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Congreso Sociedad Andaluza de Cardiología “Congreso Andaluz de las Enfermedades Cardiovasculares”

14 – 16 mayo 2015

Hotel Abades Nevada Palace - Granada



ESQUEMA DE TRATAMIENTO ACTUAL MEDIANTE REVASCULARIZACION PRECOZ EN EL INFARTO AGUDO DE MIOCARDIO

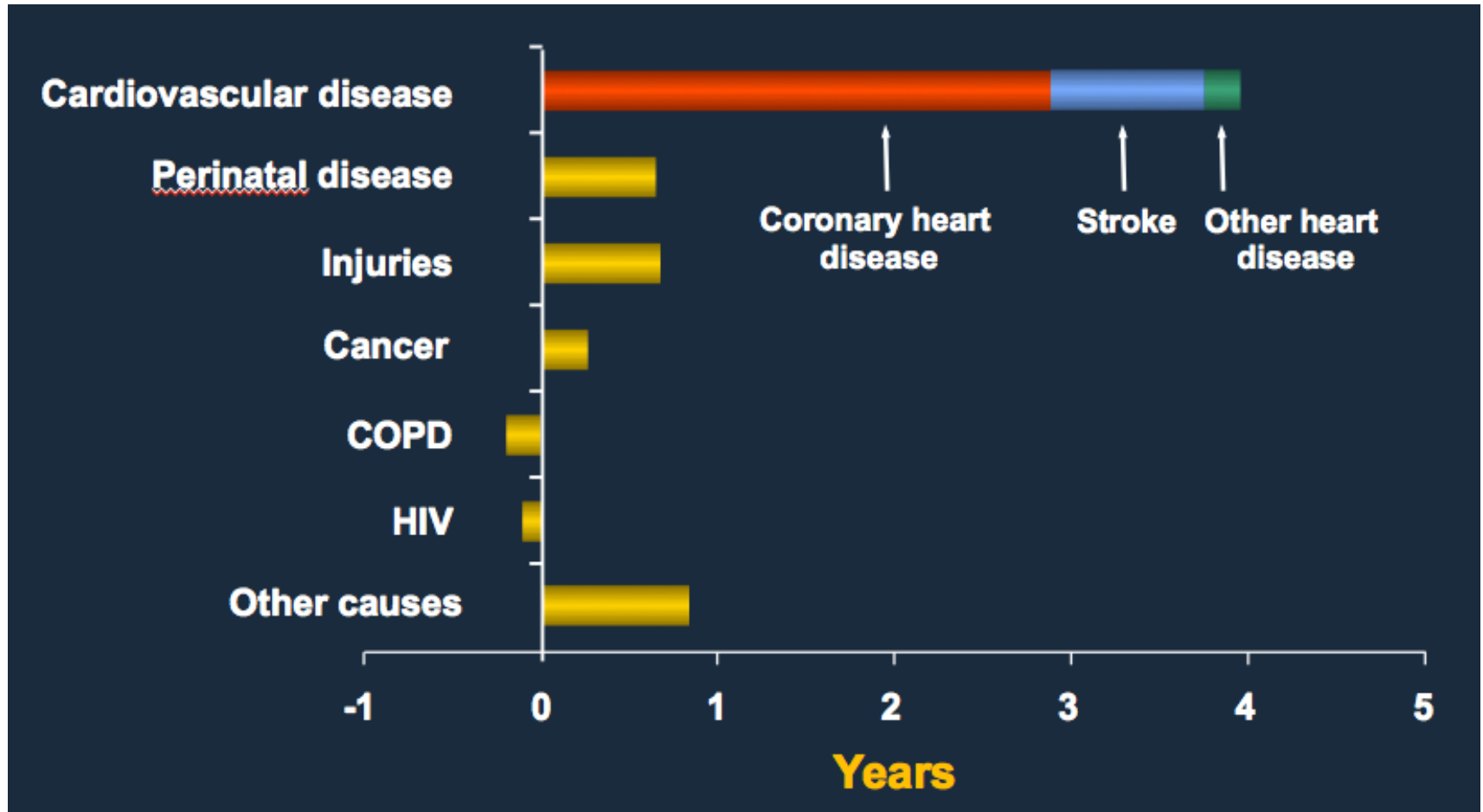
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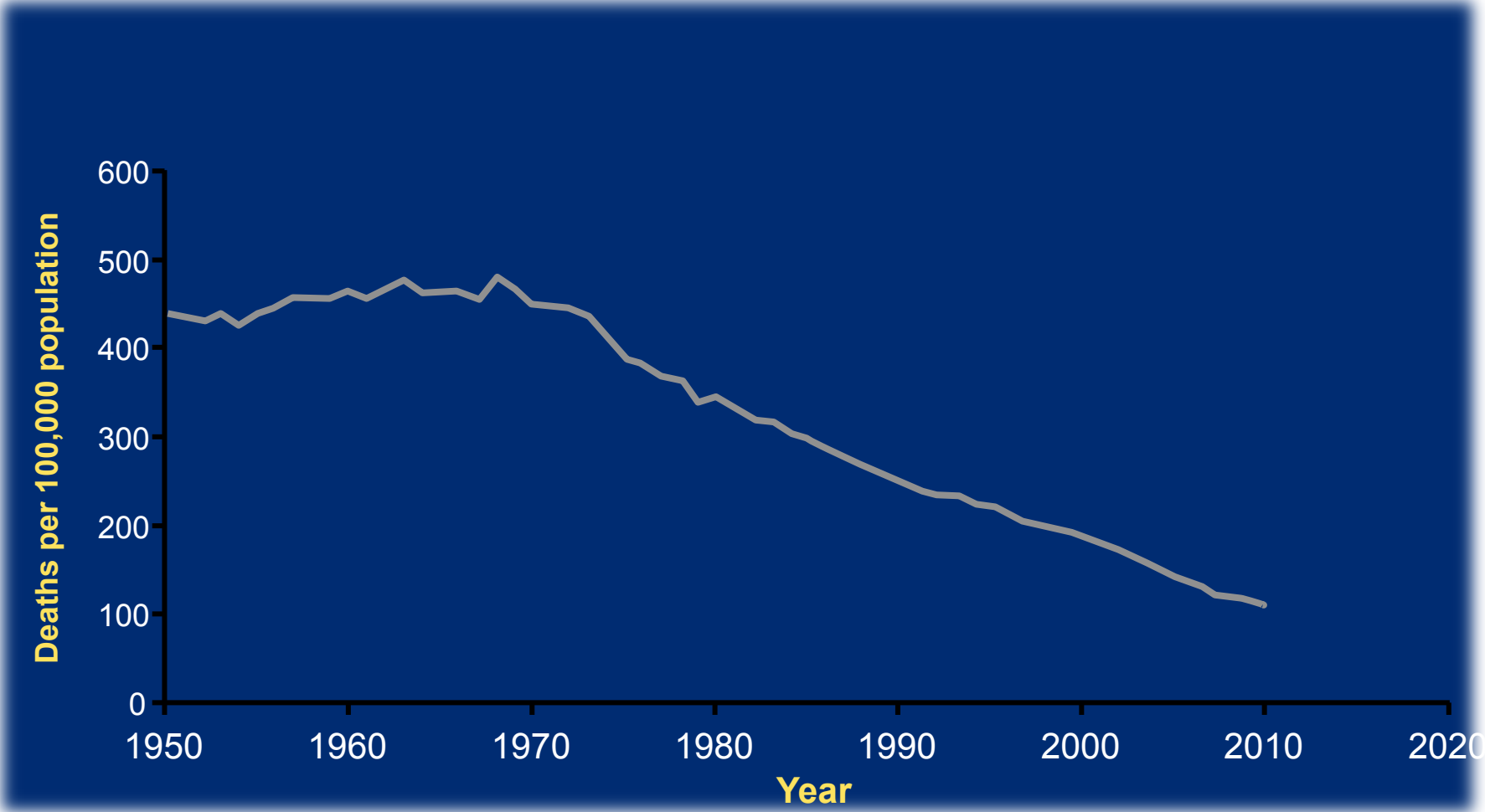


Change in US Life Expectancy Between 1970 and 2000: 8 years↑



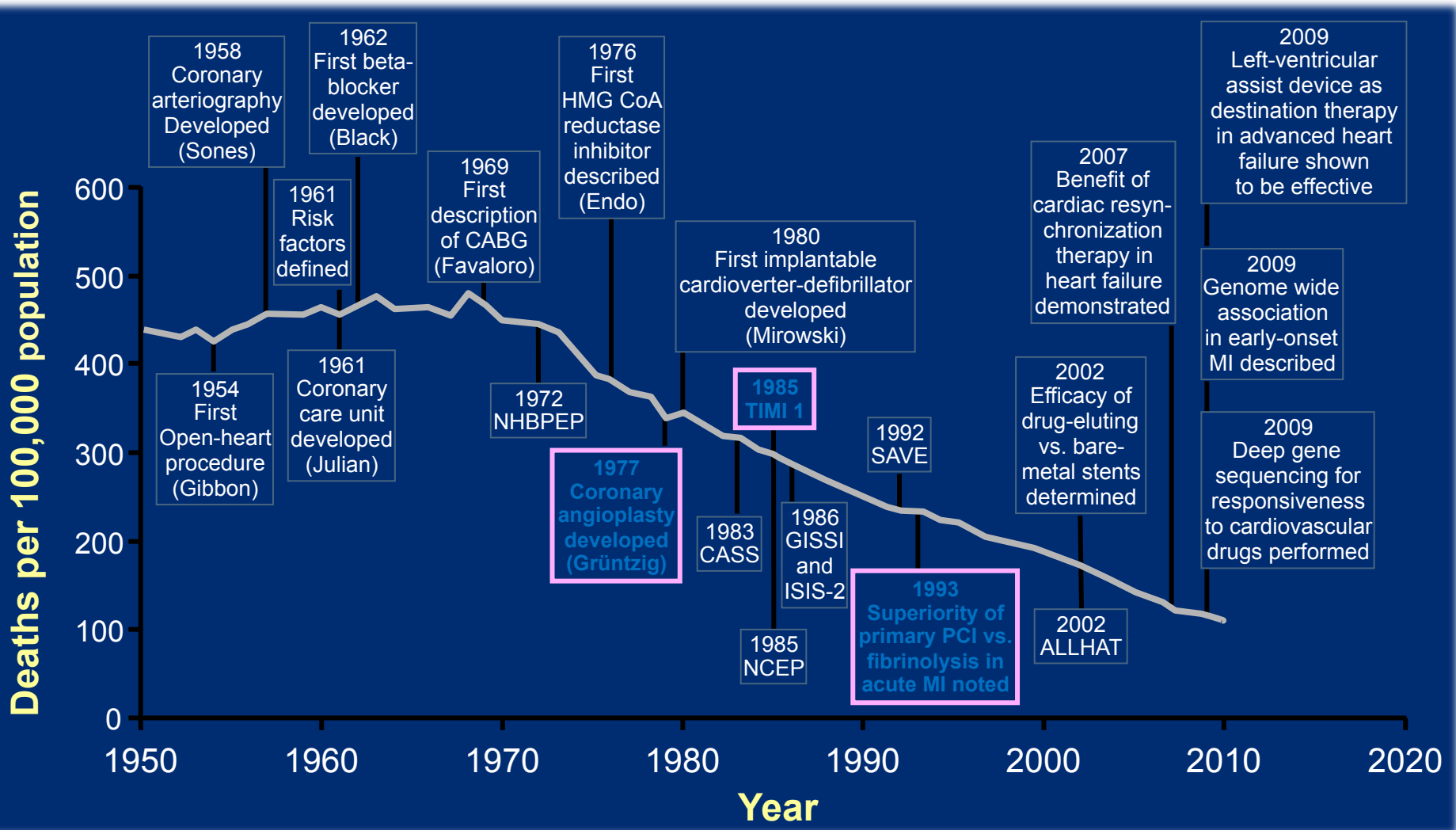
C. Lenfant et al. *NEJM* 2003;349:868–874

Decline in Deaths from Cardiovascular Disease in Relation to Scientific Advances



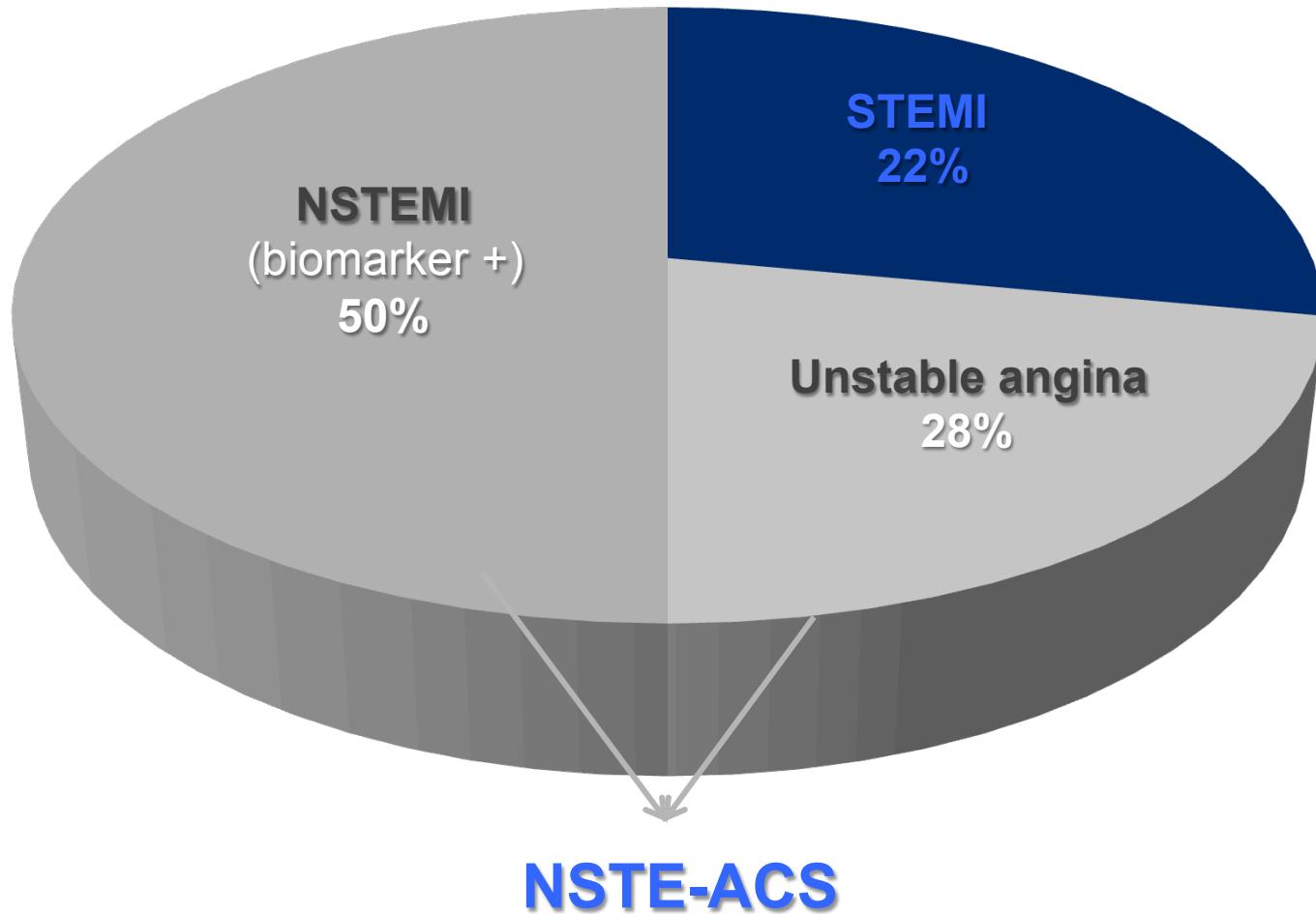
Nabel EG and Braunwald E. 2012;366:54-63

Decline in Deaths from Cardiovascular Disease in Relation to Scientific Advances



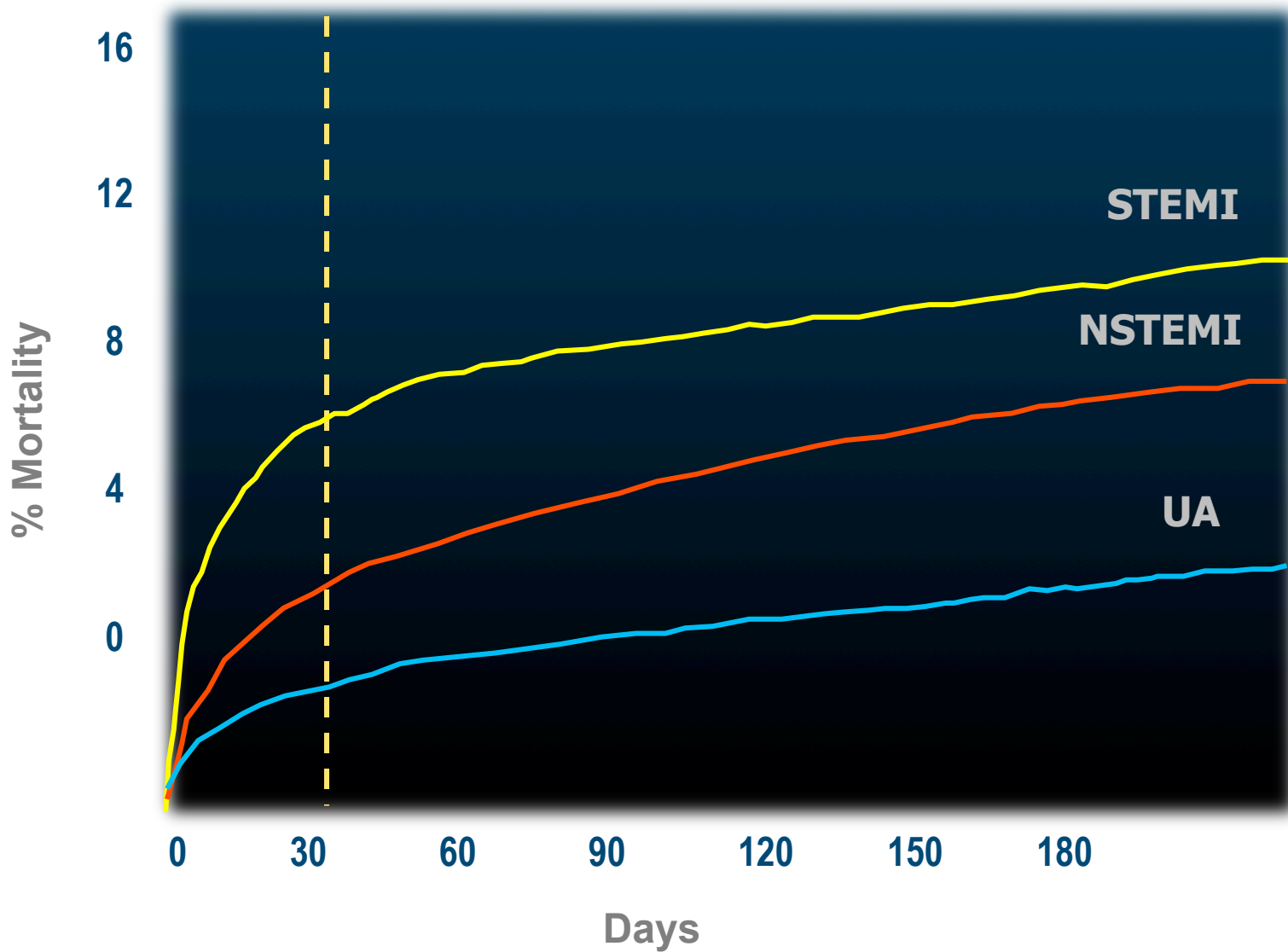
Acute Coronary Syndromes (ACS)

