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Le prétraitemen^t dans le syndrome coronaire aigu

MA BOUZID, Alger

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DÉCLARATION DE LIENS D'INTÉRÊT AVEC LA PRÉSENTATION

Nom de l'orateur : Mohammed El Amine BOUZID, Alger

Je n'ai pas de lien d'intérêt potentiel à déclarer

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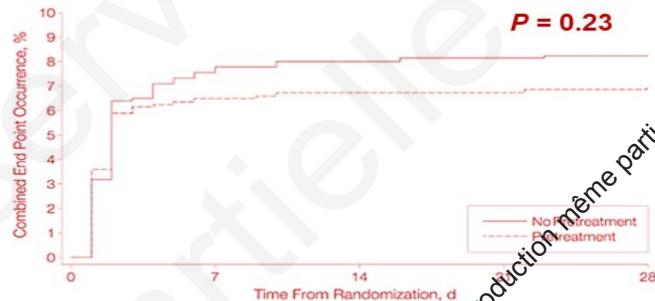
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Prétraitement

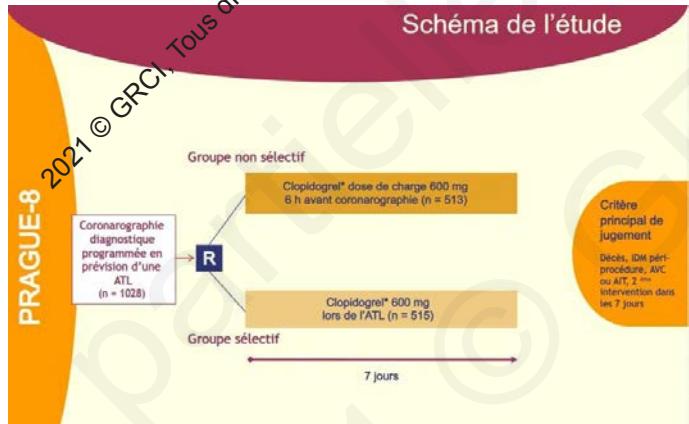
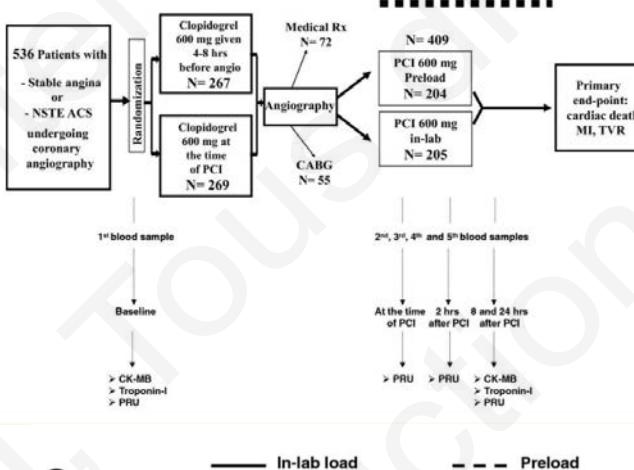
- Initiation des inhibiteurs des récepteurs P2Y₁₂, soit en pré-hospitalier, aux pavillons des urgences/USIC/ au cath-lab avant de connaître l'anatomie coronaire au décours d'un SCA
- La dose de charge d'un puissant inhibiteur du récepteur P2Y₁₂, avant la coronarographie, réduira le risque de complications thrombotiques lors d'une PCI. Ceci est d'autant plus important chez les patients souffrant d'un SCA, chez qui les plaquettes sont déjà fort activées (plusieurs études CURE, PCA-CURE, CLARITY ...)

