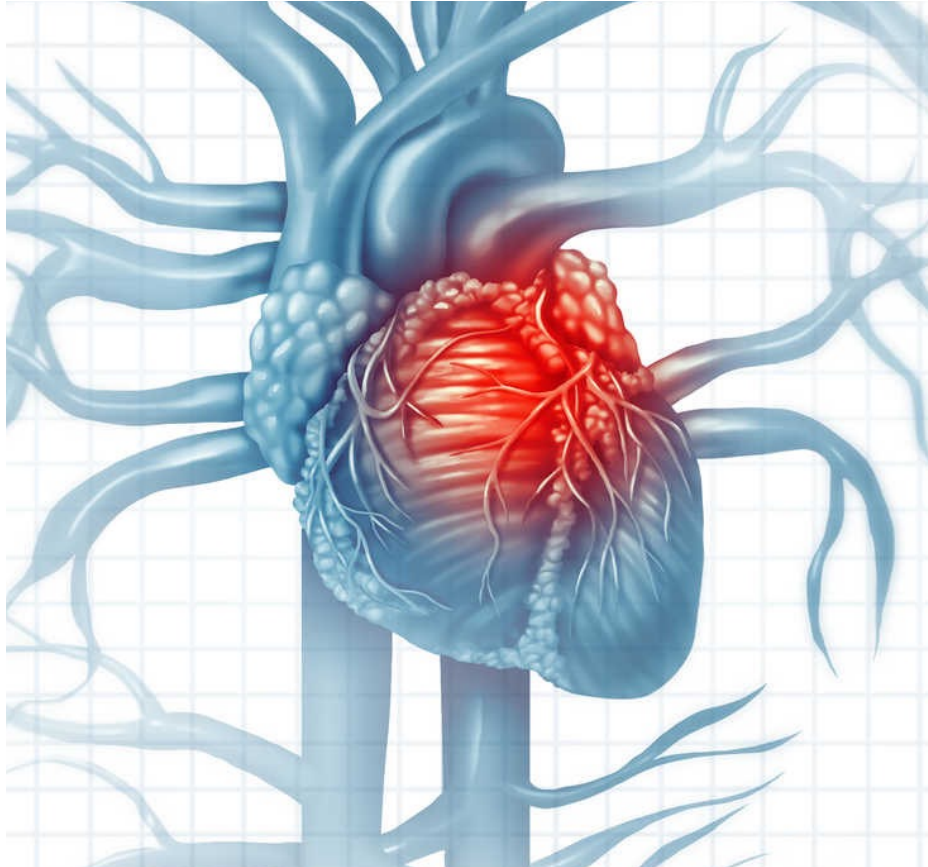


L'uso dei NAO nella cardiopatia ischemica



SERVIZIO SANITARIO REGIONALE
EMILIA-ROMAGNA
Azienda Unità Sanitaria Locale di Imola

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European Heart Journal
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ESC GUIDELINES

2016 ESC Guidelines for the management of atrial fibrillation developed in collaboration with EACTS

The Task Force for the management of atrial fibrillation of the European Society of Cardiology (ESC)

Developed with the special contribution of the European Heart Rhythm Association (EHRA)

Endorsed by the European Association of Cardio-Thoracic Surgeons (EACTS)



European Heart Journal (2016) 37, 1–64
European Society of Cardiology doi:10.1093/eurheartj/ehw126

SPECIAL ARTICLE

The 2018 European Heart Rhythm Association Practical Guide on the use of non-vitamin K

antagonist oral anticoagulants with a

Jan Steffel
Matthias Halperin
Holger Rottenburger
Roman Colan



European Heart Journal (2018) 39, 1–76
European Society of Cardiology doi:10.1093/eurheartj/ehy374

ESC/EACTS GUIDELINES



European Heart Journal (2018) 39, 1–193
European Society of Cardiology doi:10.1093/eurheartj/ehy174

EHRA CONSENSUS DOCUMENT



European Heart Journal (2018) 39, 1–193
European Society of Cardiology doi:10.1093/eurheartj/ehy174

2018 Joint European consensus document on

Circulation

WHITE PAPER

Antithrombotic Therapy in Patients With Atrial Fibrillation Undergoing Percutaneous Coronary Intervention: A North American

Circulation

ACC/AHA/HRS GUIDELINE

2019 AHA/ACC/HRS Focus Update on the Management of Atrial Fibrillation: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines

A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines



European Heart Journal (2020) 41, 1–126
European Society of Cardiology doi:10.1093/eurheartj/ehaa121

ESC GUIDELINES

2020 ESC Guidelines for the diagnosis and management of atrial fibrillation developed in collaboration with the European Association of Cardio-Thoracic Surgeons (EACTS)

The Task Force for the diagnosis and management of atrial fibrillation of the European Society of Cardiology (ESC)