1.41 million hospital admissions for ACS in 2010

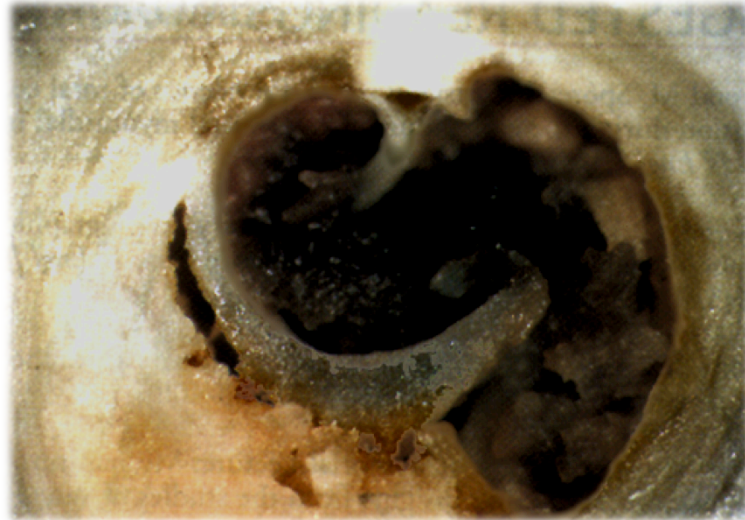
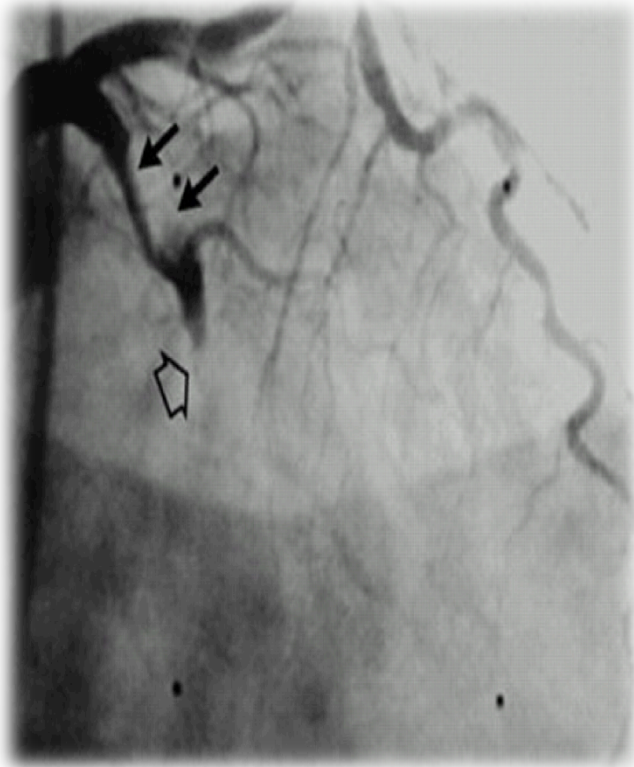


Mortality in Acute Coronary Syndromes

Death from admission to 6 mos (GRACE n=43,810)



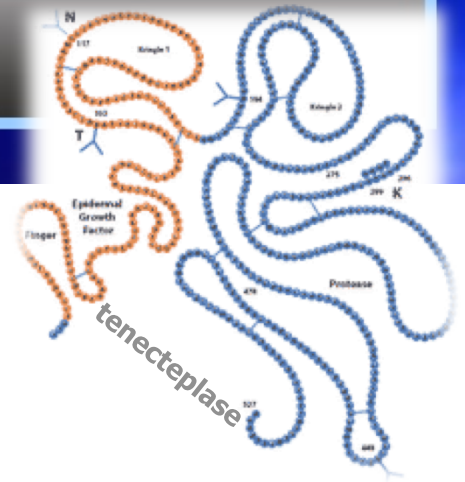
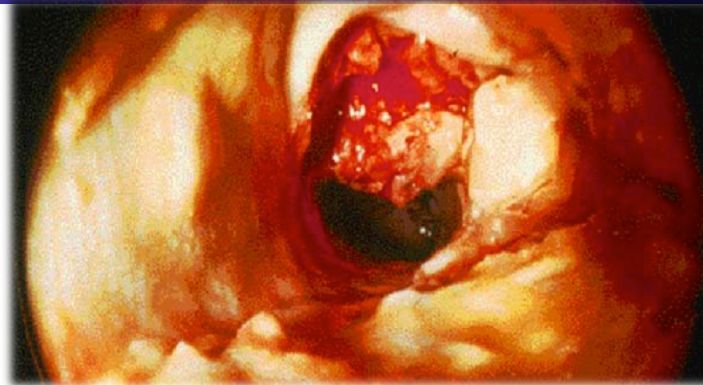
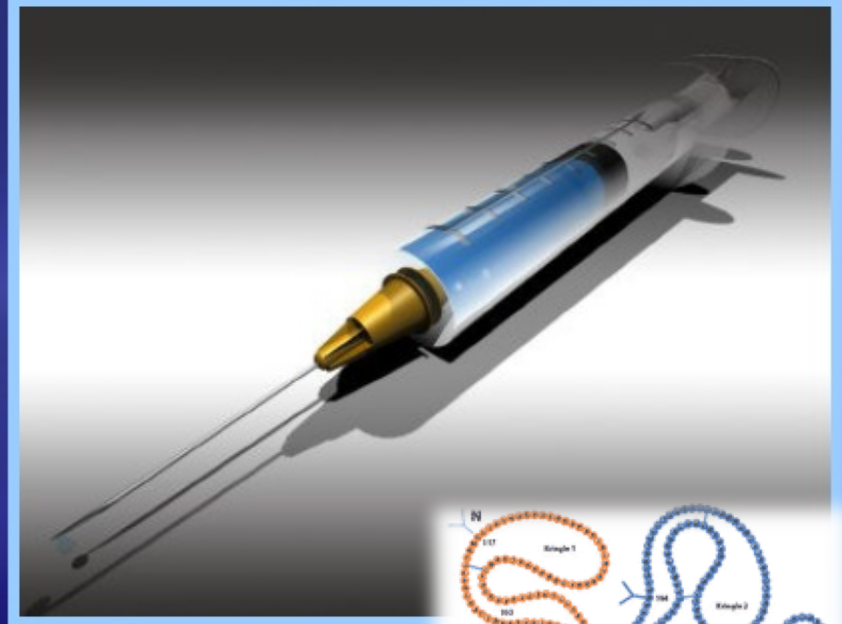
AMI: pathophysiology



ruptured plaque with occlusive thrombus

Angioplastia Primaria

Fibrinolisis



Fibrinolytic therapy

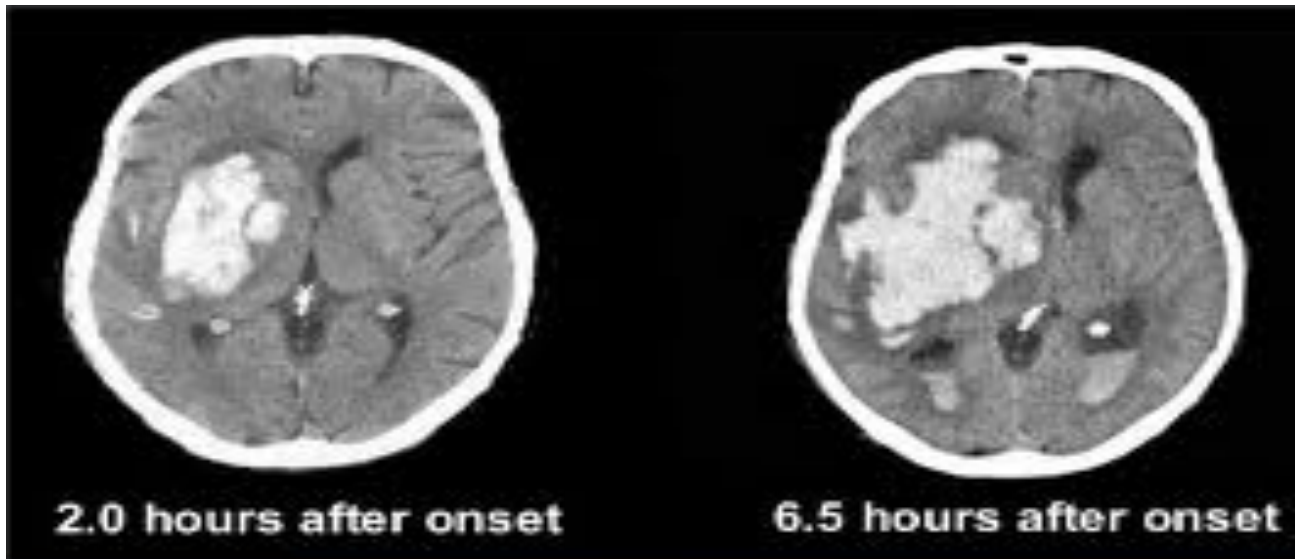
... did save lives compared to placebo, BUT

- at best, restored TIMI 3 flow in 55% (rt-PA)
- incidence of recurrent ischemia and reinfarction

+

ICH

0.5-1.0% of pts



2.0 hours after onset

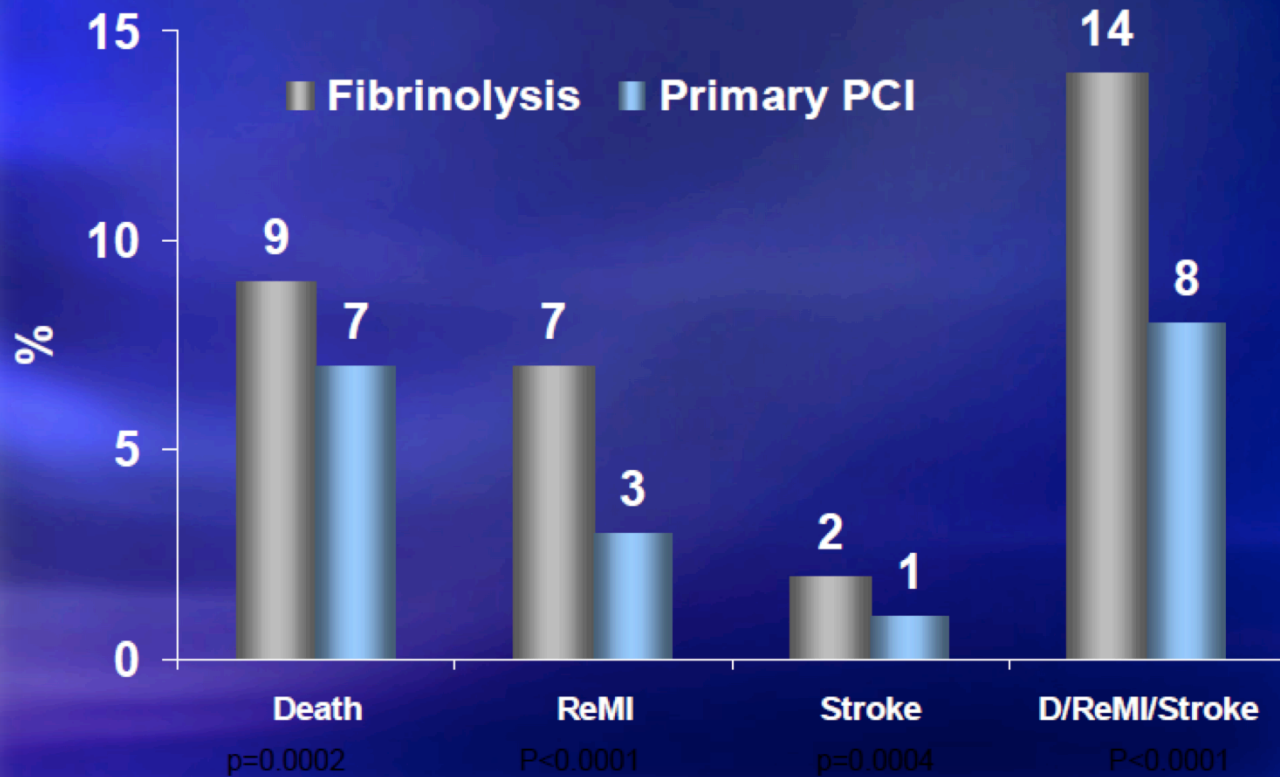
6.5 hours after onset

2 hours
after t-PA

6 hours
after t-PA

Primary PCI vs. Fibrinolysis

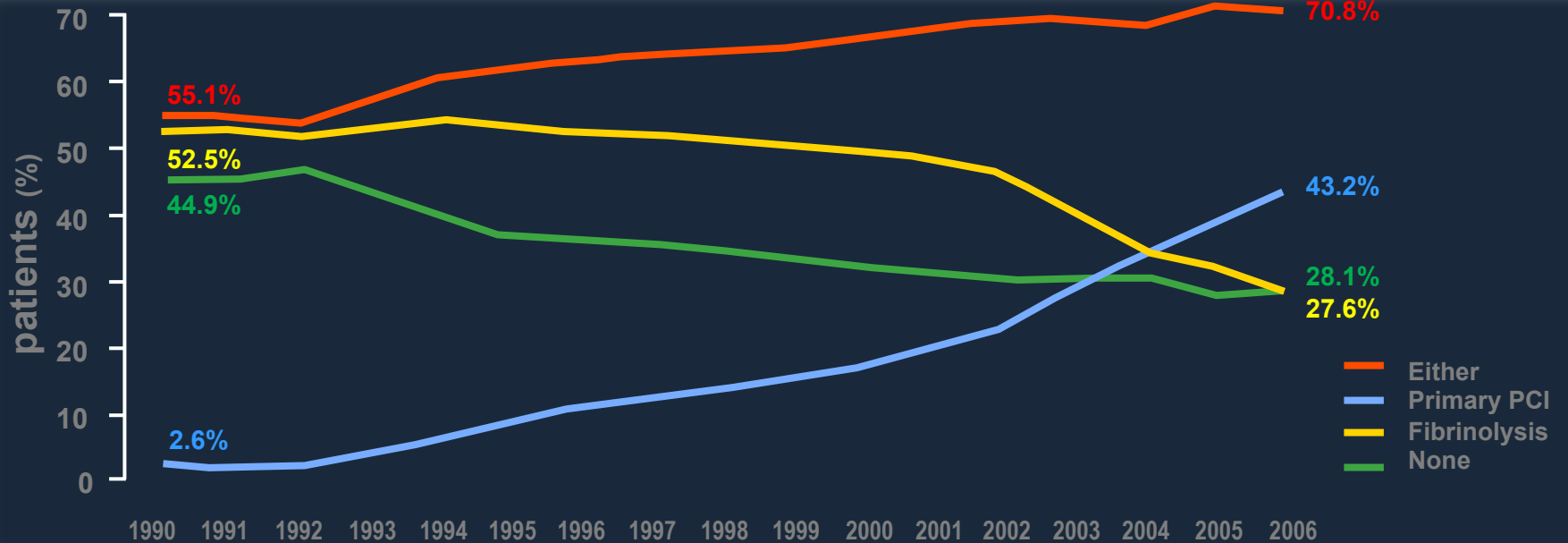
Meta-Analysis of 23 trials (n=7739)