ARMYDA-5 Preload

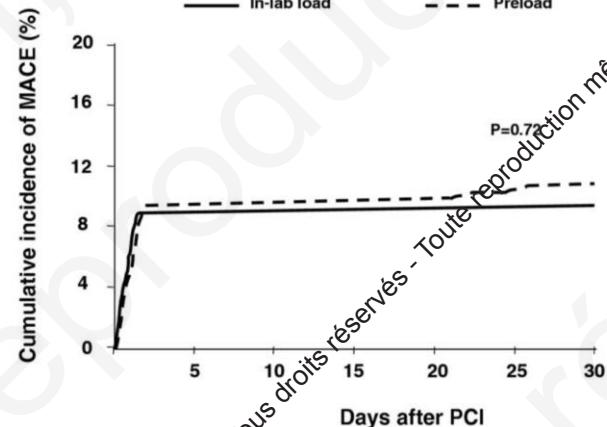
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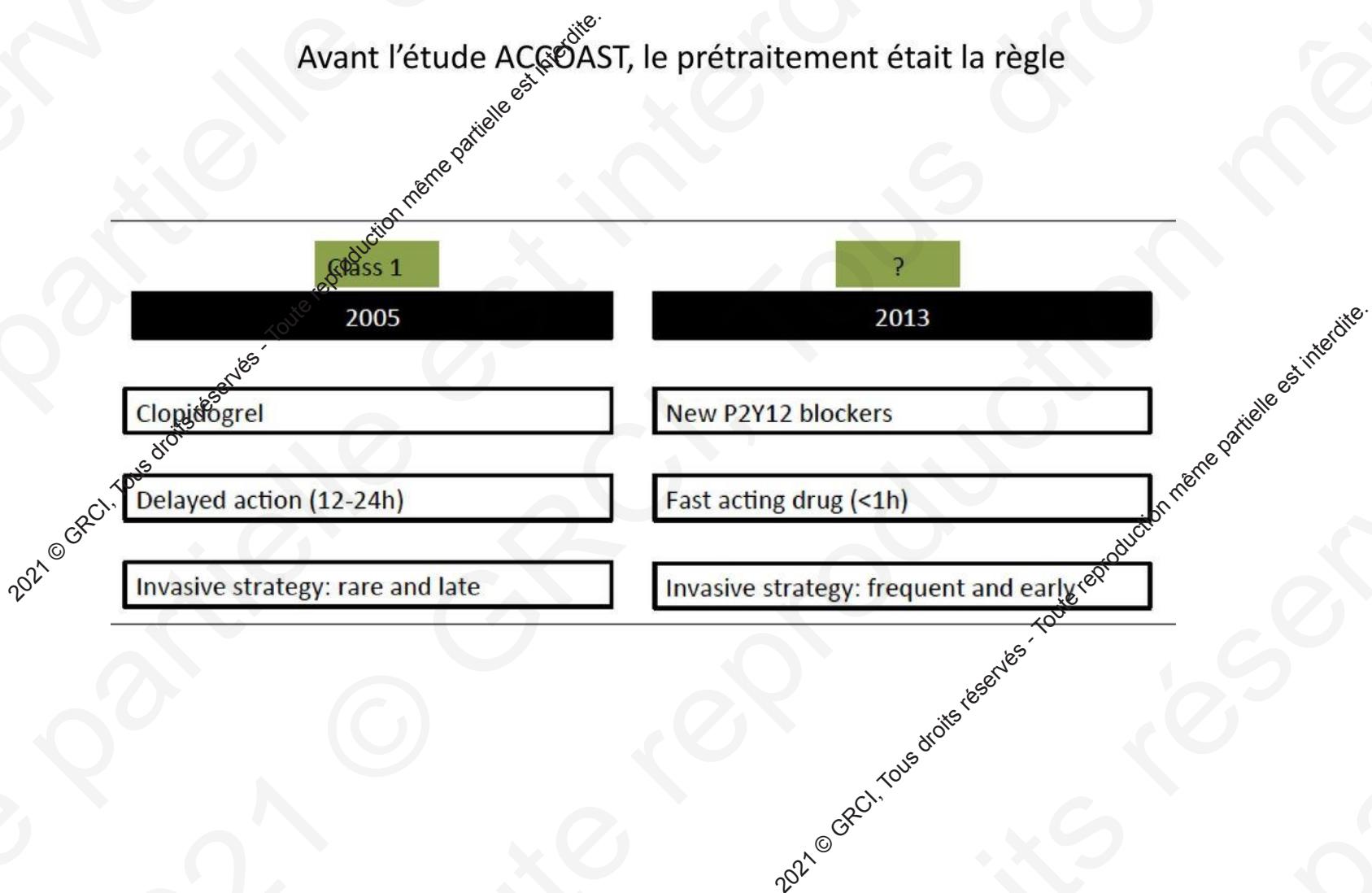
Steinhubl S et al. JAMA. 2002;288(19):2411-2420



