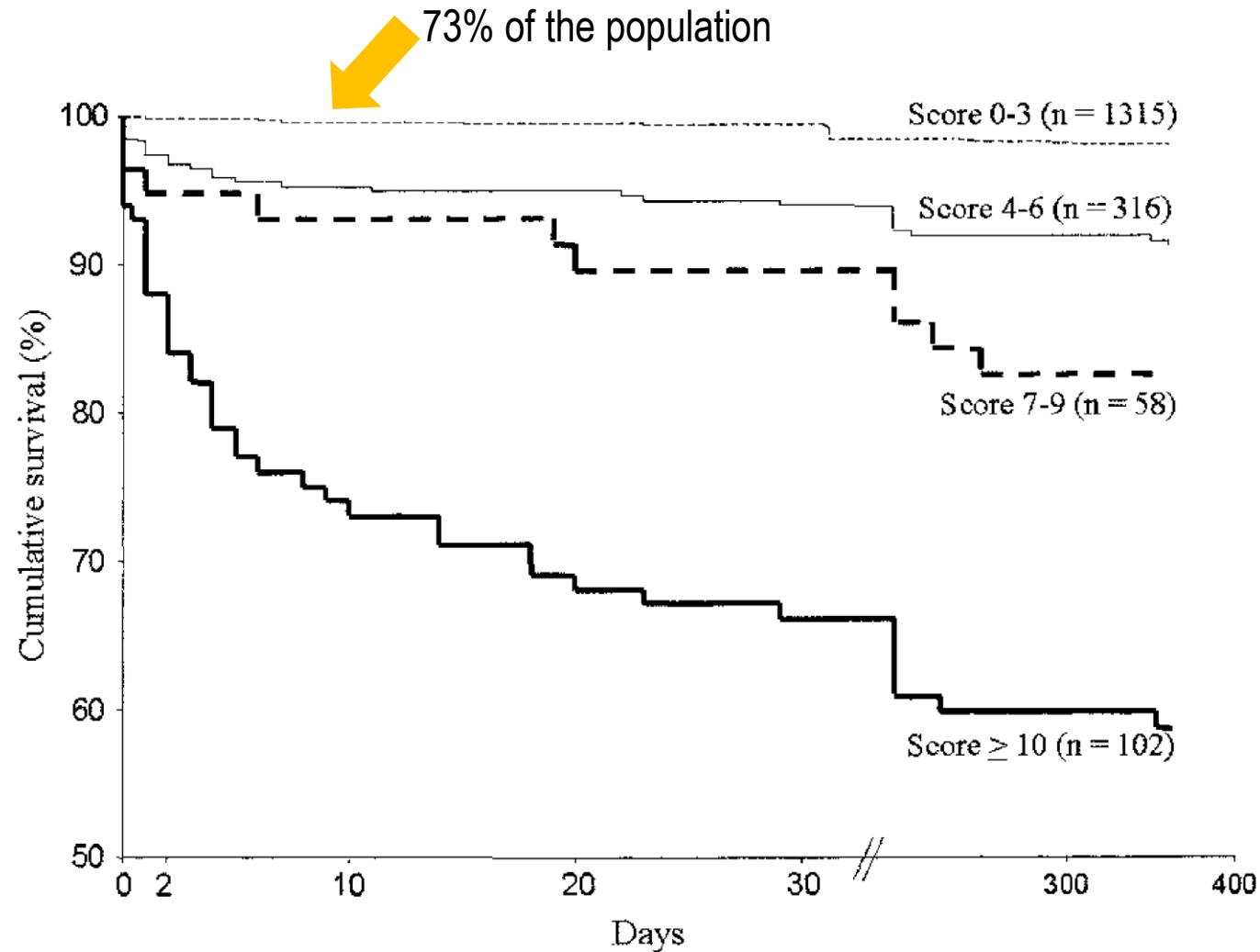
PAMI Low Risk

- age < 70 years,
- no persistent arrhythmias after reperfusion
- one- or two-vessel disease
- LVEF > 45%
- Successful PTCA