Evolution in Reperfusion

774,279 reperfusion eligible STEMI pts at 2,157 hospitals from 1990-2006

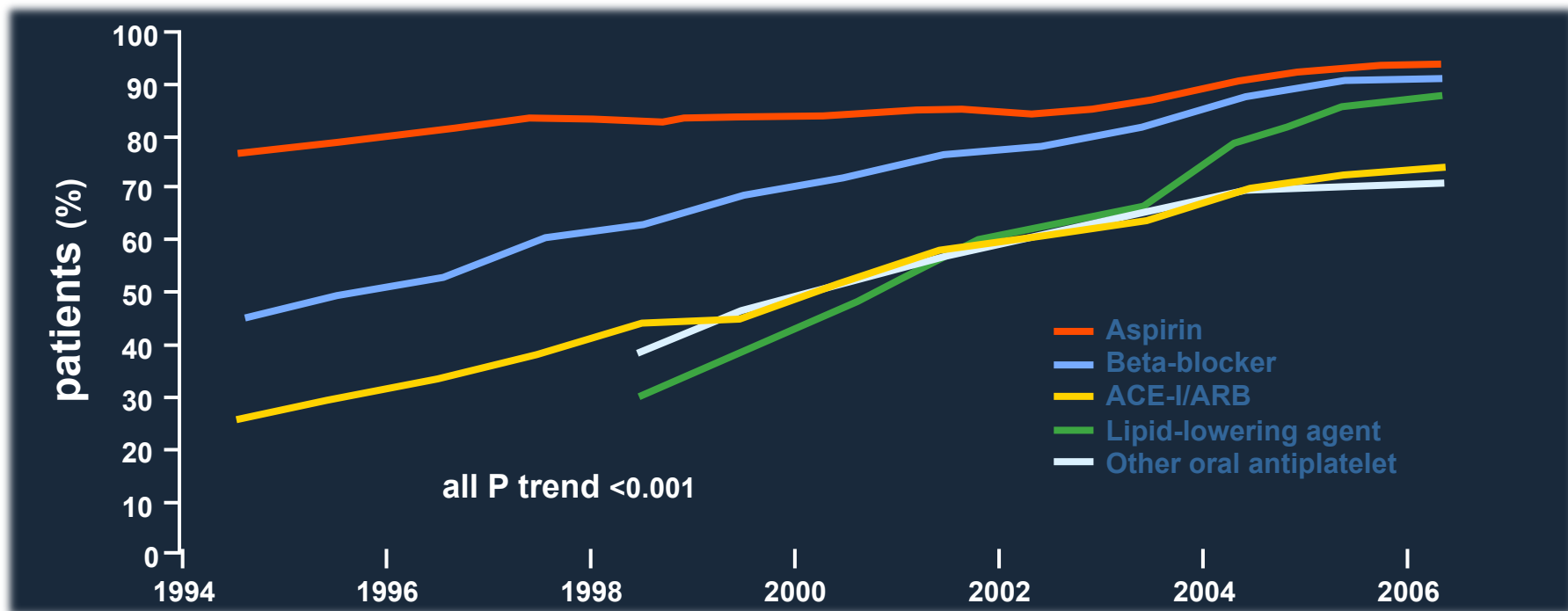
Type of Reperfusion Therapy



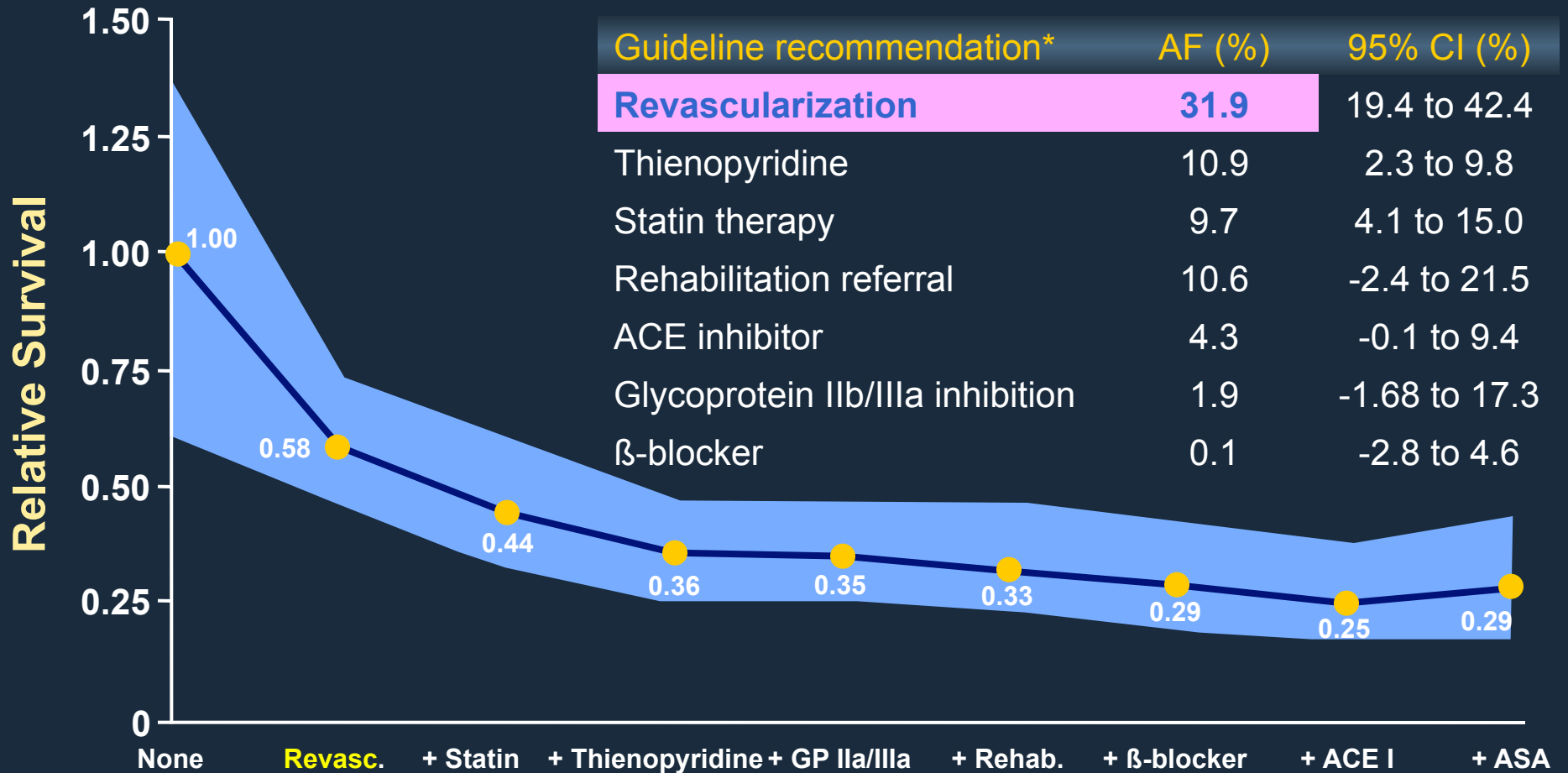
Evolution in Reperfusion

1,146,609 STEMI pts at 2,157 hospitals from 1990-2006

Improvements in Medical Therapy: Discharge Meds



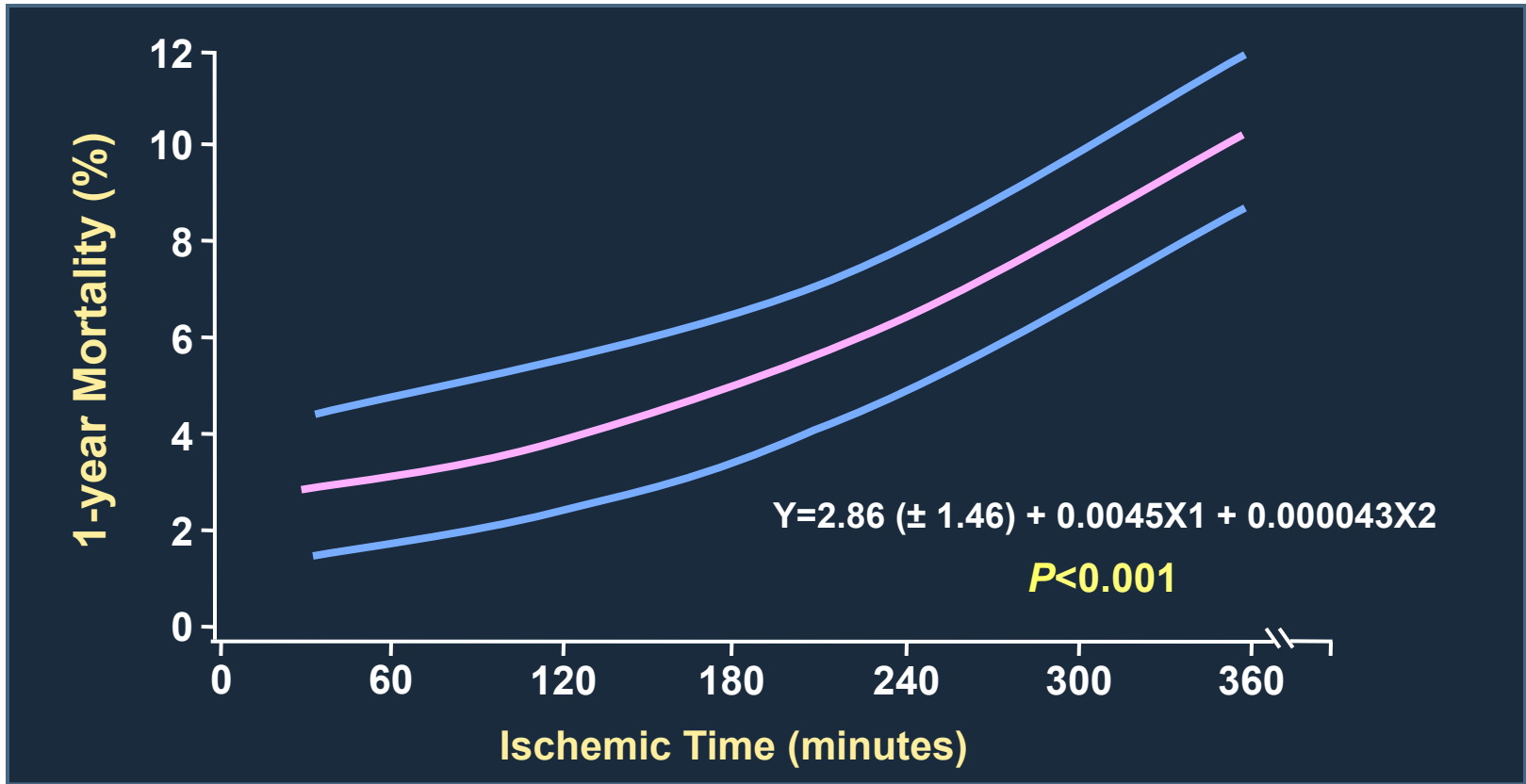
6-Month Survival after Hospital Discharge According to Guideline Recommendations in ACS (GRACE Registry): N=1,716 Cases and 3,432 Controls



Note: 37.2% STEMI; Similar effect in STEMI and NSTEMI

AF = attributable fraction (risk)

TIME from Symptom Onset to Treatment Predicts 1-year Mortality after Primary PCI



the relative risk of 1-year mortality increases by 7.5% for each 30-minute delay

do whatever it takes to **reduce time** from *symptom onset to ER arrival* and time from *ER arrival to PCI*!



Public awareness of MI symptoms

Chest pain centers of excellence
with lower DBTs and excellent
outcomes

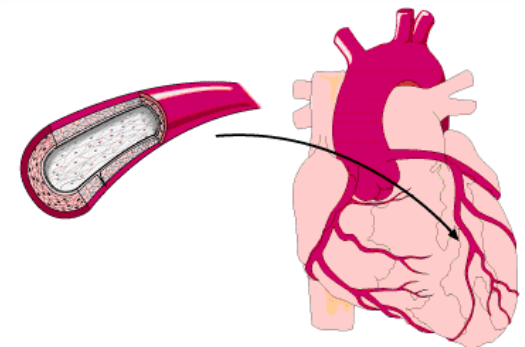
Regional coordination

Ambulance ECG telemetry

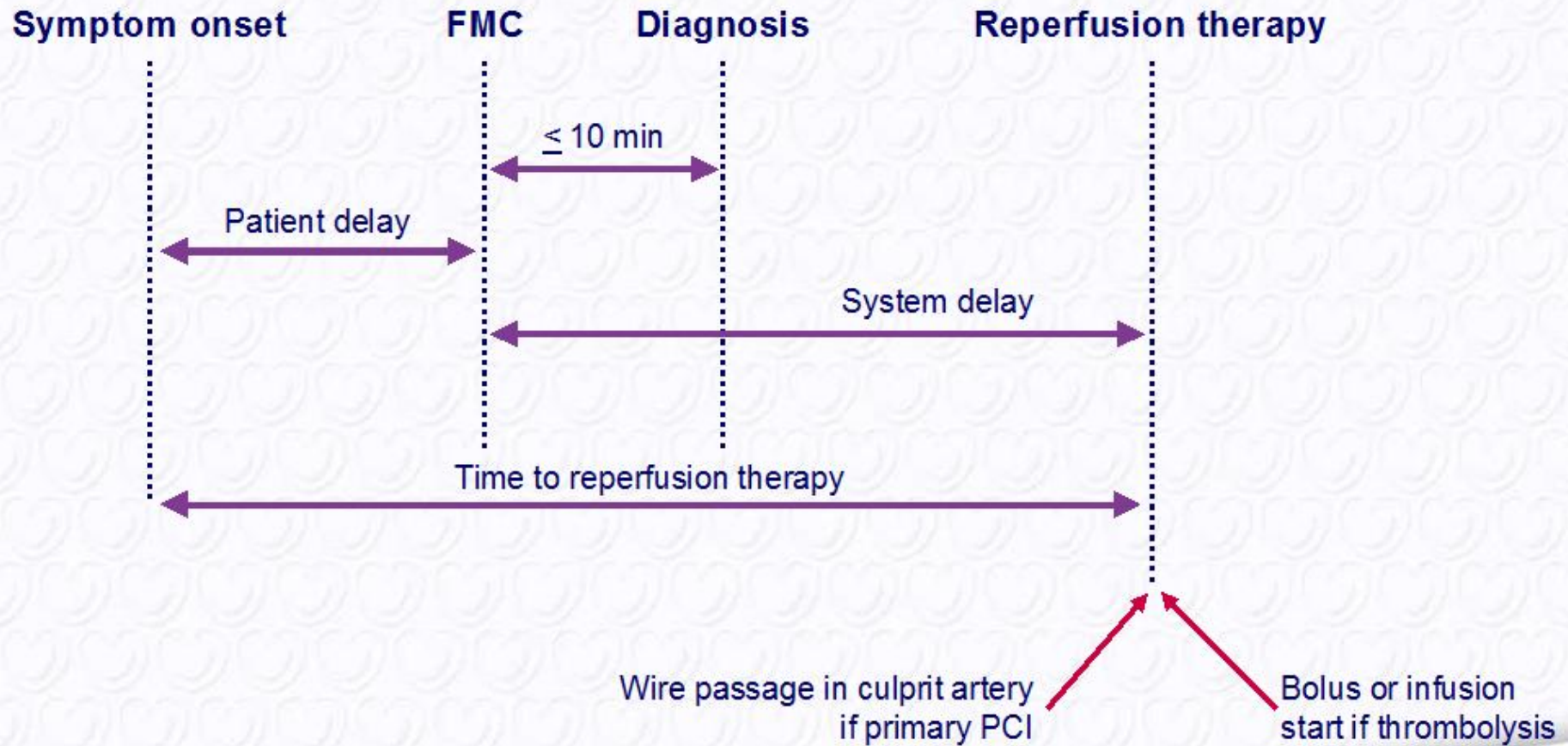
Ambulance/ER CCL activation

ICs sleep in hospital

Continual QI



Components of delay in STEMI and ideal time intervals for intervention



All delays are related to FMC (first medical contact)

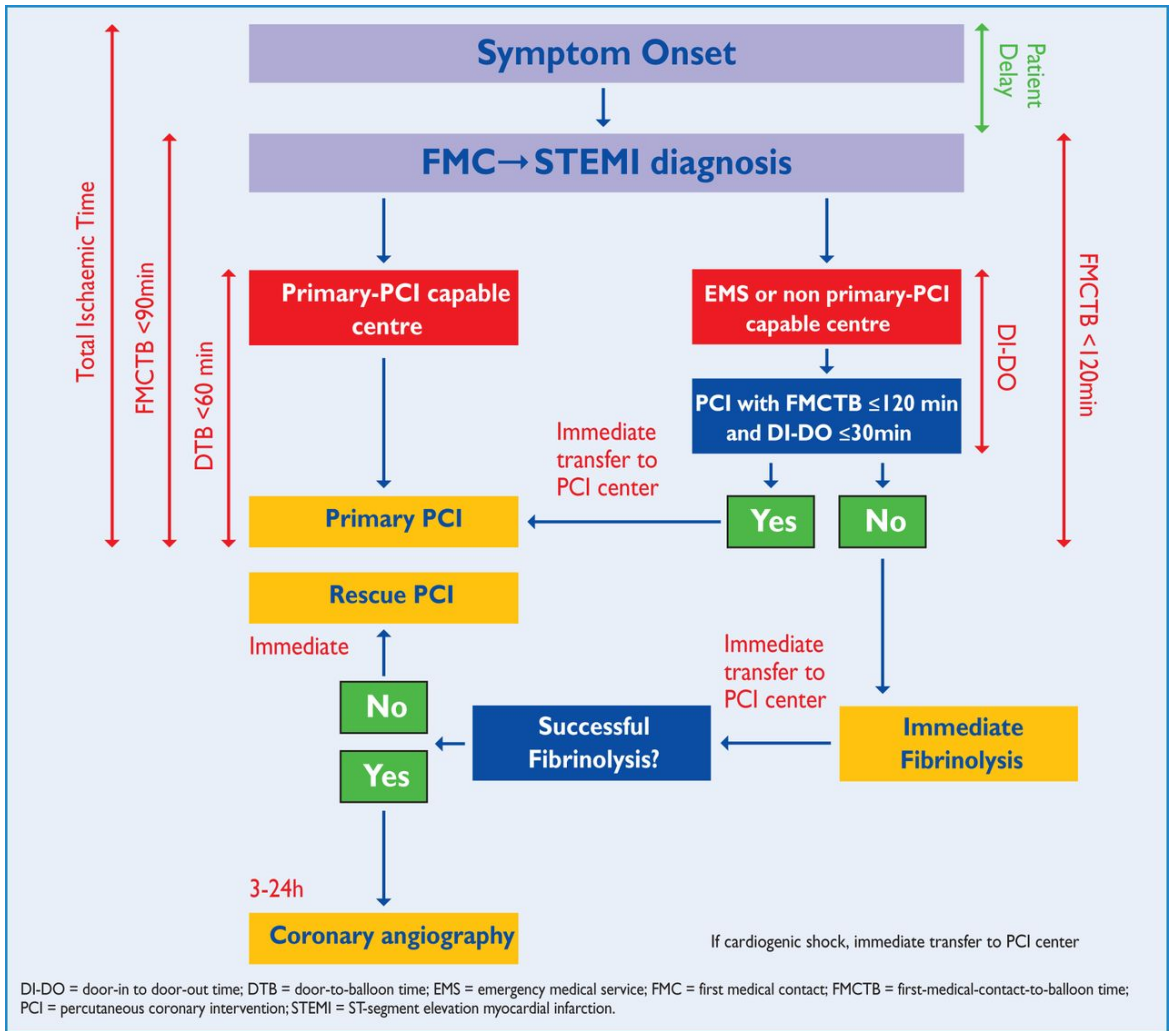
European Heart Journal (2012) 33, 2569–2619
doi:10.1093/eurheartj/ehs215

Important delays and treatment goals in the management of acute STEMI

| Delays | Target |
|--|---|
| Preferred for FMC to ECG and diagnosis. | ≤ 10 min |
| Preferred for FMC to fibrinolysis ('FMC to needle'). | ≤ 30 min |
| Preferred for FMC to primary PCI ('door to balloon') in primary PCI hospitals. | ≤ 60 min |
| Preferred for FMC to primary PCI. | ≤ 90 min (≤ 60 min if early presenter with large area at risk) if this target cannot be met, consider fibrinolysis. |
| Acceptable for primary PCI rather than fibrinolysis. | ≤ 120 min (≤ 90 min if early presenter with large area at risk) if this target cannot be met, consider fibrinolysis. |
| Preferred for successful fibrinolysis to angiography. | 3-24 h |

FMC = first medical contacts; PCI = percutaneous coronary intervention.

Organization of STEMI patient disposal describing pre- and in-hospital management and reperfusion strategies within 12 hours of first medical contact with ideal time interval for interventions.