Widimski P et al. 2008;29:1495-503



Di Sciascio G et al. JAAC 2010;10:56:550-7

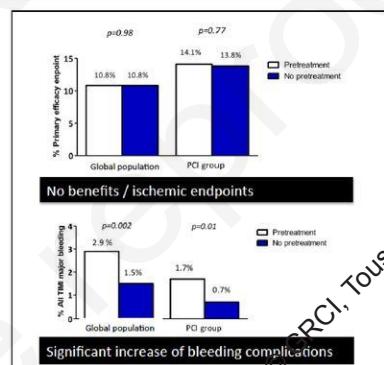
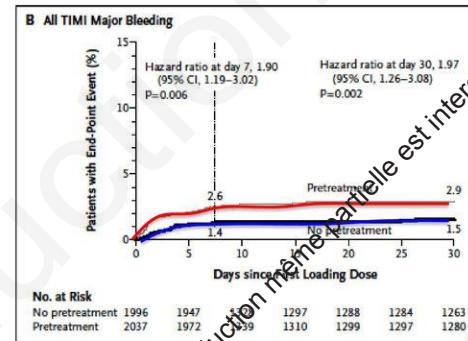
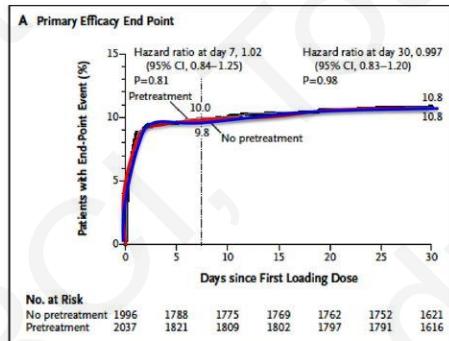
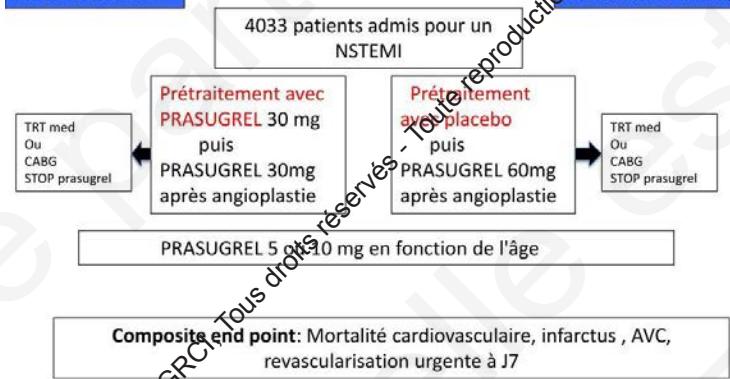


ACCOAST (Prasugrel)