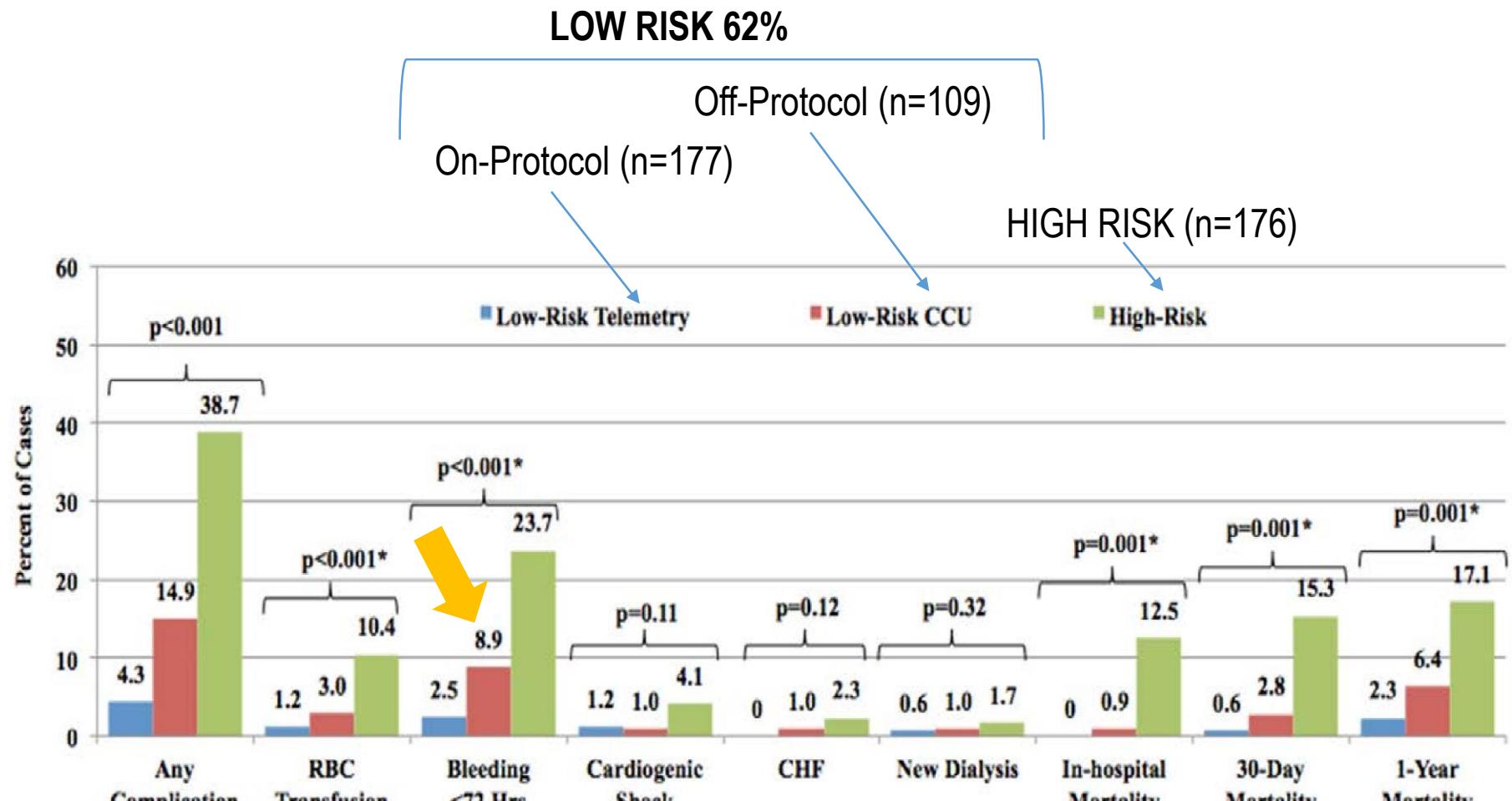
CADILLAC risk score

EF < 40%	4
Cr clearance < 60 mL/min	3
Killip class 2 or 3	3
Final TIMI flow 0-2	2
Age > 65 yrs	2
Anemia (hemoglobin < 13.0 mg/dL for males or < 12.0 mg/dL for females)	2
Three vessel disease	2

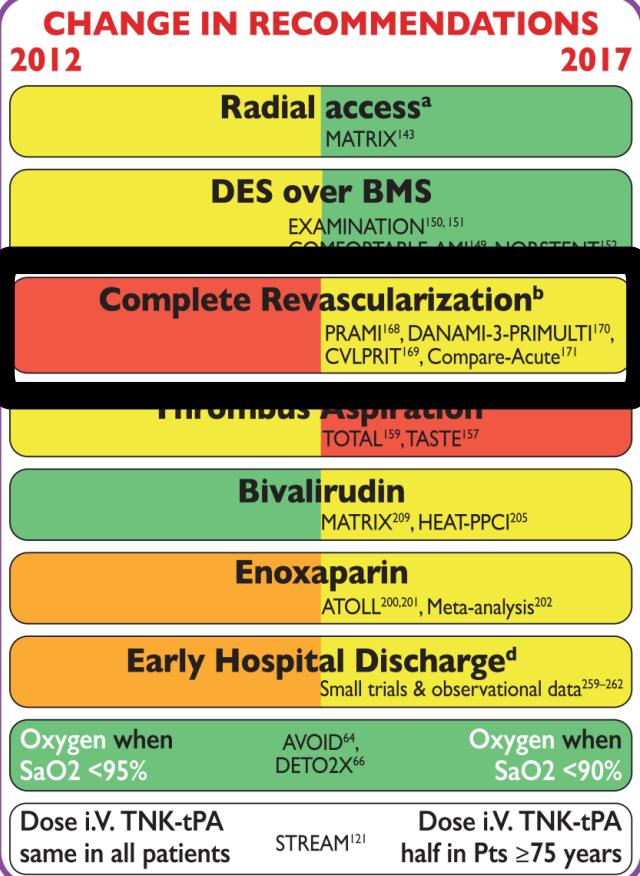
Zwolle risk score et mortalité



Plus besoin d'USIC pour les STEMI ?