Authors/Task Force members et al. Eur Heart J 2014;eurheartj.ehu278

Primary PCI for myocardial reperfusion in STEMI: indications and logistics

| Recommendations | Class ^a | Level ^b |
|-----------------|--------------------|--------------------|
|-----------------|--------------------|--------------------|

Indication

| | | |
|---|-----|---|
| Reperfusion therapy is indicated in all patients with time from symptom onset <12 hours duration and persistent ST-segment elevation or (presumed) new LBBB. | I | A |
| Primary PCI is the recommended reperfusion therapy over fibrinolysis if performed by an experienced team in a timely fashion. | I | A |
| In patients with time from symptom onset >12 hours, primary PCI is indicated in the presence of continuing ischaemia, life-threatening arrhythmias or if pain and ECG changes have been stuttering. | I | C |
| Primary PCI is indicated for patients with severe acute heart failure or cardiogenic shock due to STEMI independent from time delay of symptom onset. | I | B |
| Reperfusion therapy with primary PCI should be considered in patients presenting late (12–48 hours) after symptom onset. | IIa | B |

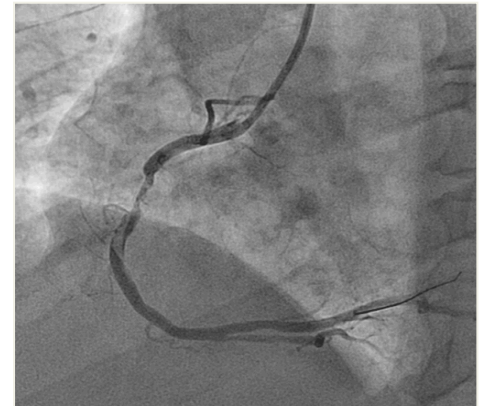
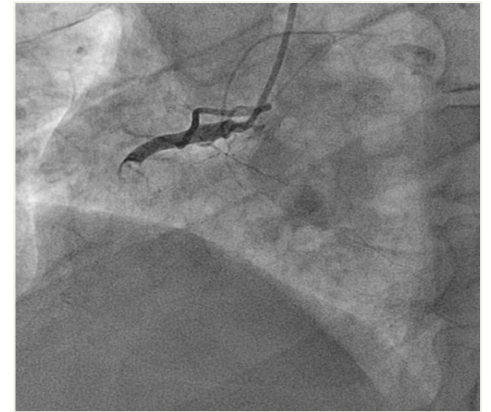
| Recommendations | Class ^a | Level ^b |
|-----------------|--------------------|--------------------|
|-----------------|--------------------|--------------------|

Logistics

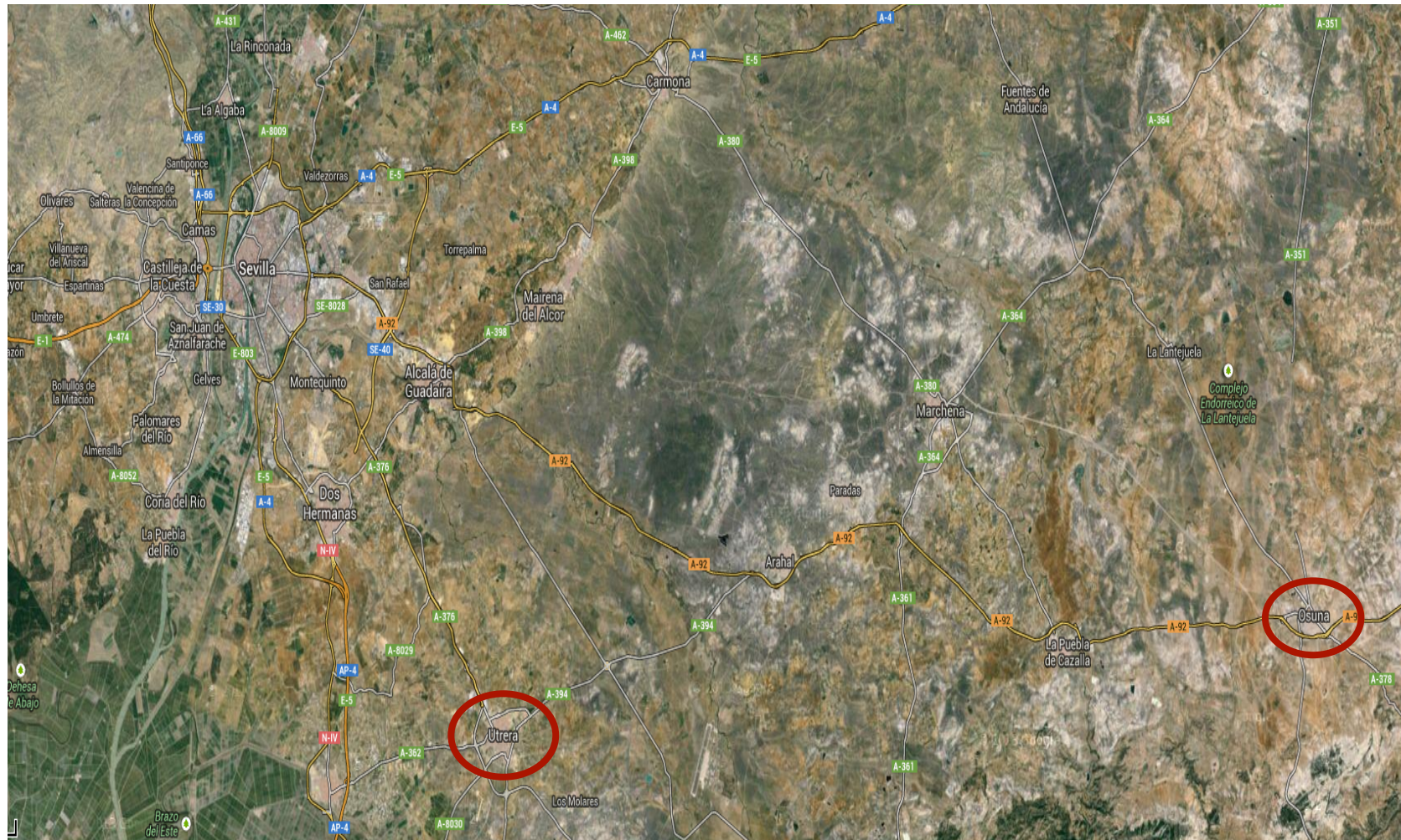
| | | |
|--|-----|---|
| It is recommended that the pre-hospital management of STEMI patients be based on regional networks designed to deliver reperfusion therapy timely and effectively, and to offer primary PCI to as many patients as possible. | I | B |
| It is recommended that all EMSs, emergency departments, coronary care units, and catheterization laboratories have a written updated STEMI management protocol, preferably shared within geographic networks. | I | C |
| It is recommended that primary PCI-capable centres deliver a 24-hour/7-day service and ensure for primary PCI to be performed as fast as possible and at the latest within 60 minutes of hospital arrival. | I | B |
| Patients transferred to a PCI-capable centre for primary PCI should bypass the emergency department and be transferred directly to the catheterization laboratory. | IIa | B |



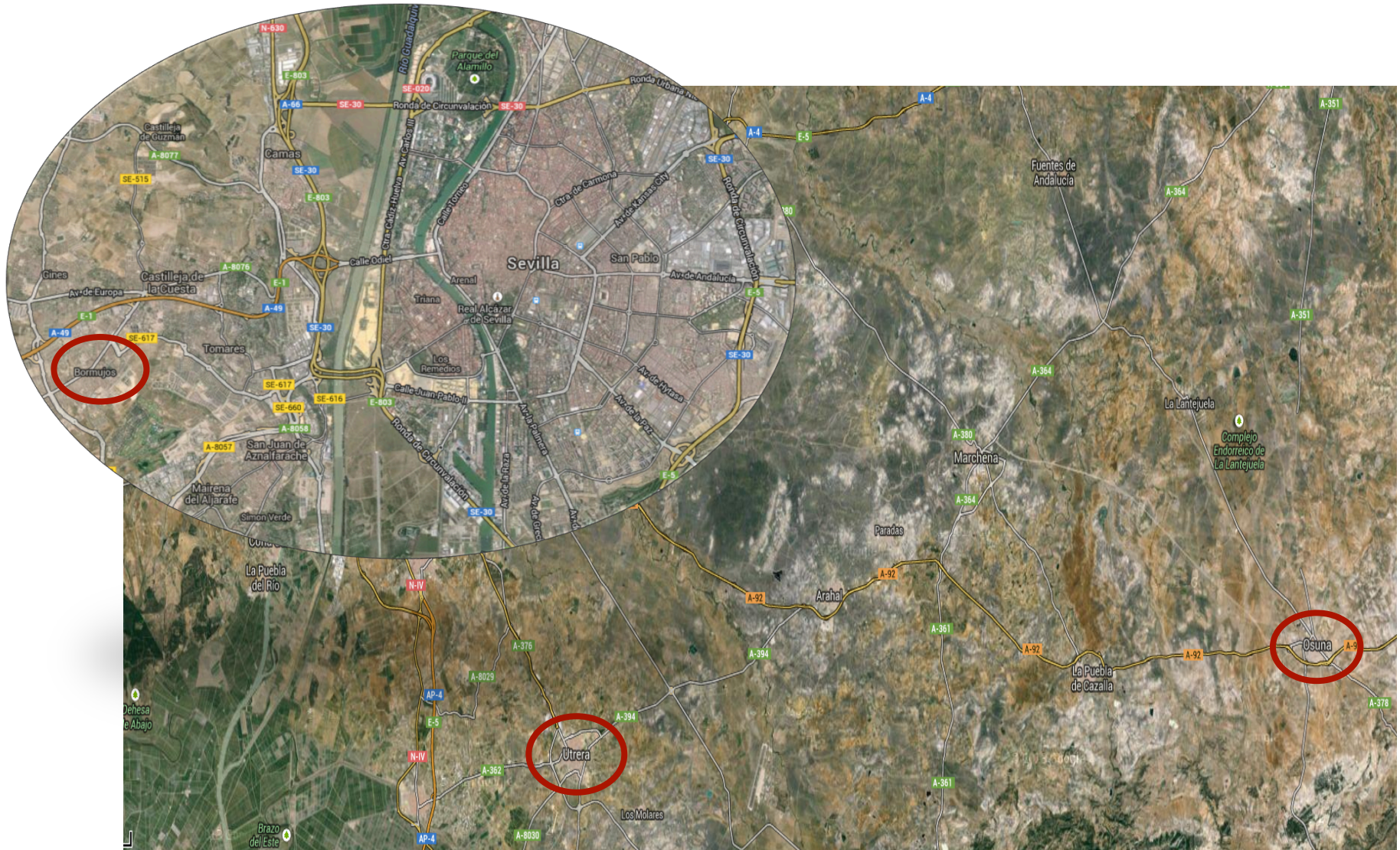
El *Área del Corazón Hospital Virgen del Rocío* posee un *programa de ACTP Primaria* que funciona 24 horas los 365 días del año, de forma que es el método rutinario de reperfusión para todos los infartos con elevación del ST

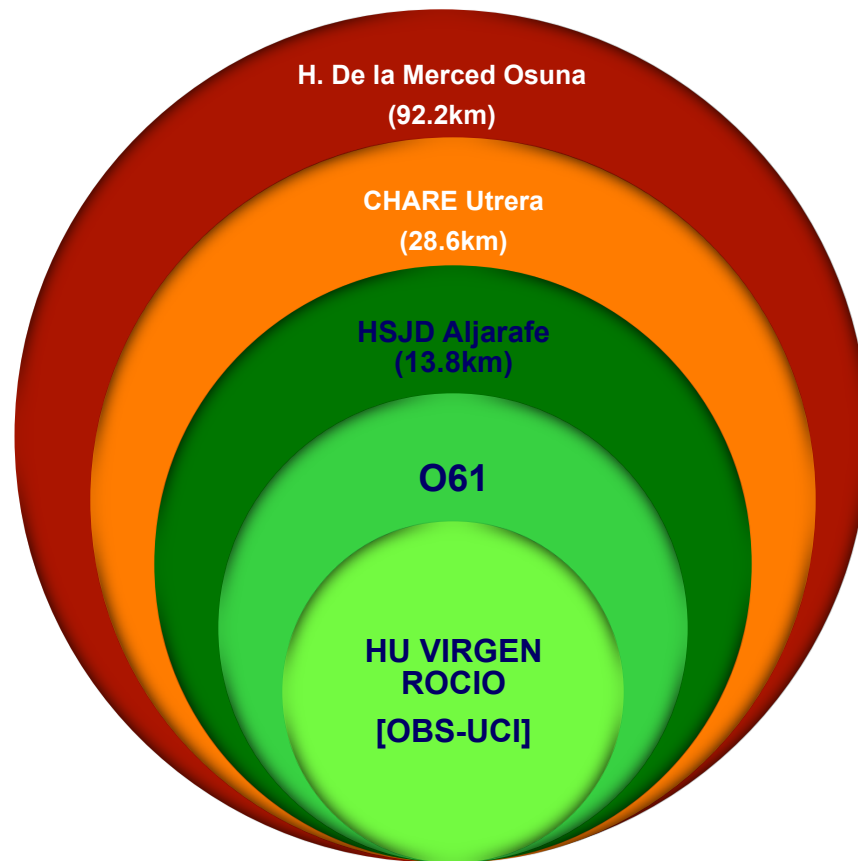


área sanitaria, población asignada 875 331 habitantes



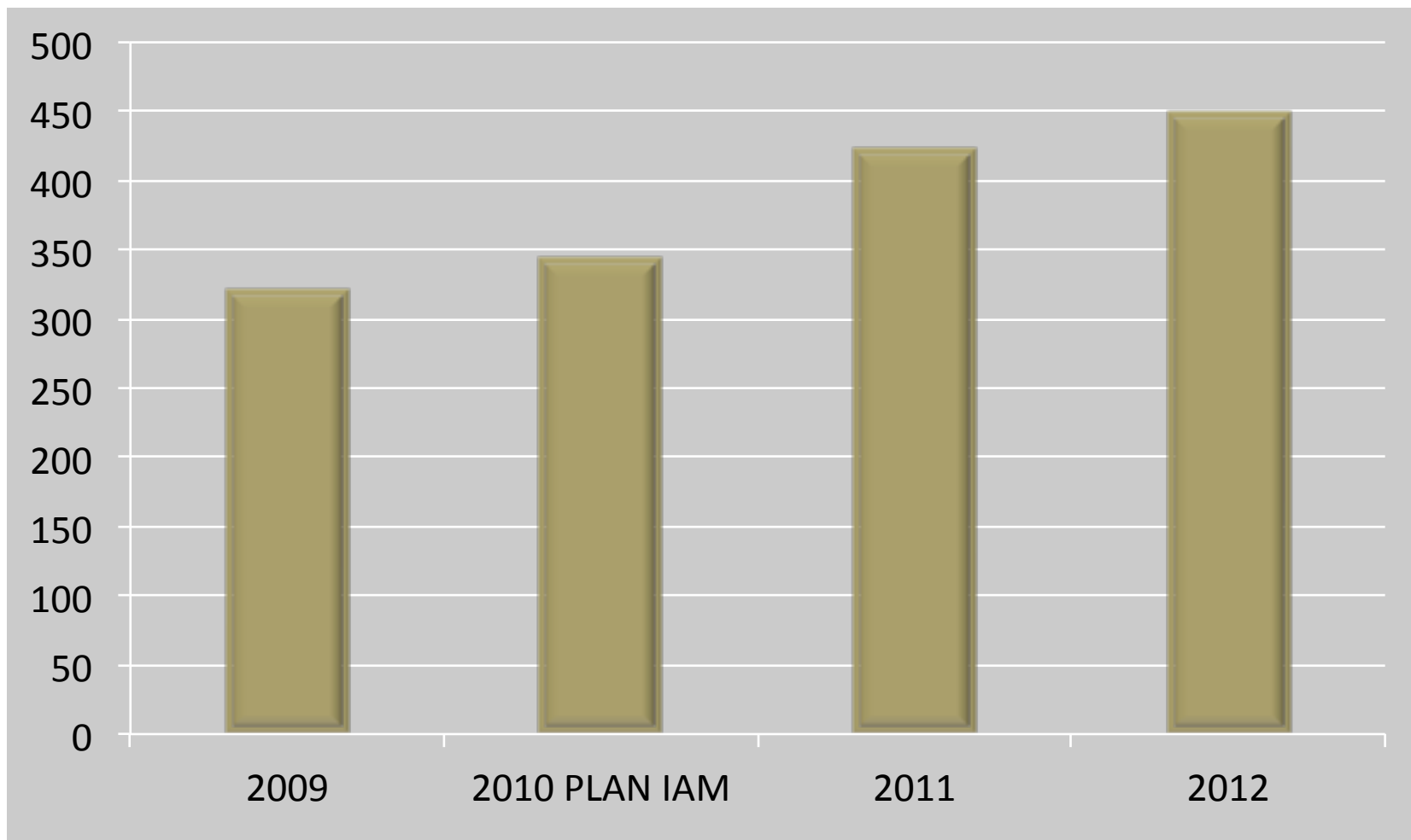
área sanitaria, población asignada 875 331 habitantes