Pretreatment with Prasugrel in Non-ST-Segment Elevation Acute Coronary Syndrome

ACCOAST STUDY

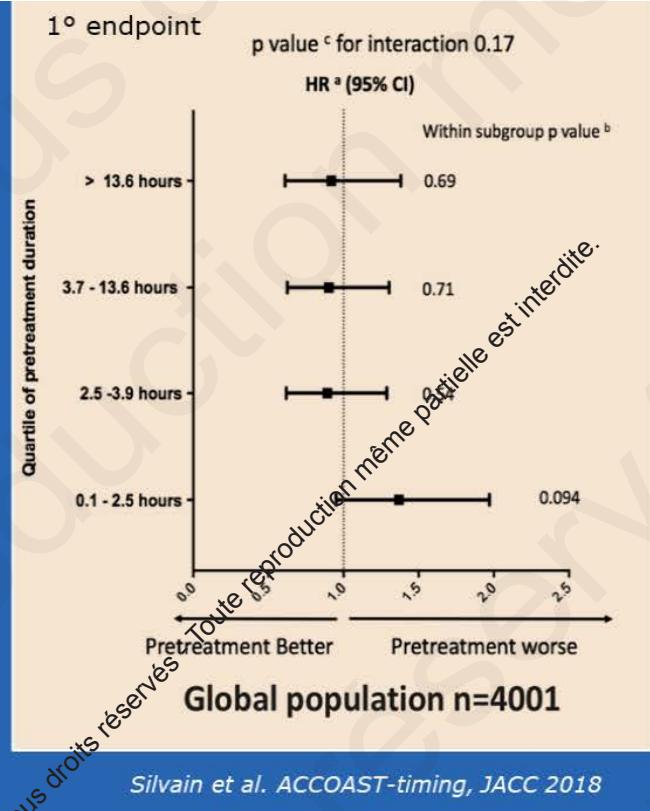
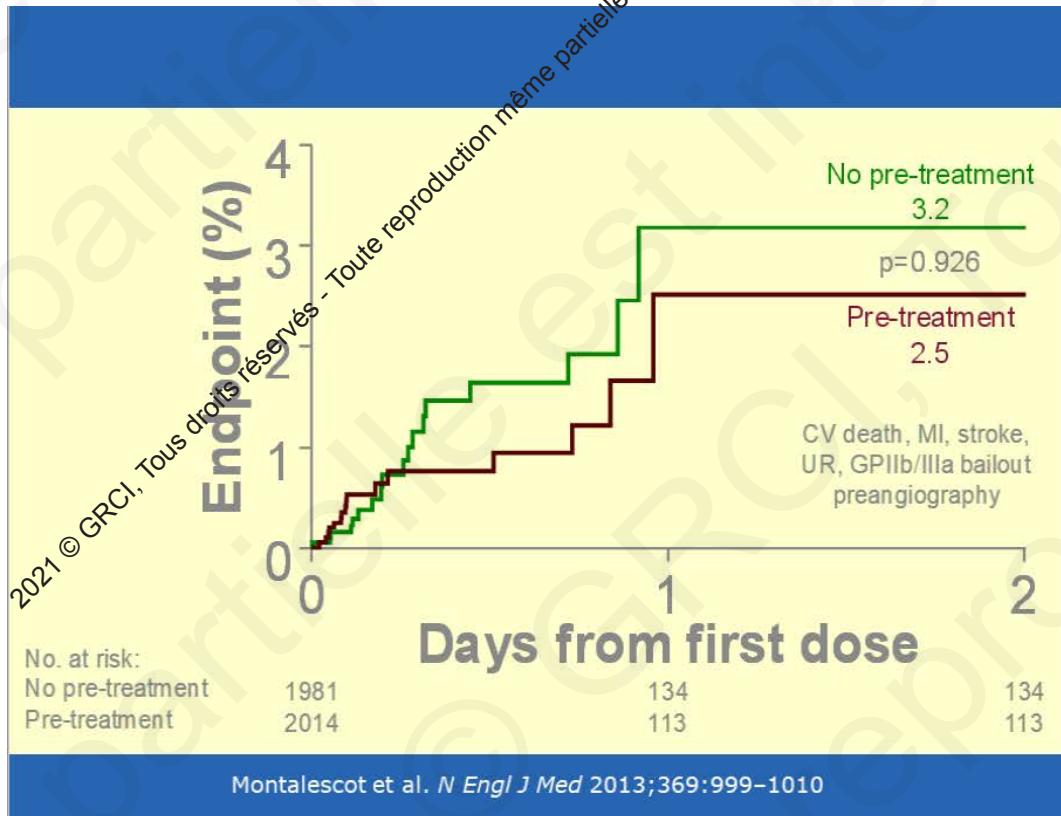
Montalescot and al, NEJM 2013



CONCLUSION ACCOAST

- Pas de bénéfice du pré-traitement par PRASUGREL
- PAS DE bénéfice dans le groupe PCI
- Augmentation des complications hémorragiques

ACCOAST (Prasugrel)



Pas de sur-risque en attente de la coronarographie

DUBIUS (Ticagrelor)



Clinical Topics Latest in Cardiology Education and Meetings Tools and Practice Support

Downstream or Upstream P2Y12 Receptor Blockers in NSTE-ACS: Primary Results of the DUBIUS Trial

Mar 15, 2021 | Giuseppe Tarantini; Marco Mojoli, MD

Expert Analysis

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Quick Takes

- The available evidence related to the clinical impact of pretreatment has been incomplete and heterogeneous regarding the different P2Y12 inhibitors.
- In the DUBIUS (Downstream Versus Upstream Strategy for the Administration of P2Y12 Receptor Blockers in Non-ST Elevated Acute Coronary Syndromes With Initial Invasive Indication) trial, downstream and upstream oral P2Y12 inhibitor administration strategies were associated with low incidence of ischemic and bleeding events and minimal numeric difference of event rates between treatment groups.
- Broad use of a radial approach may have contributed to the low observed adverse event rates.

Related Content

Downstream versus Upstream administration of P2Y12 receptor Blockers in non-ST elevated acute coronary Syndromes with initial invasive Indication

The Clopidogrel in Unstable Angina to Prevent Recurrent Events Trial: Investigators' Effects of Clopidogrel in Addition to Aspirin in Patients With Acute Coronary Syndromes Without ST-Segment Elevation