Prospective cohort	Low-risk on-protocol (n=177)	Low-risk off-protocol (n=109)	High risk (n=176)
Patient costs	\$6090 (\$4730, \$7356)	\$8412 (\$6728, \$10920)	\$11 783 (\$7953, \$25 359)



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- Left and right bundle branch block considered equal for recommending urgent angiography if ischemic symptoms.

TIME TO ANGIOGRAPHY AFTER FIBRINOLYSIS:

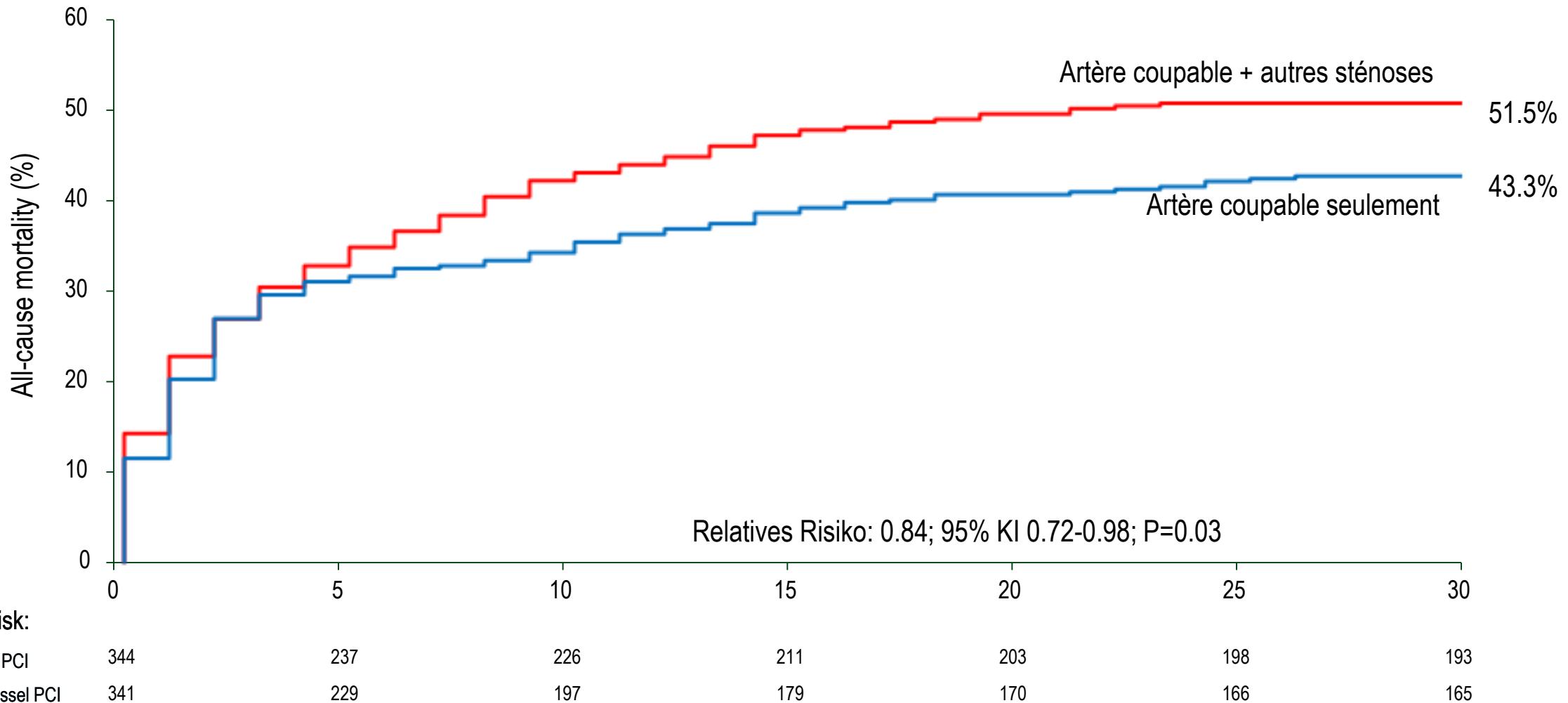
- Timeframe is set in 2–24h after successful fibrinolysis.

PATIENTS TAKING ANTICOAGULANTS:

- Acute and chronic management presented.



Mortalité totale



Thiele et al. NEJM 2017; 377:2419-2432

Merci de votre attention