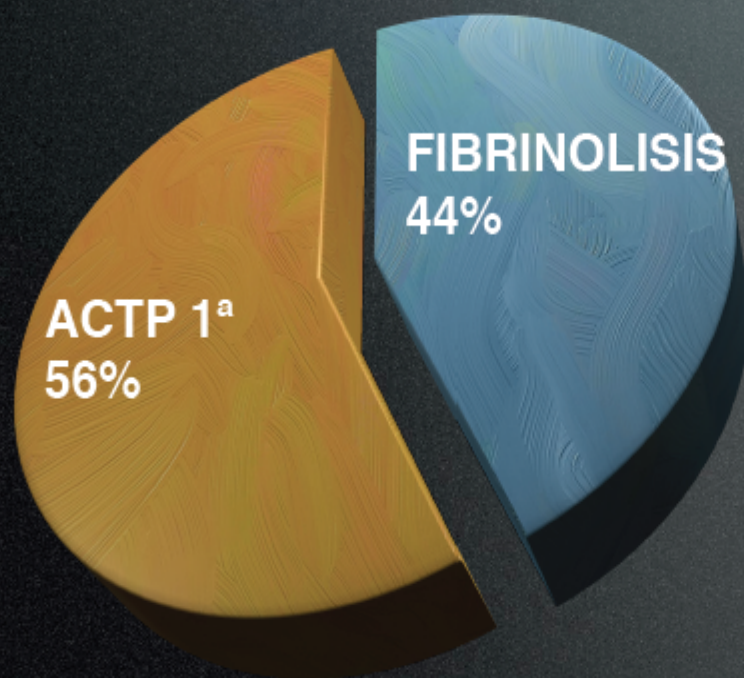
APTC PRIMARIAS PROVINCIA SEVILLA

2009-2012 : INCREMENTO 40 % APTC PRIMARIA

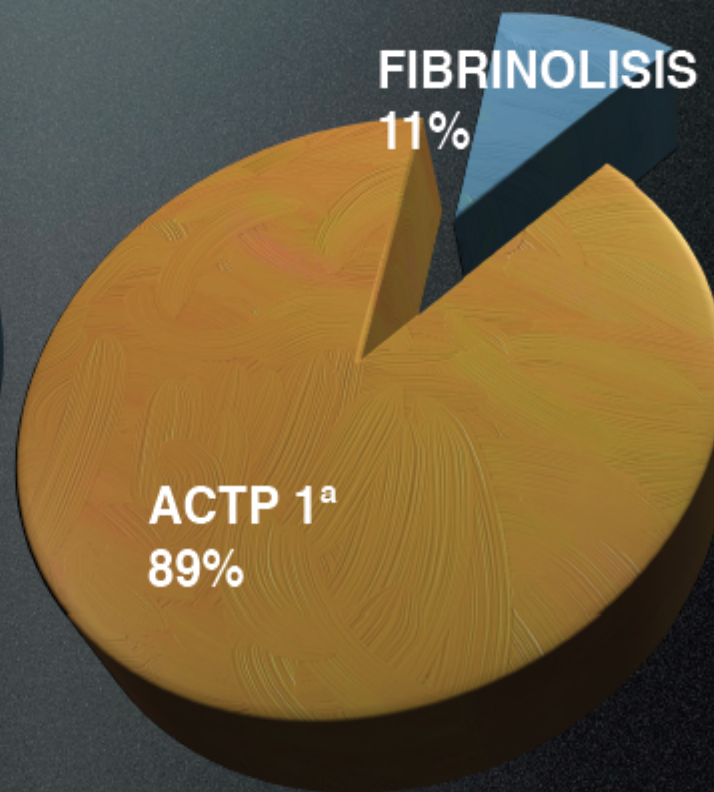


IMPACTO PROGRAMA REPERFUSIÓN

2009

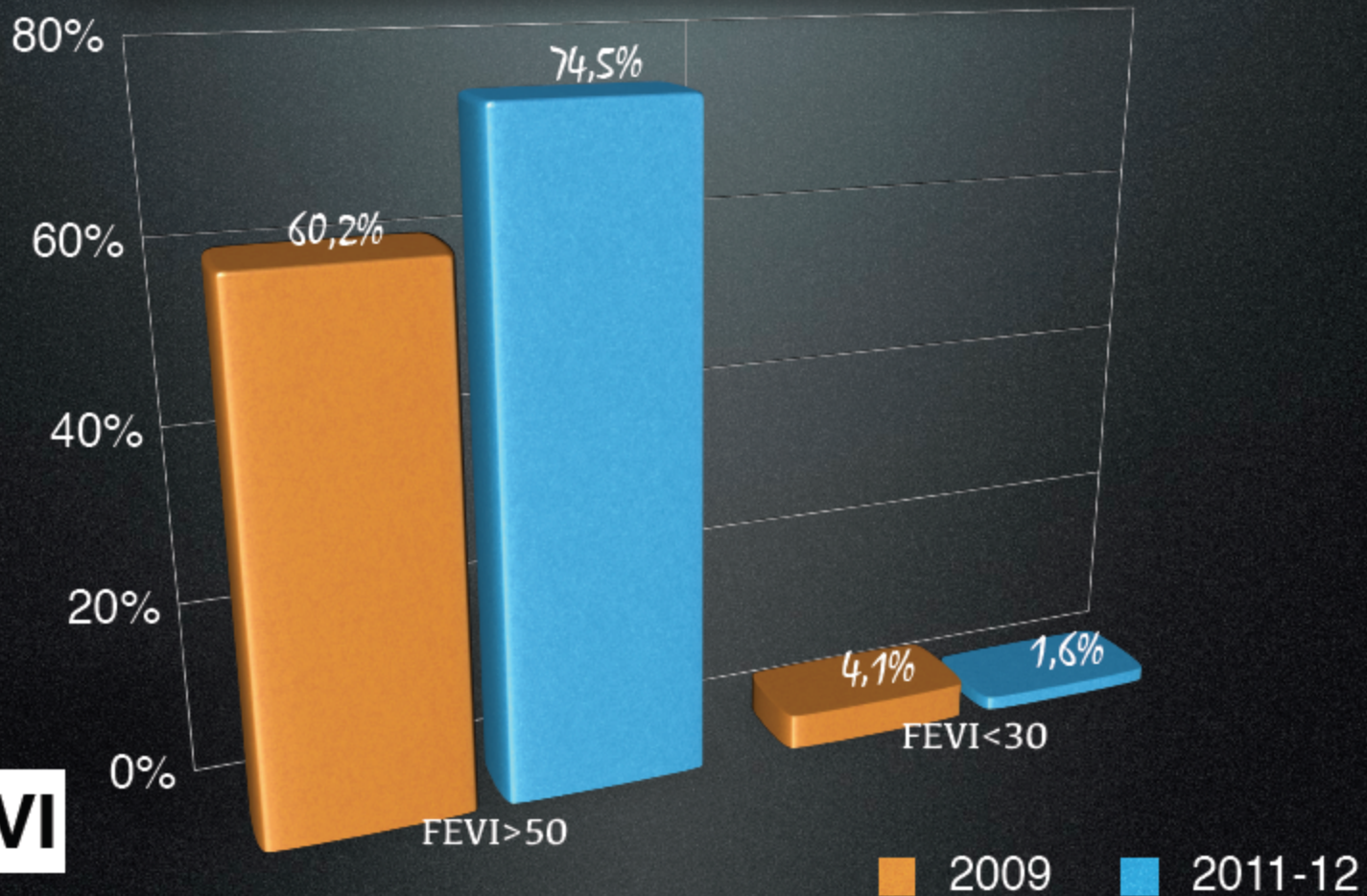


2011-12



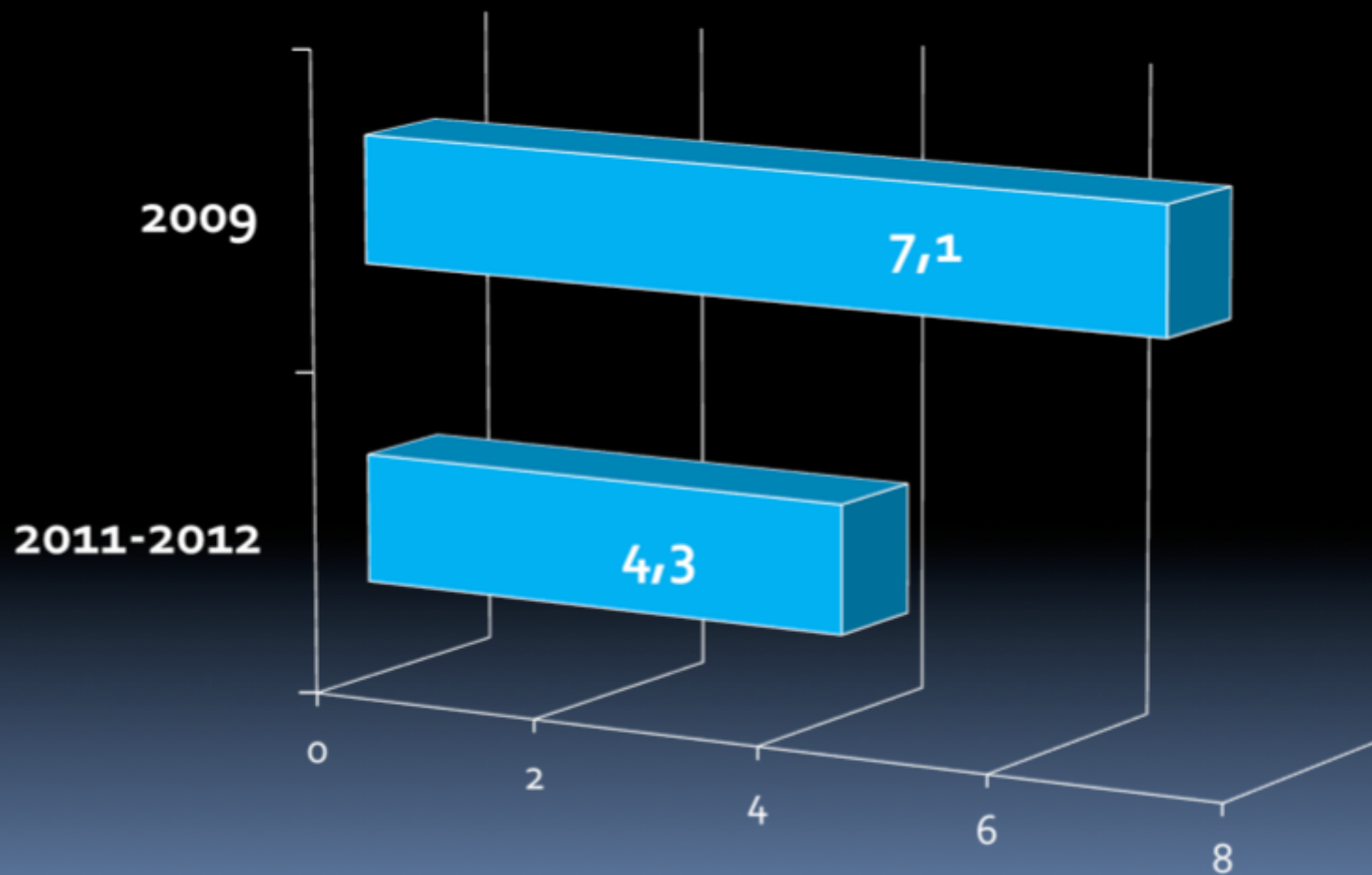
TERAPIA APLICADA

IMPACTO PROGRAMA REPERFUSIÓN



FEVI

MORTALIDAD ALTA UCI



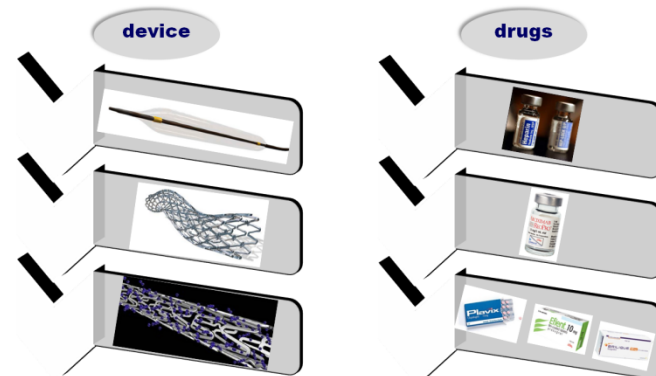
Primary PCI for myocardial reperfusion in STEMI: procedural aspects (strategy and technique)

| Recommendations | Class ^a | Level ^b |
|--|--------------------|--------------------|
| Strategy | | |
| Primary PCI should be limited to the culprit vessel with the exception of cardiogenic shock and persistent ischaemia after PCI of the supposed culprit lesion. | IIa | B |
| Staged revascularization of non-culprit lesions should be considered in STEMI patients with multivessel disease in case of symptoms or ischaemia within days to weeks after primary PCI. | IIa | B |
| Immediate revascularization of significant non-culprit lesions during the same procedure as primary PCI of the culprit vessel may be considered in selected patients. | IIb | B |
| In patients with continuing ischaemia and in whom PCI of the infarct-related artery cannot be performed, CABG should be considered. | IIa | C |

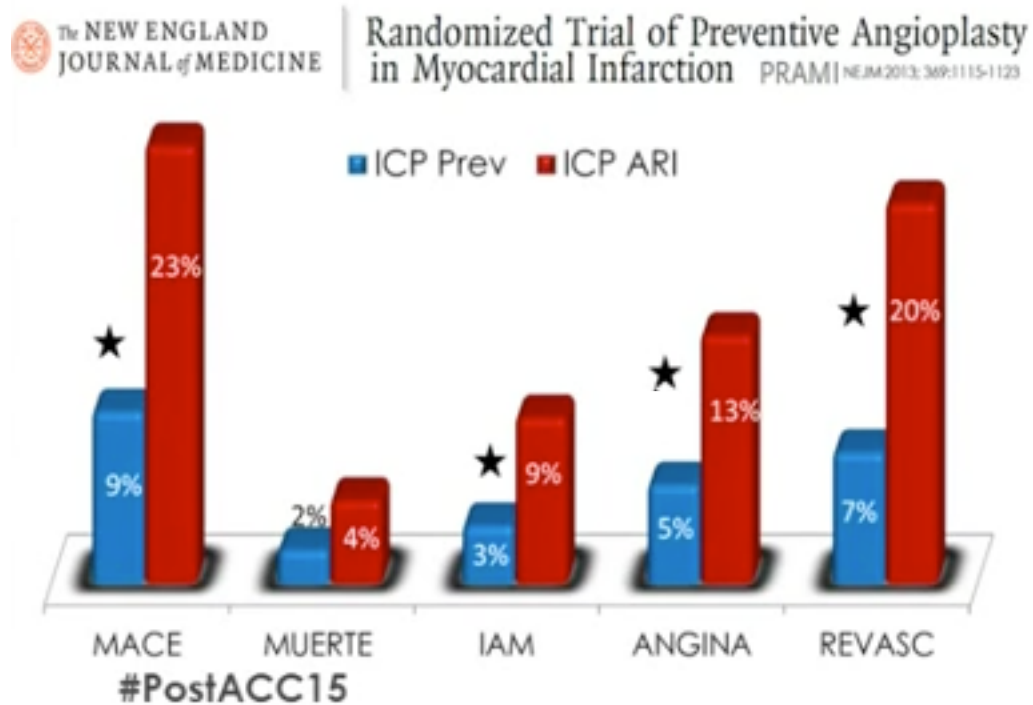
Achieve all four parameters of successful reperfusion

- Relief of chest pain
 - ST-segment resolution
 - Restoration of TIMI-3 flow
 - Myocardial perfusion grade 3-4
- Hemodynamic and electrical instability
 - Must act quickly

Evolution of Primary PCI



PRAMI: “Preventative” PCI of non-culprit lesions after culprit lesion primary PCI in STEMI

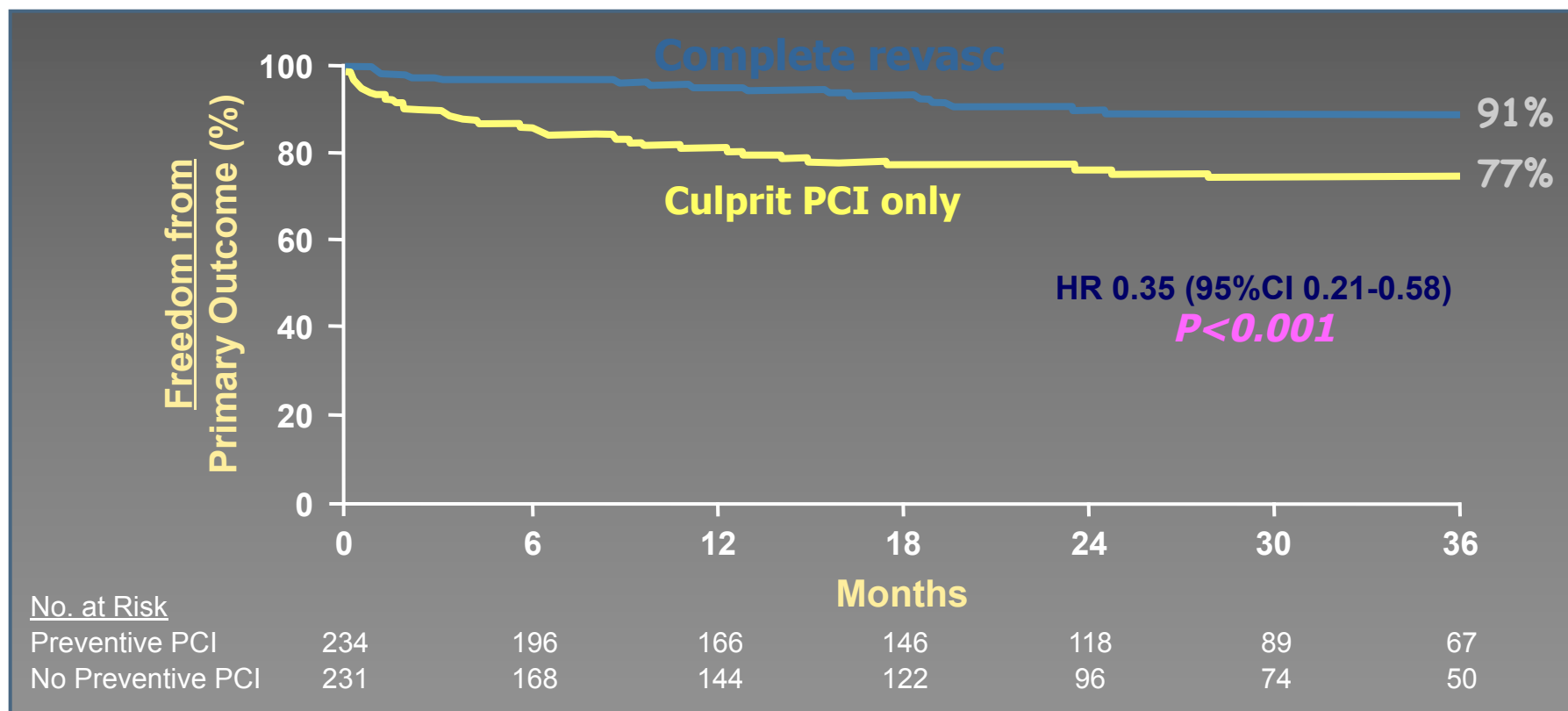


Gershlick et al. ESC 2014

PRAMI: "Preventative" PCI of non-culprit lesions after culprit lesion primary PCI in STEMI

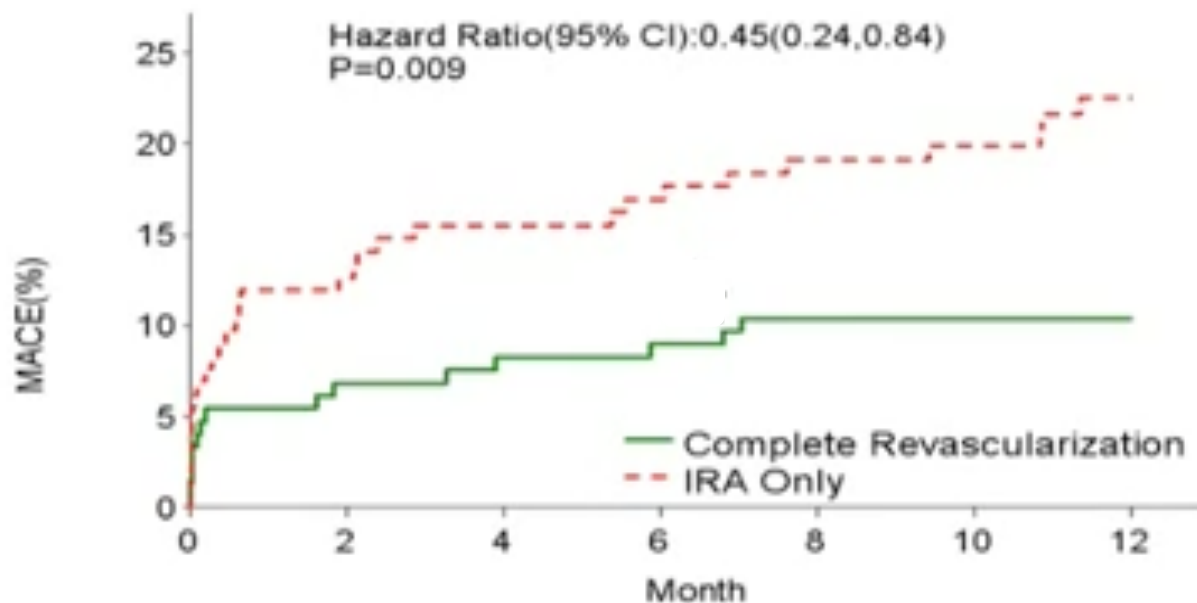
465 non-shock pts at 5 UK sites with MVD after successful primary PCI randomized to NCL PCI of non-LM DS 50-99% stenoses vs. conservative care
600 pts planned; DSMB stopped trial early after 465 pts enrolled (2008-2013)

Primary endpoint: Cardiac death, MI or refractory angina



CULPRIT - MACE

(total mortality, recurrent MI, heart failure, revascularisation)

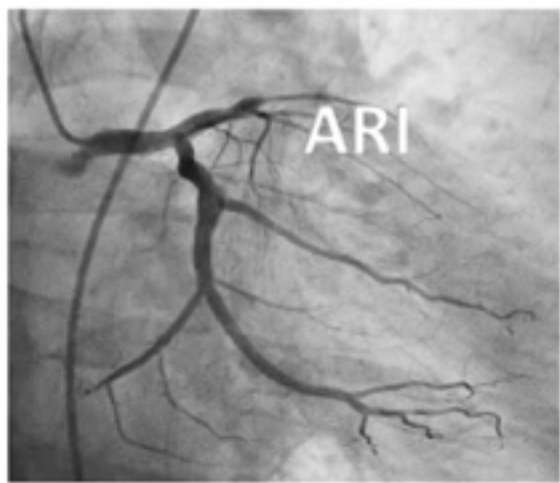


Number at risk:

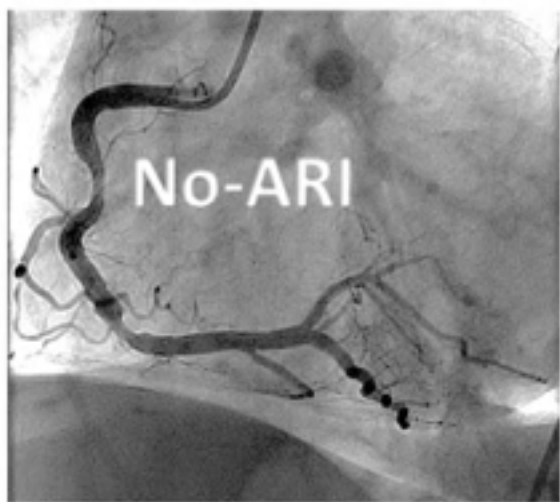
| | | | | | | |
|--------------|-----|-----|-----|-----|-----|----|
| Complete:150 | 131 | 129 | 128 | 125 | 108 | 73 |
| IRA Only:146 | 122 | 118 | 116 | 111 | 98 | 68 |

Gershlick *et al.* ESC 2014

DANAMI3-PRIMULTI



- **30-50%** pacientes sometidos ICPP presentan enfermedad multivaso.
- Las guías recomiendan ICP sólo de la ARI.
- Existen evidencias a favor de revascularizar el resto de lesiones.