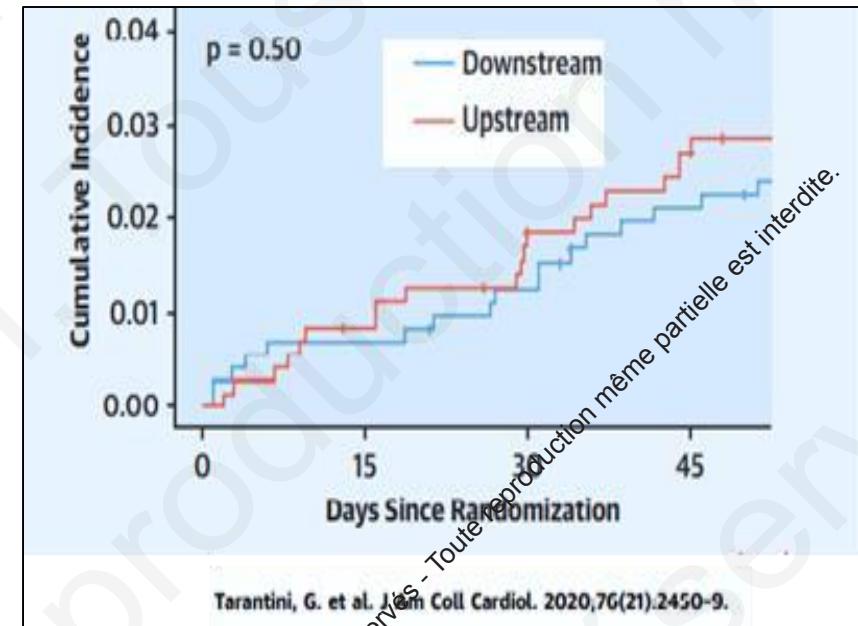
Clopidogrel for Reduction of Events During Observation

PLATelet inhibition and patient Outcomes

Trial to Assess Improvement in Therapeutic Outcomes by Optimizing Platelet Inhibition With Prasugrel-Thrombolysis in Myocardial Infarction 38

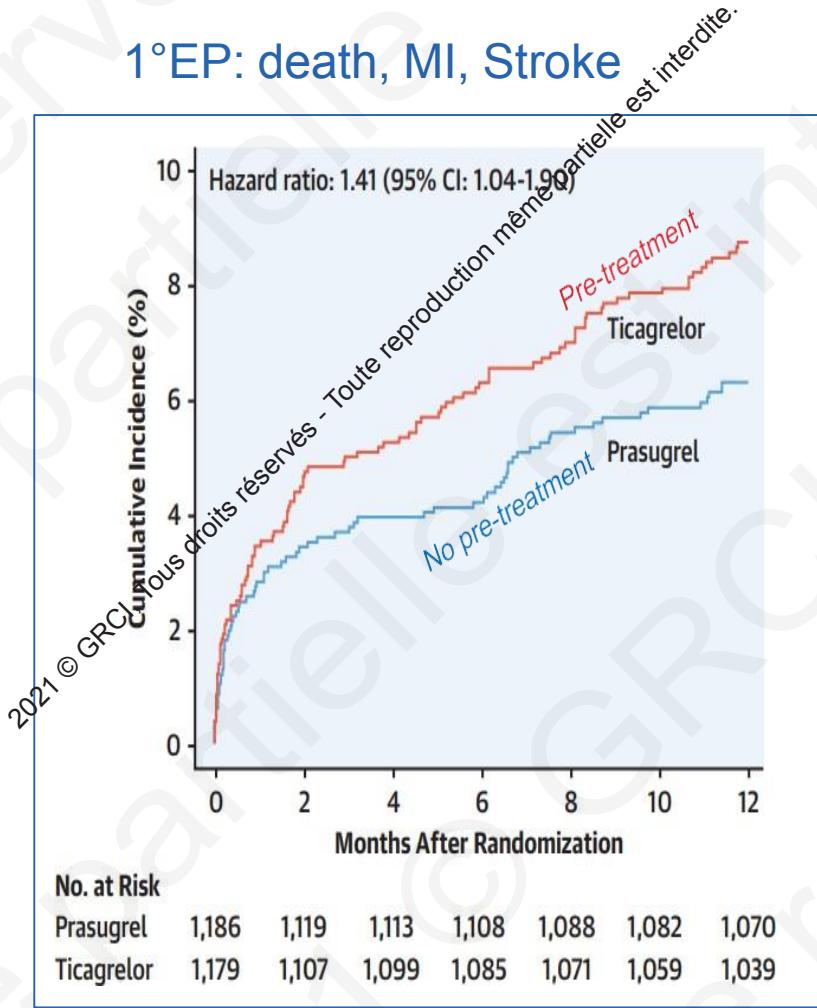
Comparison of Prasugrel at the Time of PCI or as Pretreatment at the Time of Diagnosis in Patients With Non-ST Elevation Myocardial Infarction

Intracoronary Stenting and Antithrombotic Regimen 5

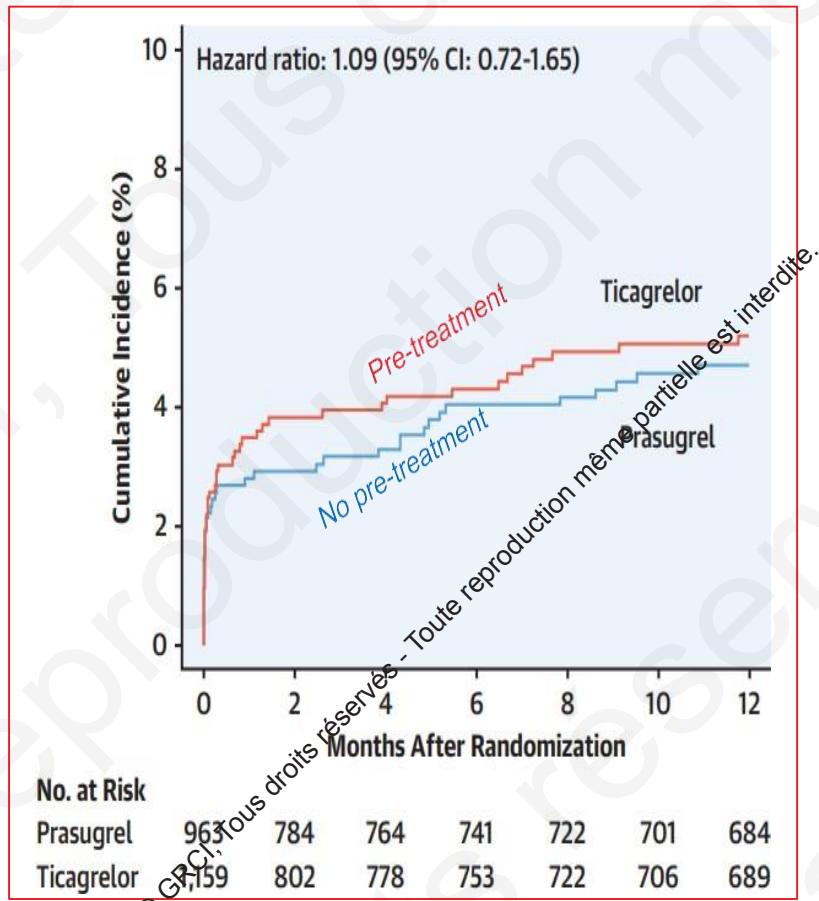


ISAR REACT 5 - NSTE-ACS

1°EP: death, MI, Stroke



Safety EP: BARC 3-5



Valina C et al. J Am Coll Cardiol 2020;76:2436-46

2020 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation

The Task Force for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation of the European Society of Cardiology (ESC)

Authors/Task Force Members: Jean-Philippe Collet  * (Chairperson) (France), Holger Thiele  * (Chairperson) (Germany), Emanuele Barbato (Italy), Olivier Barthélémy (France), Johann Bauersachs (Germany), Deepak L. Bhatt (United States of America), Paul Dendale (Belgium), Maria Dorobantu (Romania), Thor Edvardsen (Norway), Thierry Folliguet (France), Chris P. Gale (United Kingdom), Martine Géard (France), Alexander Jobs (Germany), Peter Jüni (Canada), Ekaterini Lambrinou (Cyprus), Basil S. Lewis (Israel), Julinda Mehilli (Germany), Emanuele Meliga (Italy), Béla Merkely (Hungary), Christian Mueller (Switzerland), Marco Roffi (Switzerland), Frans H. Rutten (Netherlands), Dirk Sibbing (Germany), George C.M. Siontis (Switzerland)