Guías Europeas
de
Revascularización

| Enf. Multivaso | | |
|---------------------------|-----|---|
| ICPP Solo ARI | IIa | B |
| Diferir el resto de vasos | IIa | B |
| ICPP todos los vasos | IIb | B |

#PostACC15



627

IAM Multivaso
Aleat tras éxito ARI

313

ICPP ARI



314

ICPP + Revasc
guiada x FFR
diferida

DANAMI3-PRIMULTI

| | IRA only (n = 313) | Complete revascularisation (n = 314) |
|--------------------------|-----------------------|--|
| Age (years) | 64 (range 34 – 92) | 64 (range 37 – 94) |
| Male | 255 (82%) | 251 (80%) |
| Medical history | | |
| Diabetes | 42 (13%) | 29 (9%) |
| Hypertension | 146 (47%) | 130 (41%) |
| Current smoking | 151 (48%) | 160 (51%) |
| Previous MI | 27 (9%) | 17 (5%) |
| Infarct location | | |
| Anterior | 112 (36%) | 105 (33%) |
| Inferior | 179 (57%) | 195 (62%) |
| Posterior | 20 (6%) | 10 (3%) |
| Three vessel disease | 100 (32%) | 97 (31%) |
| Stenosis on proximal LAD | 86 (28%) | 80 (26%) |

#PostACC15



627

IAM Multivaso
Aleat tras éxito ARI

313

ICPP ARI



314

ICPP + Revasc
guiada x FFR
diferida

DANAMI3-PRIMULTI

| | IRA only (n = 313) | Complete revascularisation (n = 314) | P |
|--|-----------------------|--|---------|
| Procedure duration (min) | 42 (31 – 59) | 76 (56 – 100) | <0.0001 |
| Contrast volume (ml) | 170 (125 – 220) | 280 (215 – 365) | <0.0001 |
| Fluoroscopy dose (Gycm ²) | 49 (33 – 74) | 77 (52 – 115) | <0.0001 |
| Nº arteries treated per patient | 1 (1–2) | 2 (1–3) | <0.0001 |
| Number of implanted stents | 1 (1–1) | 2 (1–3) | <0.0001 |
| Stent diameter (mm) | 3.5 (2.75–3.5) | 3.0 (2.75–3.5) | 0.005 |
| Total stent length (mm) | 18 (15–28) | 33 (18–51) | <0.0001 |
| Stent type | | | 0.5 |
| No stenting | 18 (6%) | 12 (4%) | |
| Bare-metal | 5 (2%) | 3 (1%) | |
| Drug-eluting | 290 (93%) | 298 (96%) | |
| Use of Glycoprotein IIb/IIIa inhibitor | 72 (23%) | 64 (20%) | 0.4 |
| Use of Bivalirudin | 234 (75%) | 237 (76%) | 0.8 |

#PostACC15



627

IAM Multivaso
Aleat tras éxito ARI

313

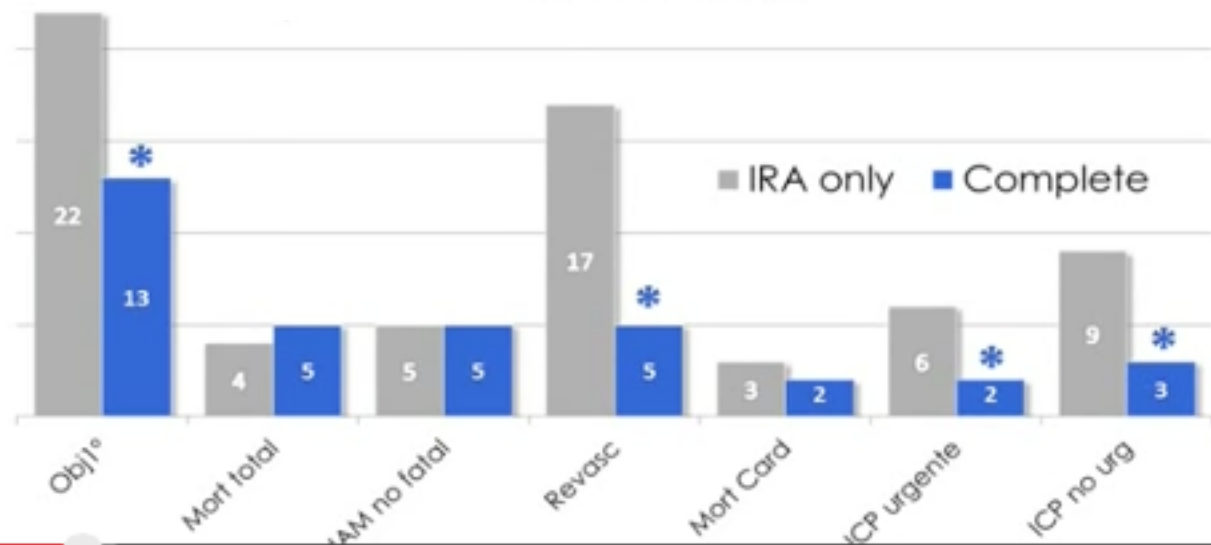
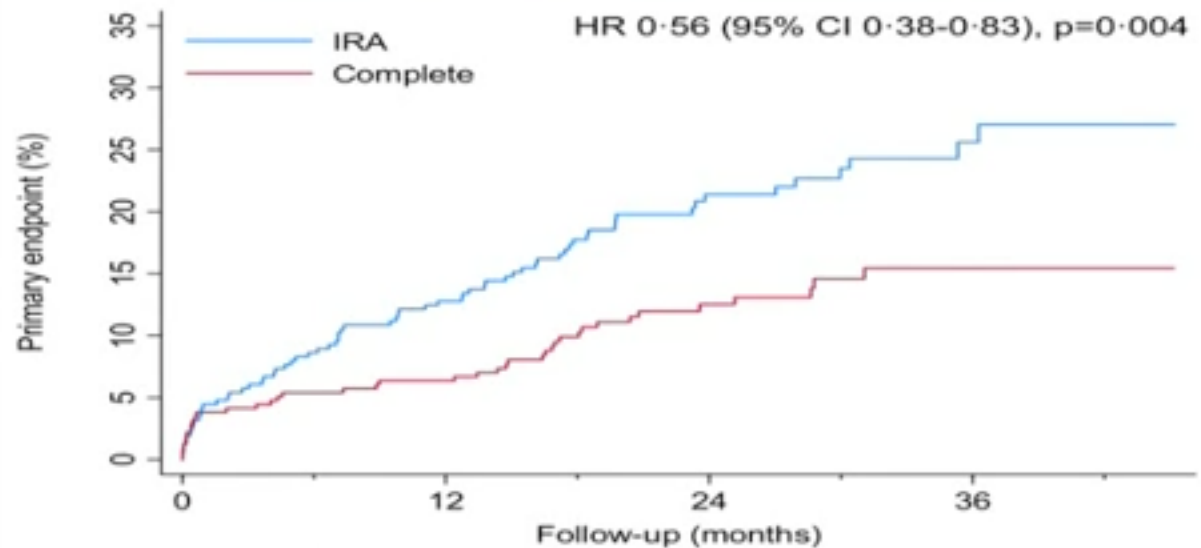
ICPP ARI



314

ICPP + Revasc
guiada x FFR
diferida

DANAMI 3-PRIMULTI



DANAMI 3-PRIMULTI

CONCLUSIONES

La revascularización guiada por FFR en pacientes

- **Multivaso**
- ICPP con **éxito** de **ARI**

1. Disminuye el objetivo 1º
2. Disminuye 40% la tasa de revasc. Urgente
3. **No** disminuye los eventos duros.

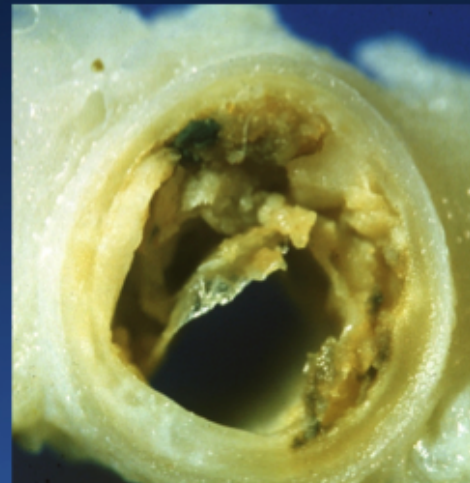
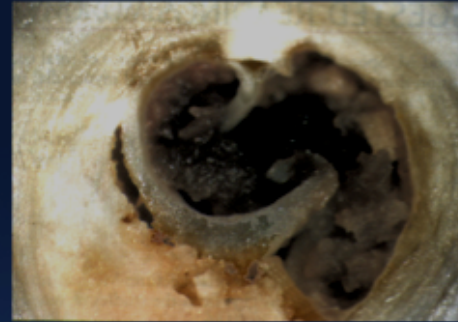
Por tanto

La estrategia de solo ARI es recomendable ante cualquier dificultad anticipada sobre No-ARI

| Recommendations | Class ^a | Level ^b |
|---|--------------------|--------------------|
| Technique | | |
| Stenting is recommended (over balloon angioplasty) for primary PCI. | I | A |
| New-generation DES are recommended over BMS in primary PCI. | I | A |
| Radial access should be preferred over femoral access if performed by an experienced radial operator. | IIa | A |
| Thrombus aspiration may be considered in selected patients | IIb | A |



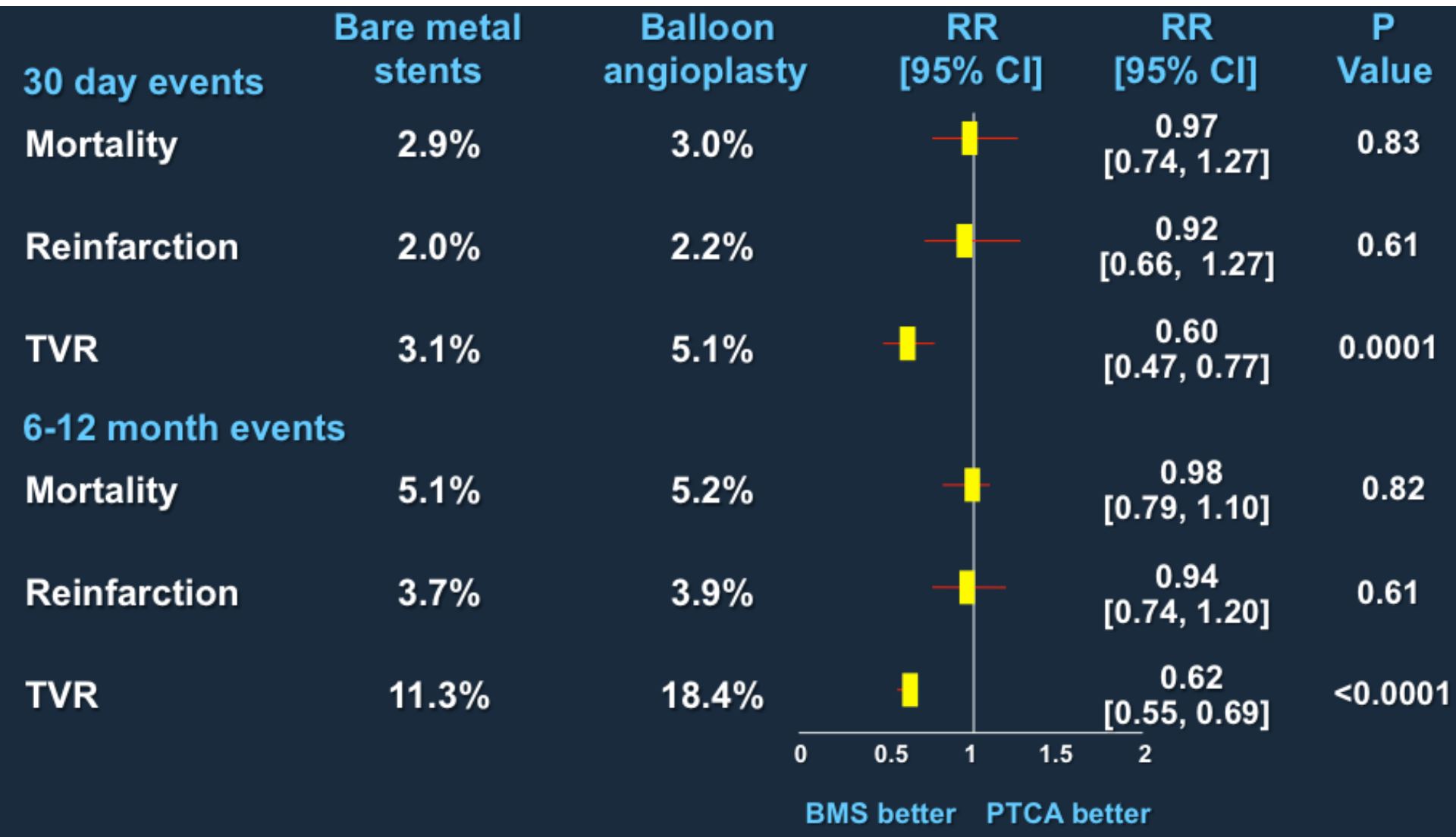
Limitations of Balloon Angioplasty in AMI



Post POBA

13 RCTs of BMS vs. Balloon PTCA in AMI

N=6922



DESERT: Pooled pt-level meta-analysis from 11 of 13 RCTs of SES/PES vs. BMS in primary PCI

N=6298. mean FU 1201 ± 440 days

Cox models with piecewise time-constant regression coefficients

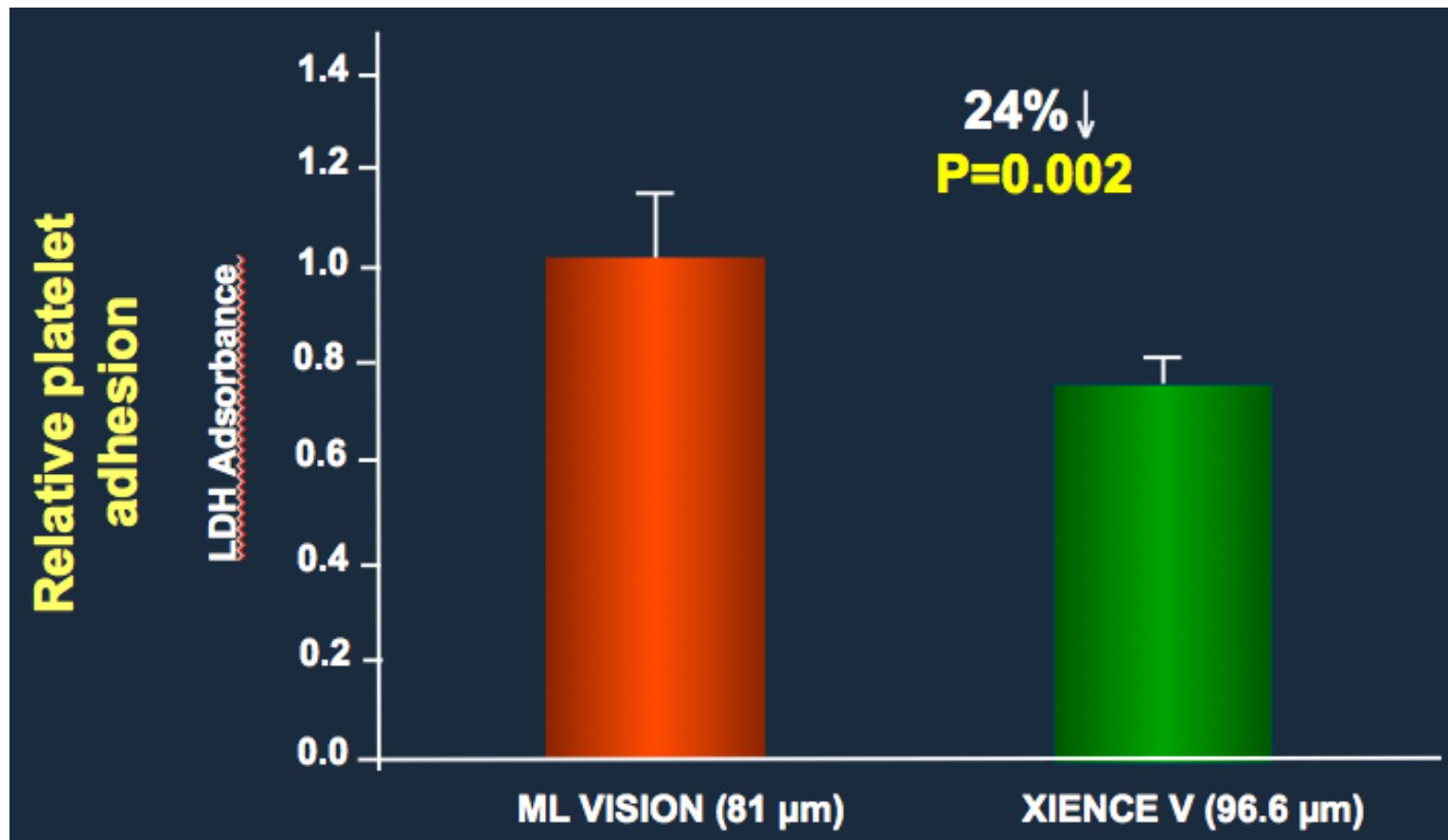
| Outcome | Interval | HR (95%CI)* | p-value |
|------------------|-----------|------------------|------------------|
| Death | 0-1 year | 0.93 (0.70-1.22) | 0.14 |
| | 1-2 years | 0.58 (0.37-0.93) | 0.23 |
| | >2 years | 0.93 (0.65-1.33) | 0.18 |
| Reinfarction | 0-1 year | 0.85 (0.63-1.16) | 0.31 |
| | 1-2 years | 1.34 (0.81-2.23) | 0.25 |
| | >2 years | 2.06 (1.22-3.49) | 0.03 |
| Stent thrombosis | 0-1 year | 0.90 (0.66-1.23) | 0.52 |
| | 1-2 years | 1.38 (0.70-2.71) | 0.35 |
| | >2 years | 2.81 (1.28-6.19) | 0.04 |
| TVR | 0-1 year | 0.48 (0.41-0.58) | <0.001 |
| | 1-2 years | 0.66 (0.49-0.87) | 0.01 |
| | >2 years | 0.93 (0.62-1.38) | 0.71 |