Downloaded from https://academic.oup.com/europace/article/42/14/1289/55

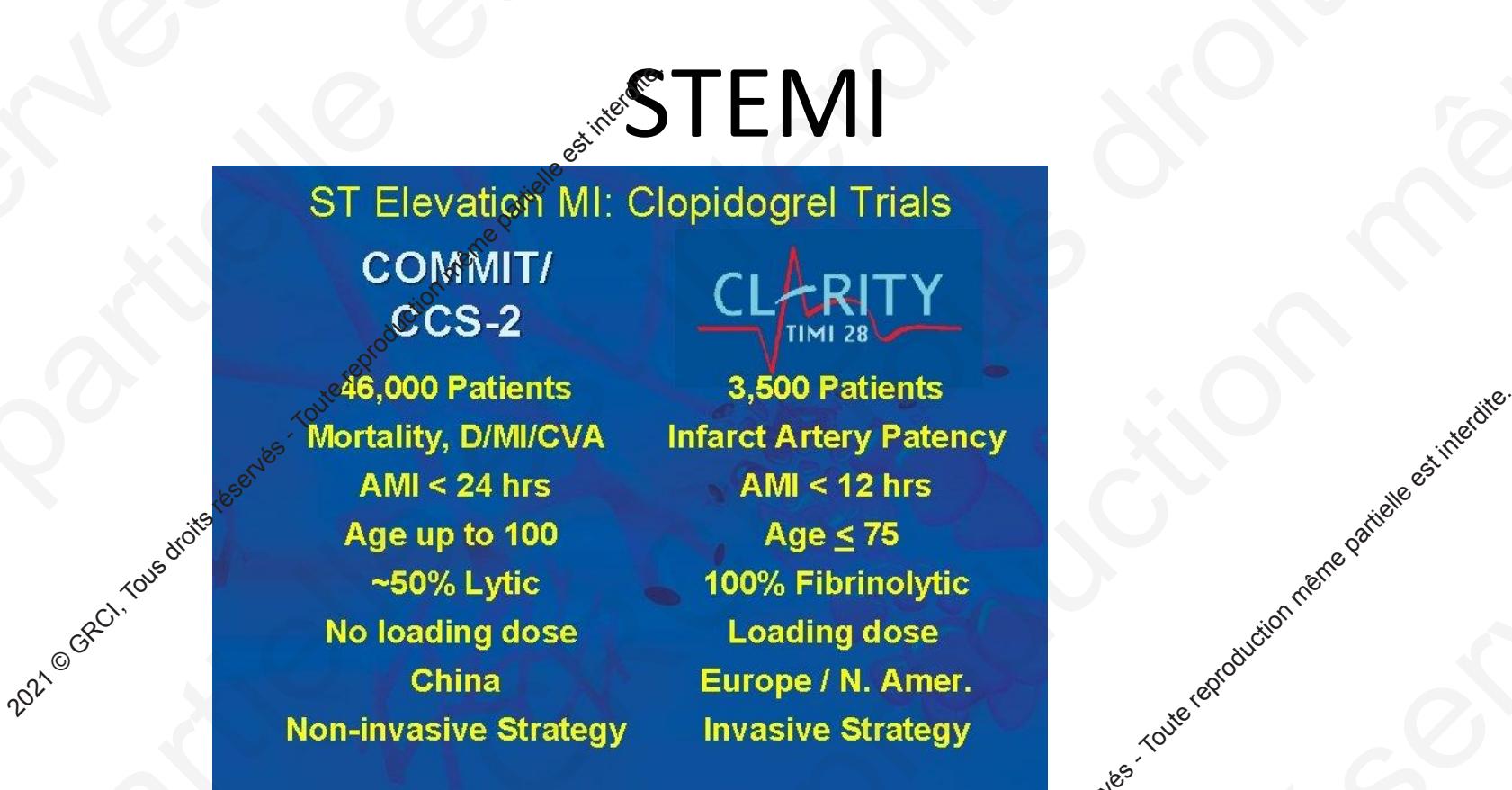
Treatment with GP IIb/IIIa antagonists in patients in whom coronary anatomy is not known is not recommended 

It is not recommended to administer routine pre-treatment with a P2Y₁₂ receptor inhibitor in patients in whom coronary anatomy is not known and an early invasive management is planned. 

III	A
III	A

Messages clés (NSTEMI)

- Il n'est pas recommandé de prétraiter de routine par un inhibiteur du récepteur P2Y12 chez les patients dont l'anatomie coronaire n'est pas connue et pour lesquels une prise en charge invasive précoce est prévue (classe III A)
- Patients NSTEMI qui ne peuvent pas bénéficier d'une stratégie invasive précoce, un préTRT peut être envisagé chez les patients à faible risque hémorragique (classe II b) >>> **prise en charge individualisée**
- Le Prasugrel doit être préféré au Ticagrelor chez les patients atteints de NSTE-ACS qui vont bénéficier d'angioplastie (classe IIa b)



Ces 2 études ont validé (bien qu'imparfaitement) l'utilité d'un traitement d'emblée par aspirine et clopidogrel dans le STEMI, qu'il y ait ou non angioplastie primaire

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STE MI

ATLANTIC

Administration of Ticagrelor in the cath Lab or in the Ambulance for New ST elevation myocardial Infarction to open the Coronary artery

Atlantic Population

- Documented evidence of STEMI
- Planned for angioplasty (PCI)
- onset of ischaemic symptoms within 6 h
- initially managed by ambulance physician/personnel; also concerning patients not pre-treated for STEMI in emergency rooms of non-PCI hospitals

STE-ACS planned for PCI (N = 1862)

```

graph TD
    Start(( )) --> Randomised[Randomised, double-blind]
    Randomised --> PreHospital[Pre-hospital]
    Randomised --> InHospital[In-Hospital]
    
    subgraph PreHospital [Pre-hospital]
        direction TB
        A1[Ticagrelor  
180 mg loading dose] --> B1[Placebo  
loading dose]
        A2[Placebo  
loading dose] --> B2[Ticagrelor  
180 mg loading dose]
    end
    
    subgraph InHospital [In-Hospital]
        direction TB
        B1 --> C1[Primary Objectives]
        B2 --> C1
    end
    
    C1[≥ 70% ST-segment elevation resolution pre-PCI  
OR  
TIMI flow grade 3 of MI culprit vessel at initial angiography] --> D1[Ticagrelor 90 mg/bid 30 days]
    C1 --> D2[Placebo 90 mg/bid 30 days]

```

End Point	Interventional		Control		P Value†	Difference (95% CI)‡
	In-hospital Transfer (N = 906) as fast as possible after arrival	Out-hospital Transfer (N = 952)	Odds Ratio (95% CI)§	P Value†		
Coprimary end points						
Absence of ST-segment elevation resolution ≥ 70% before PCI	672/774 (86.8)	722/824 (88.0)	0.93 (0.69 to 1.25)	0.63	-0.008 (-0.041 to 0.025)	
Absence of TIMI flow grade 3 in infarct-related artery at initial angiography	681/824 (82.6)	717/777 (83.1)	0.97 (0.75 to 1.25)	0.82	-0.004 (-0.040 to 0.032)	
Both						
Both	541/744 (72.1)	577/777 (73.3)	0.96 (0.77 to 1.21)	0.73	-0.008 (-0.052 to 0.037)	
One or both	677/779 (87.4)	710/751 (94.5)	0.93 (0.60 to 1.45)	0.75	-0.004 (-0.027 to 0.032)	
Secondary end points						
Absence of ST-segment elevation resolution ≥ 70% after PCI	517/742 (69.1)	535/743 (71.5)	0.82 (0.66 to 1.04)	0.05	-0.050 (-0.101 to 0.001)	
Absence of TIMI flow grade 3 in infarct-related artery after PCI	135/760 (17.8)	154/784 (19.6)	0.88 (0.68 to 1.14)	0.34	-0.019 (-0.038 to 0.020)	
Met one or both secondary end points						
Both	73/763 (9.6)	87/775 (11.2)	0.84 (0.60 to 1.16)	0.29	-0.017 (-0.047 to 0.064)	
One or both	339/644 (49.4)	371/703 (52.8)	0.88 (0.71 to 1.09)	0.23	-0.032 (-0.083 to 0.020)	

Recommendations	Class ^b	Level ^c
Antiplatelet therapy		
A potent P2Y ₁₂ inhibitor (prasugrel or ticagrelor), or clopidogrel if these are not available or are contraindicated, is recommended before (or at latest at the time of) PCI and maintained over 12 months, unless there are contraindications such as excessive risk of bleeding. ^{186,187}		A
Aspirin (oral or i.v. if unable to swallow) is recommended as soon as possible for all patients without contraindications. ^{213,214}	I	B
GP IIb/IIIa inhibitors should be considered for bailout if there is evidence of no-reflow or a thrombotic complication.	IIa	C
Cangeflor may be considered in patients who have not received P2Y ₁₂ receptor inhibitors. ^{192–194}	IIb	A



European Heart Journal (2017) 0, 1–48
doi:10.1093/eurheartj/ehx419

ESC GUIDELINES

2017 ESC focused update on dual antiplatelet therapy in coronary artery disease developed in collaboration with EACTS

The Task Force for dual antiplatelet therapy in coronary artery disease of the European Society of Cardiology (ESC) and of the European Association for Cardio-Thoracic Surgery (EACTS)

Seulement en bail-out, plus en systématique en cas de forte charge thrombotique

A Considérer

Conclusion

- Aspirine (dose de charge) doit être administrée dès le diagnostic de SCA (NSTEMI et STEMI)
- **NSTEMI : pas de préTRT par des inhibiteurs P2Y12 avant de connaître les lésion coronaires en cas de stratégie invasive précoce**
- Patients **NSTEMI** qui ne peuvent pas bénéficier d'une stratégie invasive précoce, **un préTRT peut être envisagé chez les patients à faible risque hémorragique (classe IIb)>>> prise en charge individualisée**
- **STEMI:** dose de charge de Prasugrel ou Ticagrelor/Clopidogrel dès le diagnostic