*HR
for DES
Vs BMS

Stent Thrombosis is Affected by Stent Design, Deployment and Polymer

Impact of Xience / Promus polymer coating

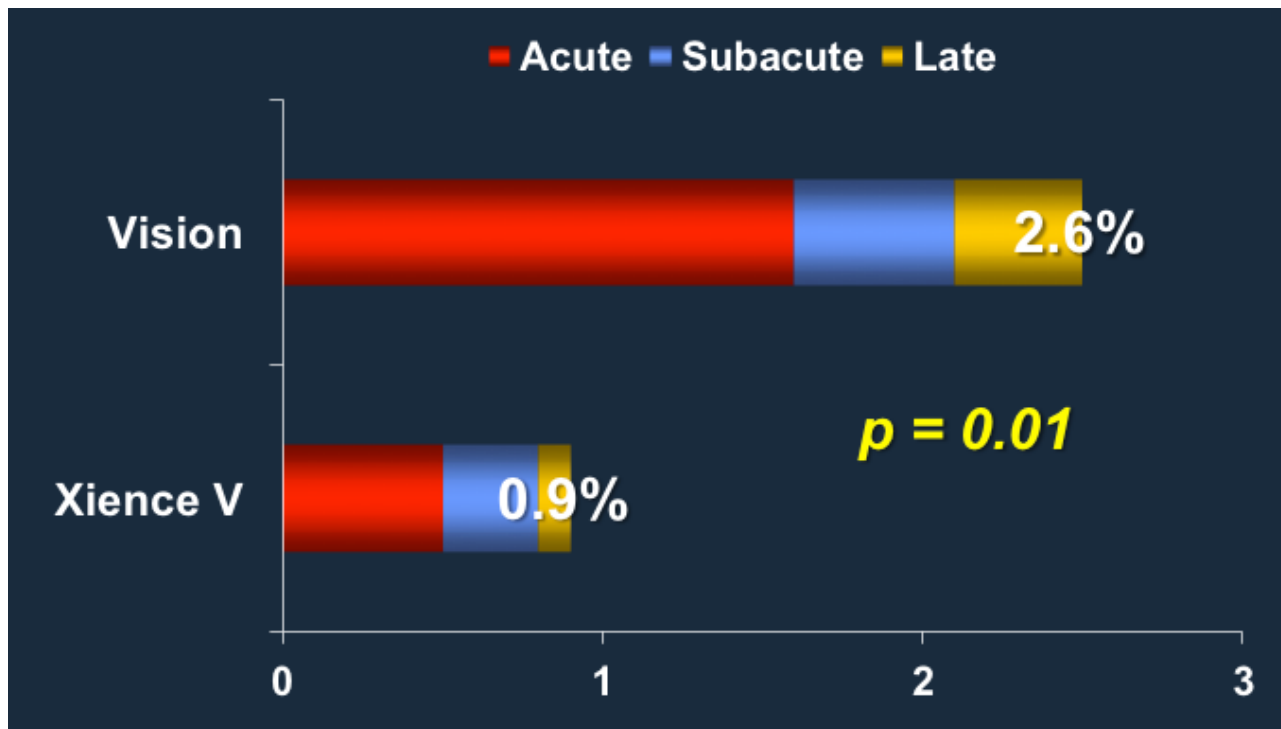
In vitro pulsatile Chandler loop model with porcine blood



EXAMINATION Trial

1504 pts with STEMI undergoing PCI within 48° (85% primary PCI within 12°) were randomized to Xience V EES vs. Vision BMS

Stent thrombosis (Def/prob) within 1 year



Definite ST was reduced with Xience V from 1.9% to 0.5%, $p=0.01$



78 



8404

SCACEST+
SCASEST
Con 1 FR
>60°, MDM(+)
Isquemia ECG

SCA sometido a ICP





78 

8404

Biv vs
HPN+IIb/IIIa



SCA sometido a ICP

Apilino®(2012)

4197

1:1

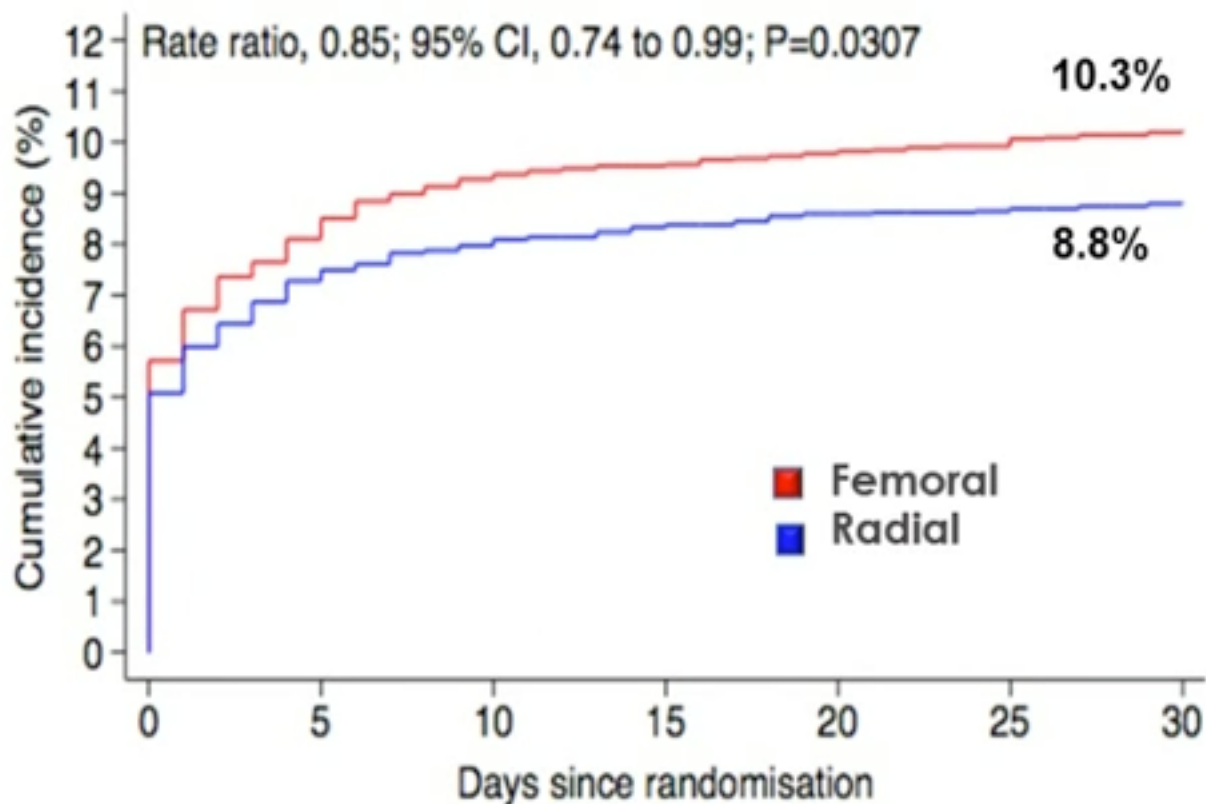
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Trans-Radial
Access

Trans-Femoral
Access

EFICACIA: MACE

Mortalidad+IAM+ACV



#PostACC15



78



8404

Biv vs
HPN+IIb/IIIa



SCA sometido a ICP

Aspirin+P2Y12

1:1

4197

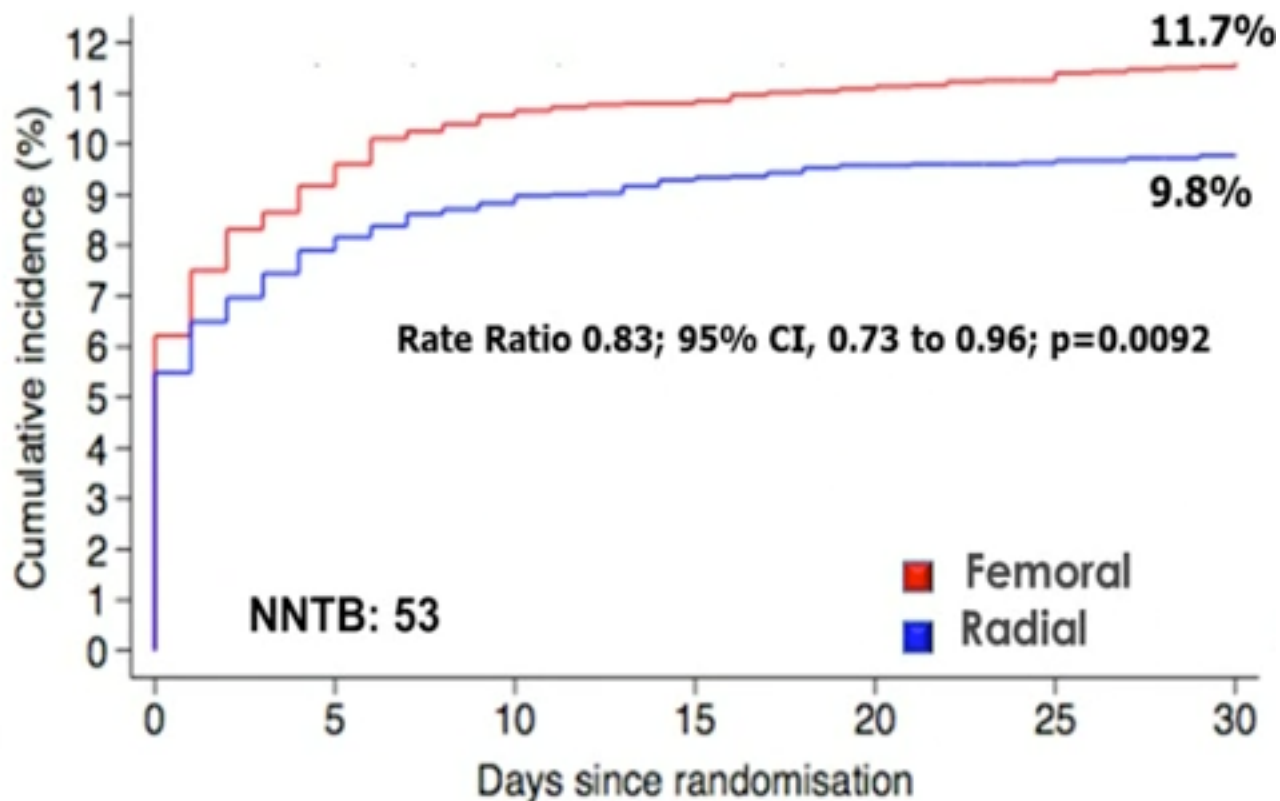
4207

Trans-Radial
Access

Trans-Femoral
Access

EFICACIA+SEGURIDAD: NACE

Mortalidad+IAM+ACV+Sangrado BARC 3-5



#PostACC15



78 

8404

Biv vs
HPN+IIb/IIIa

CONCLUSIONES

En **SCA+ICP**, el acceso radial

1. Disminuye la mortalidad
2. Disminuye los sangrados no mortales.
3. **NNTB: 53**

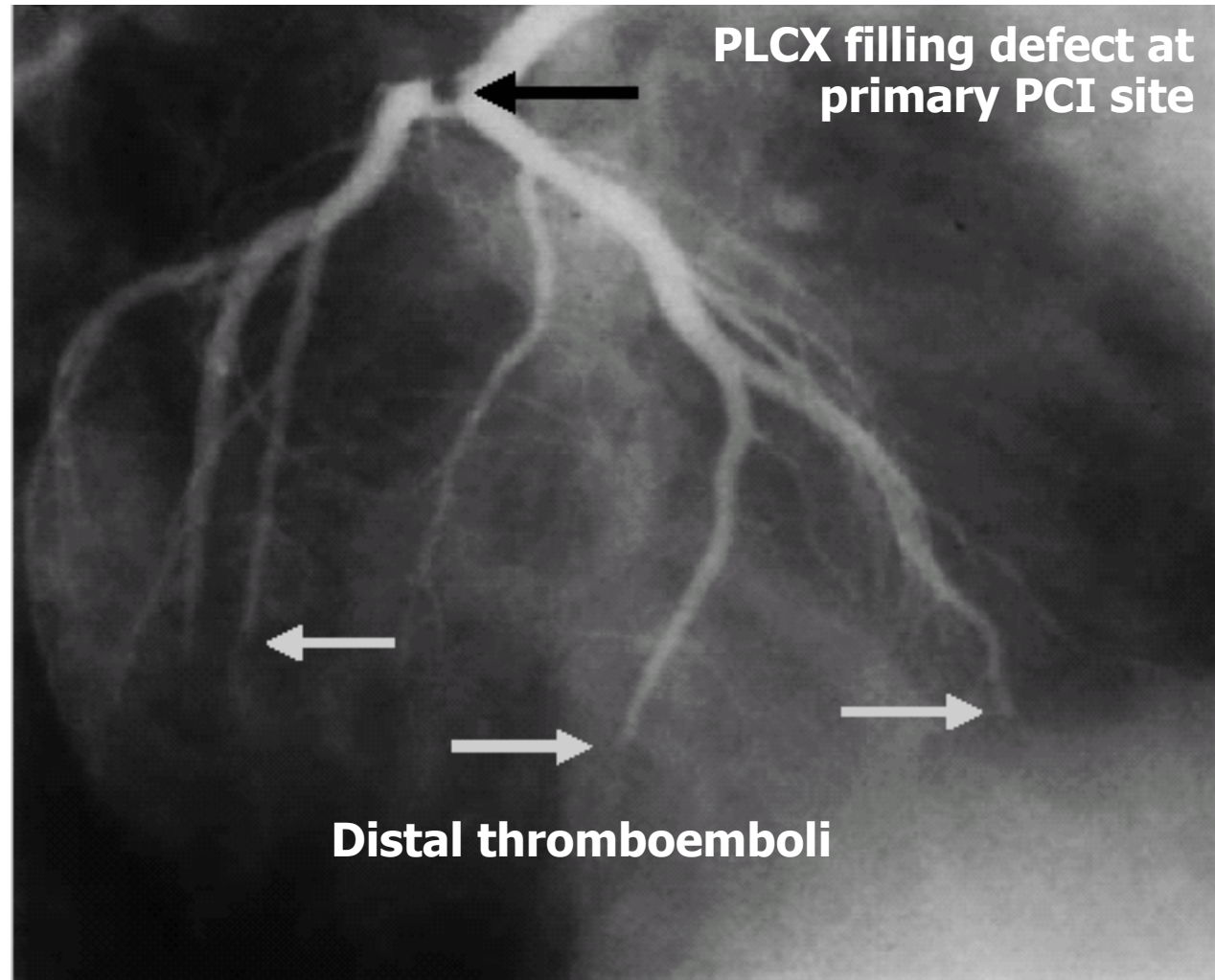
Por tanto

Estrategia de **elección** en SCA + ICP
independientemente del régimen de
anticoagulación.

Impact of Macroscopic Distal Emboli (DE)

**DE occurred in
27 of 178
(15%) pts after
primary PTCA**

- ↓ ST res
- ↑ Infarct size
- ↑ Mortality

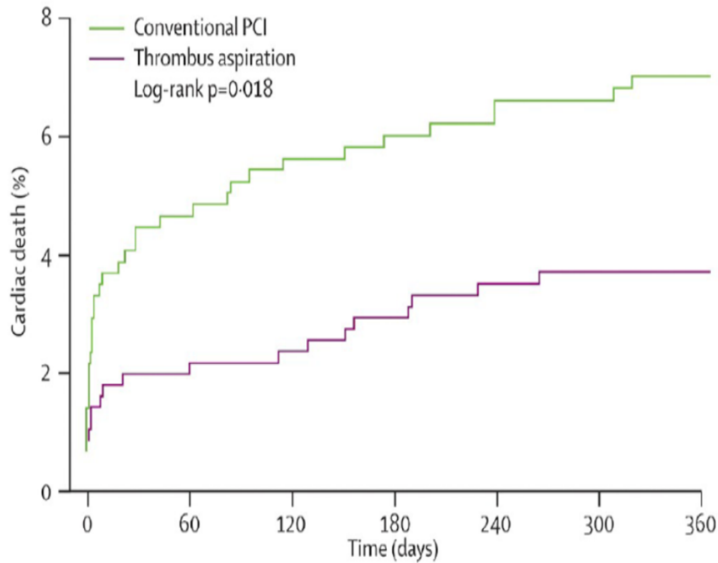


Microscopic distal emboli and no reflow



TIMI 3 flow with
absent
microvascular
perfusion

Large effect size in TAPAS (2008)

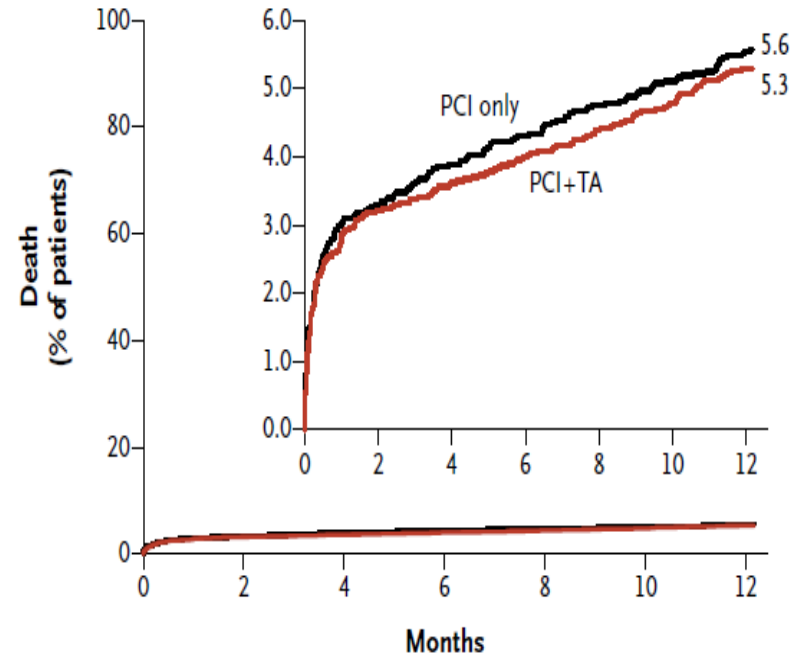


University Medical Center Groningen

Vlaar P, et al. TAPAS 1-year clinical outcome. Lancet 2008;371:1915-20



No difference in TASTE (2013) Cumulative Risk of Death



TAPAS trial (N=1071) showed a large benefit vs. TASTE (N=7244) showed no benefit of thrombus aspiration

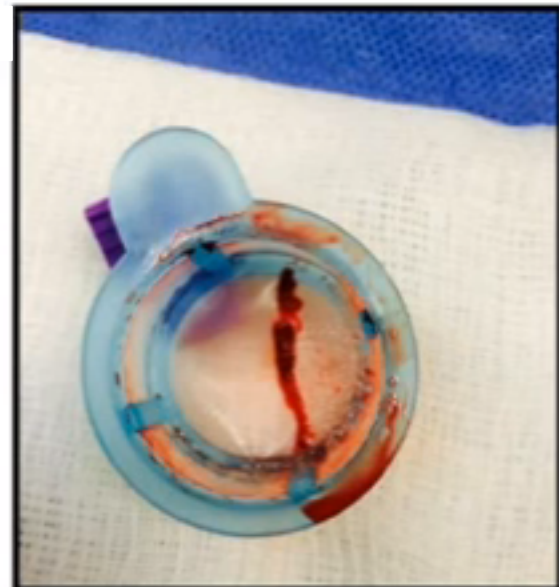
Vlaar PJ, et al. Lancet 2008;371:1915-20.
Frøbert O, et al. N Engl J Med 2013.
Lagerqvist B, et al. N Engl J Med. 2014.

Randomized trial of manual aspiration Thrombectomy + PCI vs. PCI Alone in STEMI (TOTAL)

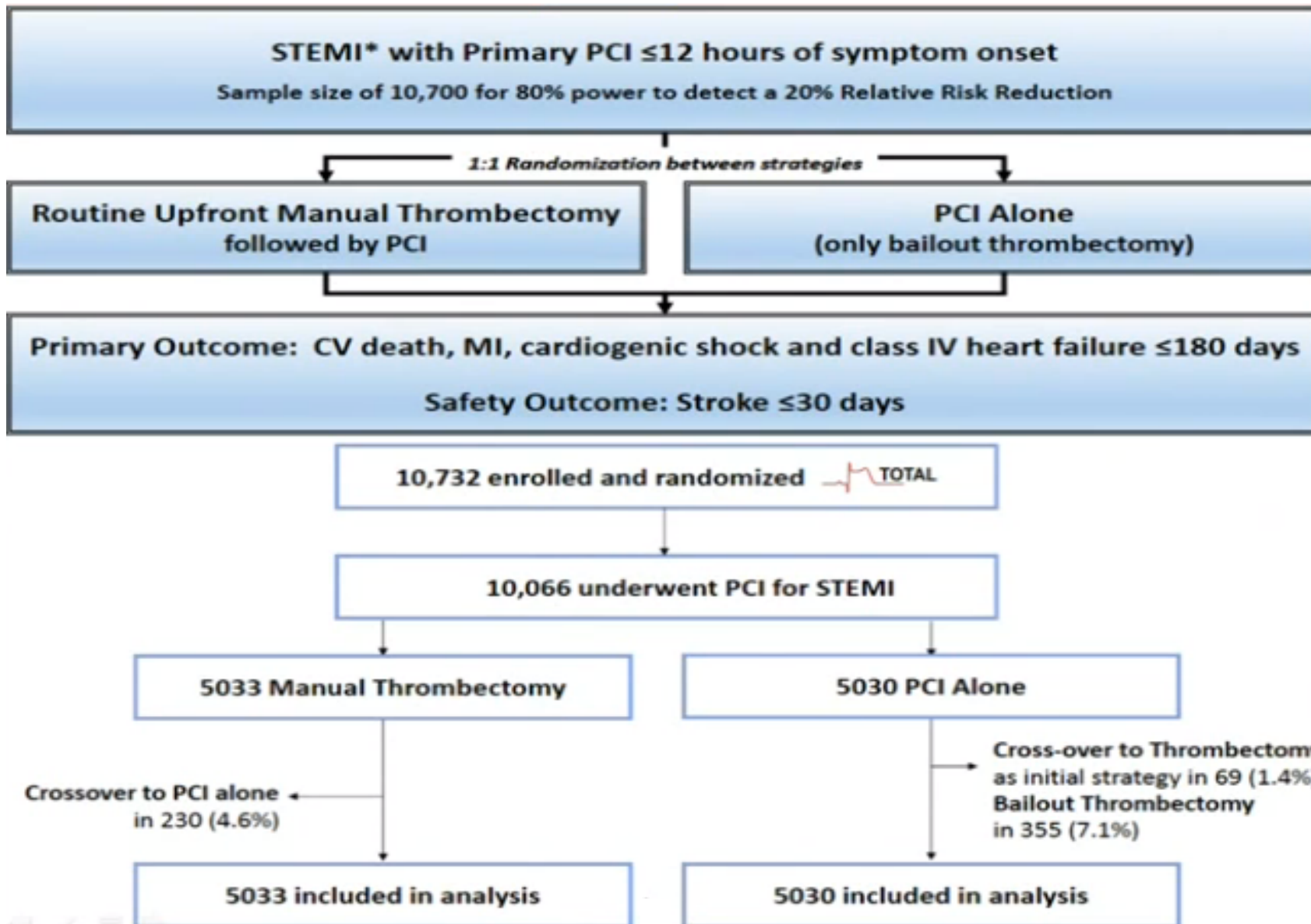
Rationale for Thrombectomy



Major Limitation of Primary PCI:
Distal Embolization and Reduced Flow

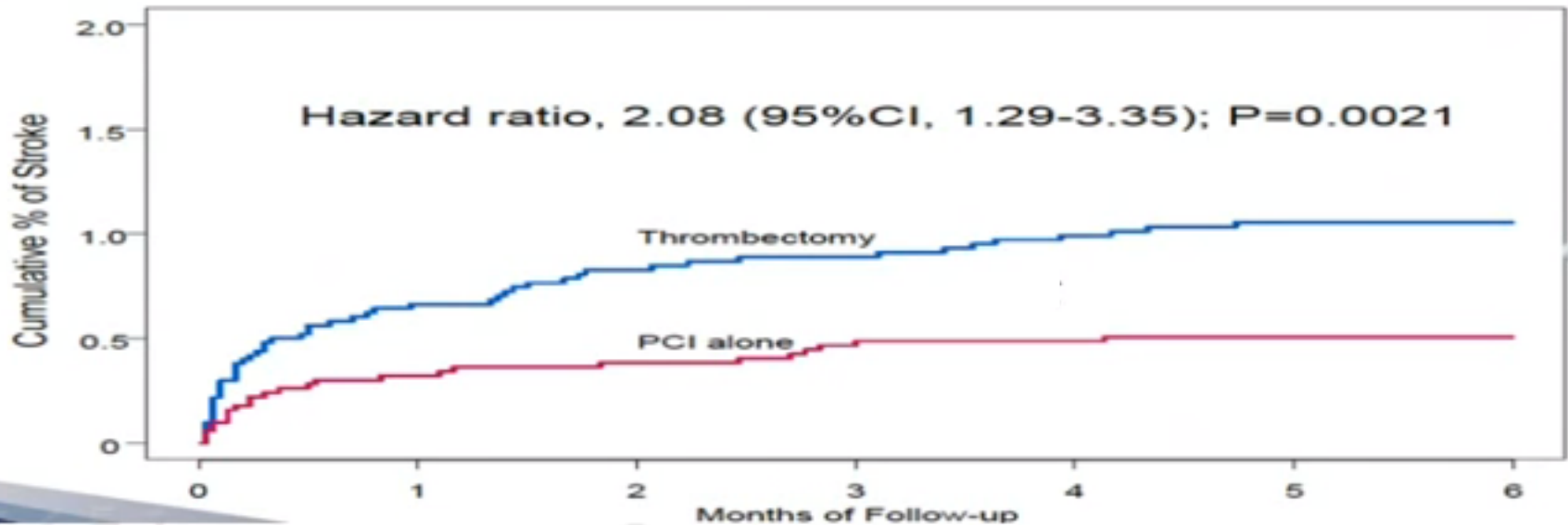


Hypothesis: Aspiration thrombectomy may
reduce embolization and
improve clinical outcomes



TOTAL

| Day 180 | Thrombectomy (N=5033) (%) | PCI alone (N=5030) (%) | HR | 95% CI | p |
|---|------------------------------|---------------------------|------|-----------|------|
| CV death, MI, shock or class IV heart failure | 347 (6.9%) | 351 (7.0%) | 0.99 | 0.85-1.15 | 0.86 |
| CV death | 157 (3.1%) | 174 (3.5%) | 0.90 | 0.73-1.12 | 0.34 |
| Recurrent MI | 99 (2.0%) | 92 (1.8%) | 1.07 | 0.81-1.43 | 0.62 |
| Cardiogenic Shock | 92 (1.8%) | 100 (2.0%) | 0.92 | 0.69-1.22 | 0.56 |
| Class IV heart failure | 98 (1.9%) | 90 (1.8%) | 1.09 | 0.82-1.45 | 0.57 |



Routine thrombectomy compared to PCI alone with only bailout thrombectomy did not reduce CV death, MI, shock or heart failure within 180 days

Routine thrombectomy was associated with increased risk of stroke within 30 days

TOTAL and TASTE emphasize the need to conduct large randomized trials of common interventions even when small trials appear positive

#PostACC15

ORIGINAL ARTICLE

Randomized Trial of Primary PCI with or without Routine Manual Thrombectomy

S.S. Jolly, J.A. Cairns, S. Yusuf, B. Meeks, J. Pogue, M.J. Rezkani, S. Kodov, L. Thabane, G. Stambouli, R. Moreno, A. Cardinale, S. Chowdhary, S. Lam, K. Narmoth, P.C. Steg, I. Bertoni, Y. Xia, W.J. Cantor, C.B. Overgaard, C.K. Naber, A.N. Chouhan, S.C. Walsh, D.F. Bertrand, A. Assoum, R. Shroff, S. Paruchy, S.V. Rao, M.K. Natarajan, J.M. ten Berg, O. Shatalovska, P. Cao, P. Widimsky, and V. Dzavik, for the TOTAL Investigators*

How effective is manual thrombus aspiration?



How effective is manual thrombus aspiration?

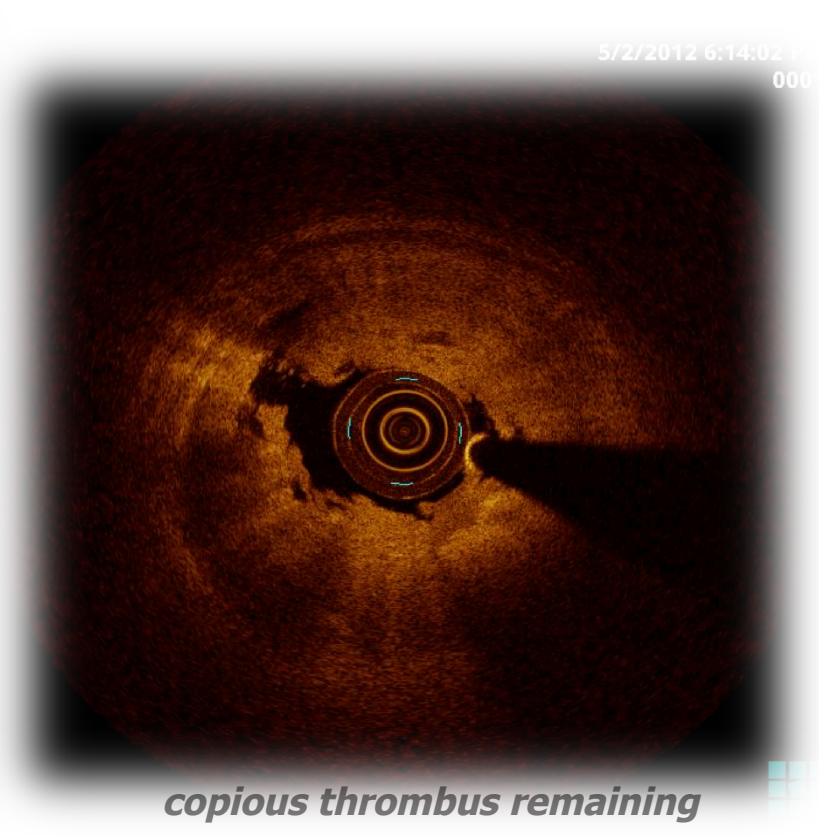
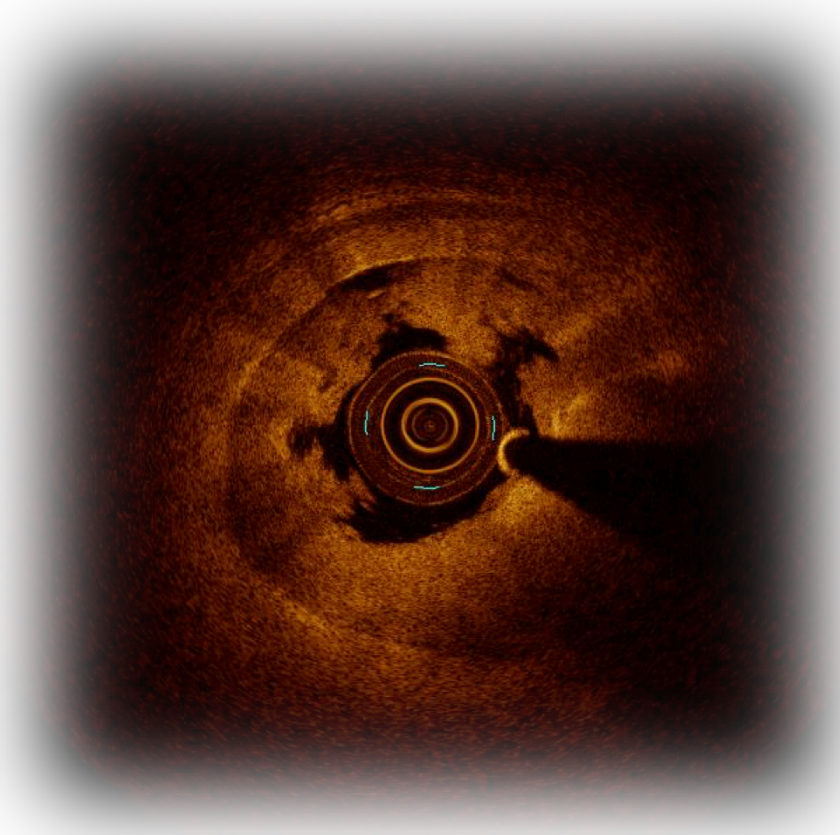
Post aspiration

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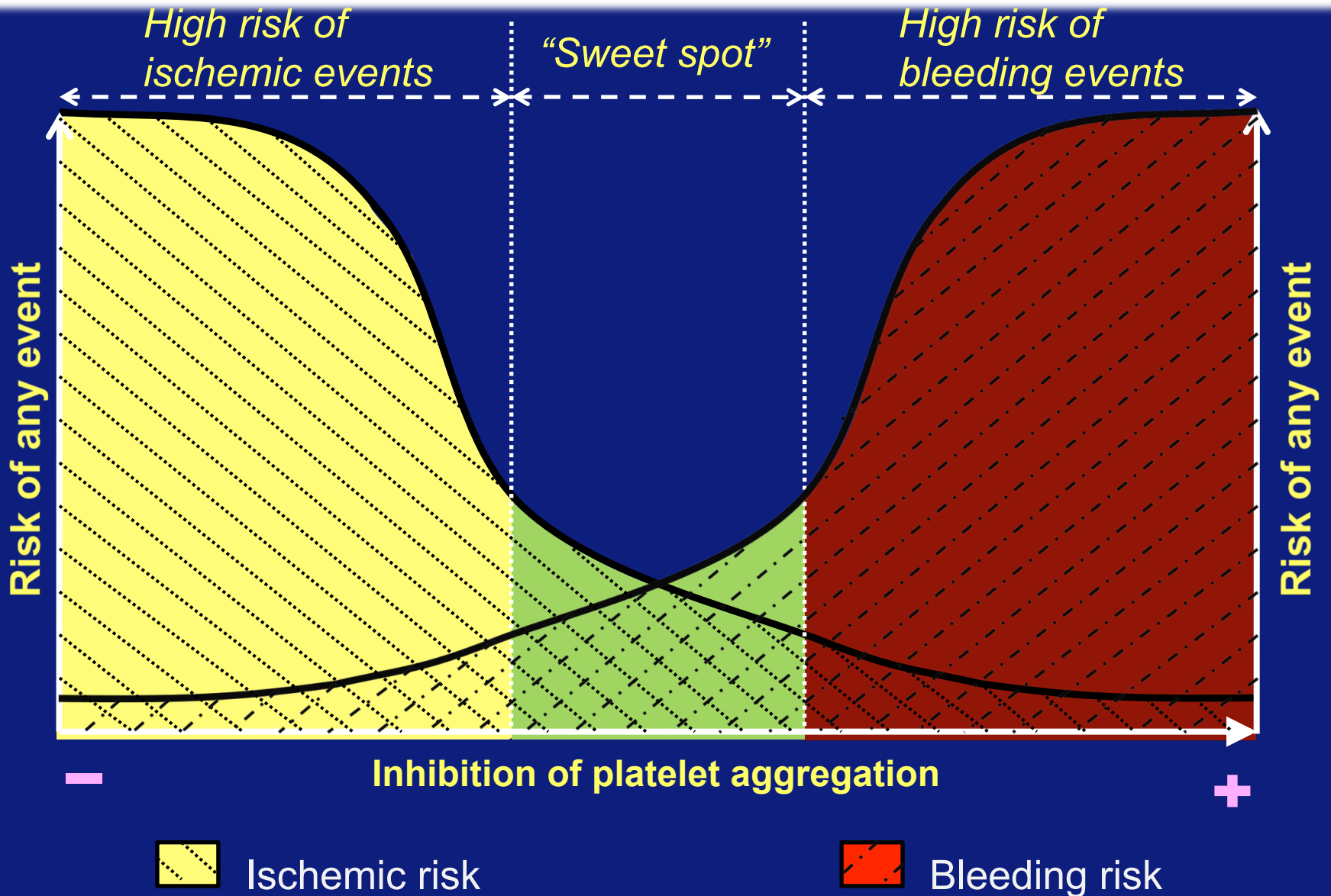


c/o Prof. Valdes, Murcia, Spain

Recommendations for antithrombotic treatment in patients with STEMI undergoing primary PCI

| Recommendations | Class ^a | Level ^b |
|--|--------------------|--------------------|
| Antiplatelet therapy | | |
| ASA is recommended for all patients without contraindications at an initial oral loading dose of 150–300 mg (or 80–150 mg i.v.) and at a maintenance dose of 75–100 mg daily long-term regardless of treatment strategy. | I | A |
| A P2Y ₁₂ inhibitor is recommended in addition to ASA and maintained over 12 months unless there are contraindications such as excessive risk of bleeding. Options are: | I | A |
| • Prasugrel (60 mg loading dose, 10 mg daily dose) if no contraindication | I | B |
| • Ticagrelor (180 mg loading dose, 90 mg twice daily) if no contraindication | I | B |
| • Clopidogrel (600 mg loading dose, 75 mg daily dose), only when prasugrel or ticagrelor are not available or are contraindicated. | I | B |
| It is recommended to give P2Y ₁₂ inhibitors at the time of first medical contact. | I | B |
| GP IIb/IIIa inhibitors should be considered for bail-out or evidence of no-reflow or a thrombotic complication. | IIa | C |
| Upstream use of a GP IIb/IIIa inhibitor (vs. in-lab use) may be considered in high-risk patients undergoing transfer for primary PCI. | IIb | B |
| Anticoagulants | | |
| Anticoagulation is recommended for all patients in addition to antiplatelet therapy during PCI. | I | A |
| The anticoagulation is selected according to both ischaemic and bleeding risks, and according to the efficacy–safety profile of the chosen agent. | I | C |
| Unfractionated heparin: 70–100 U/kg i.v. bolus when no GP IIb/IIIa inhibitor is planned; 50–70 U/kg i.v. bolus with GPIIb/IIIa inhibitor. | I | C |
| Bivalirudin 0.75 mg/kg i.v. bolus followed by i.v. infusion of 1.75 mg/kg/h for up to 4 hours after the procedure. | IIa | A |
| Enoxaparin i.v. 0.5 mg/kg with or without GP IIb/IIIa inhibitor. | IIa | B |

Balancing Safety and Efficacy



Posthospitalization Plan of Care

I IIa IIb III



Posthospital systems of care designed to prevent hospital readmissions should be used to facilitate the transition to effective, coordinated outpatient care for all patients with STEMI.

I IIa IIb III



Exercise-based cardiac rehabilitation/secondary prevention programs are recommended for patients with STEMI.



*Helping Cardiovascular Professionals
Learn. Advance. Heal.*



Posthospitalization Plan of Care

I IIa IIb III



A clear, detailed, and evidence-based plan of care that promotes medication adherence, timely follow-up with the healthcare team, appropriate dietary and physical activities, and compliance with interventions for secondary prevention should be provided to patients with STEMI.

I IIa IIb III



Encouragement and advice to stop smoking and to avoid secondhand smoke should be provided to patients with STEMI.



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Learn. Advance. Heal.*





ESQUEMA DE TRATAMIENTO ACTUAL MEDIANTE
REVASCULARIZACIÓN PRECOZ
EN EL
INFARTO AGUDO DE MIOCARDIO

CONCLUSIONES

ANGIOPLASTIA PRIMARIA

CREACIÓN DE REDES ASISTENCIALES

STENTS FARMACOACTIVOS DE SEGUNDA GENERACIÓN

ACCESO RADIAL

REVASCULARIZACIÓN COMPLETA

TROMBOASPIRACIÓN NO DE RUTINA

REHABILITACIÓN CARDÍACA Y PREVENCIÓN SECUNDARIA



muchas gracias