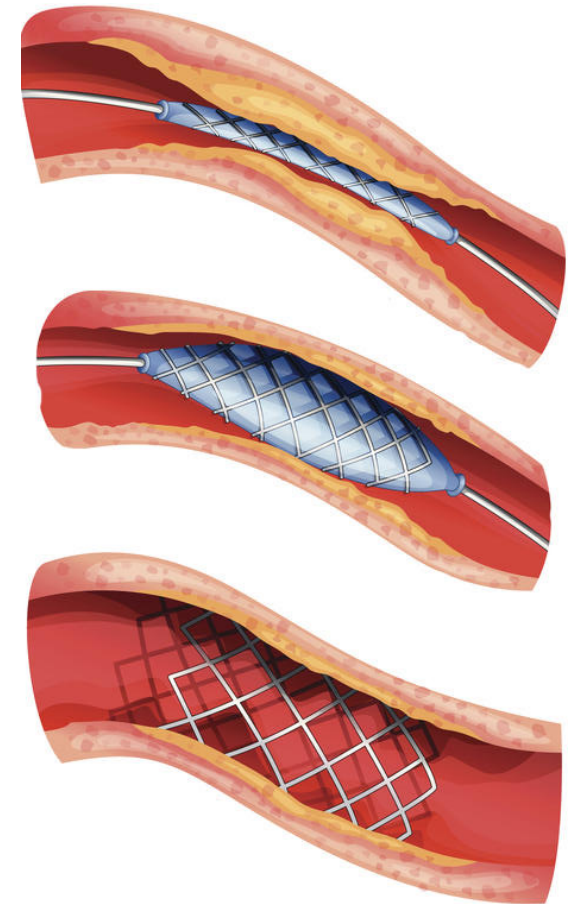
Developed with the special contribution of the European Heart Rhythm Association (EHRA) of the ESC

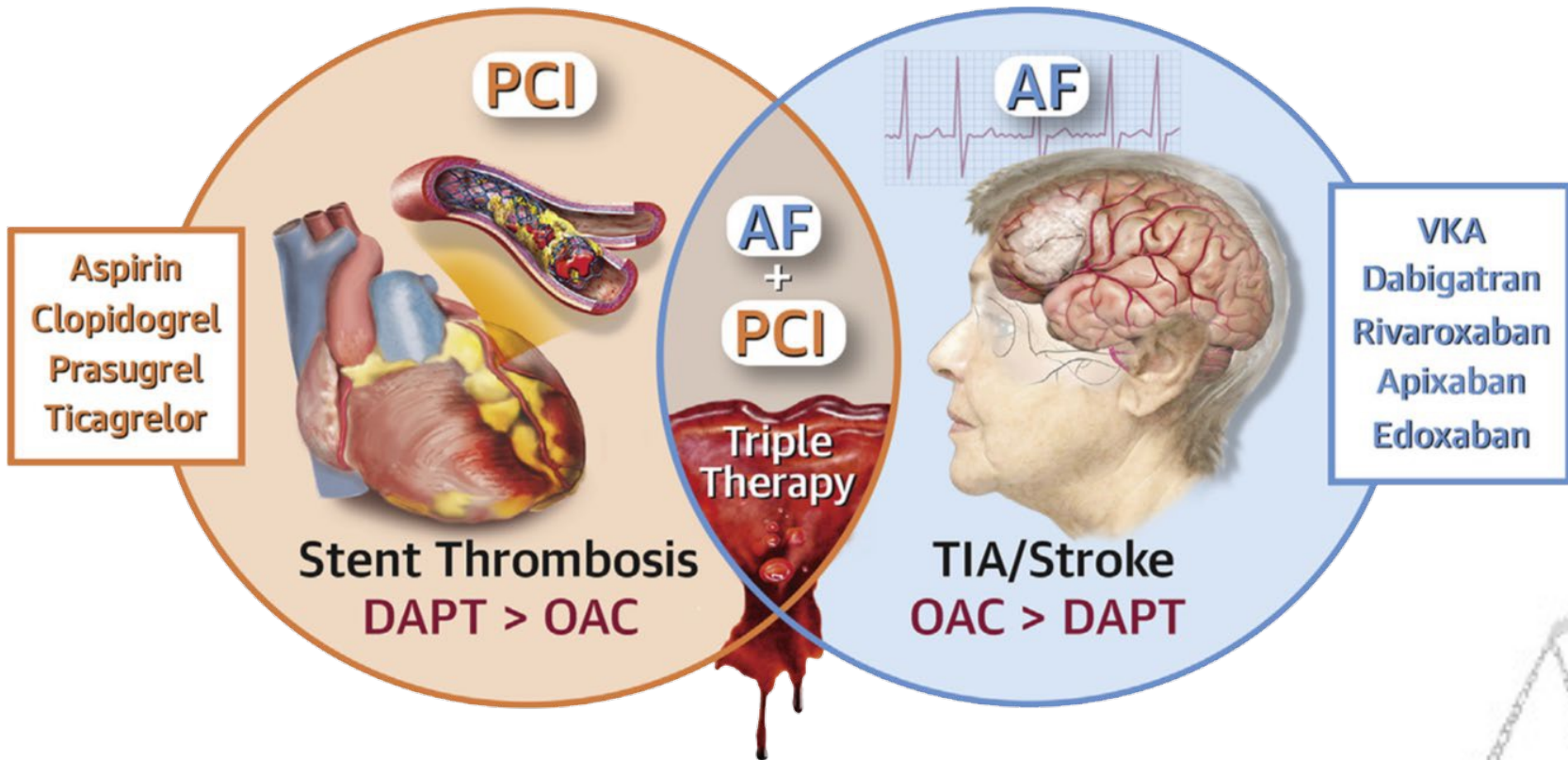


Doppia terapia antiaggregante per pazienti in anticoagulazione orale (triplice terapia)

Perché è utile

- Circa il 6–8% dei pz sottoposti a PCI hanno indicazione alla terapia anticoagulante a lungo termine per varie patologie quali FA, protesi valvolari meccaniche cardiache, tromboembolismo venoso
- Secondo le conoscenze attuali, in fase acuta la **DAPT è necessaria per prevenire la trombosi dello stent** ma non è sufficiente a prevenire lo stroke
- Viceversa, **la terapia anticoagulante orale è essenziale per la prevenzione dello stroke**, ma in fase acuta/subacuta non è idonea a prevenire nuovi eventi coronarici





Doppia terapia antiaggregante per pazienti in anticoagulazione orale (triplice terapia)

Quali rischi comporta

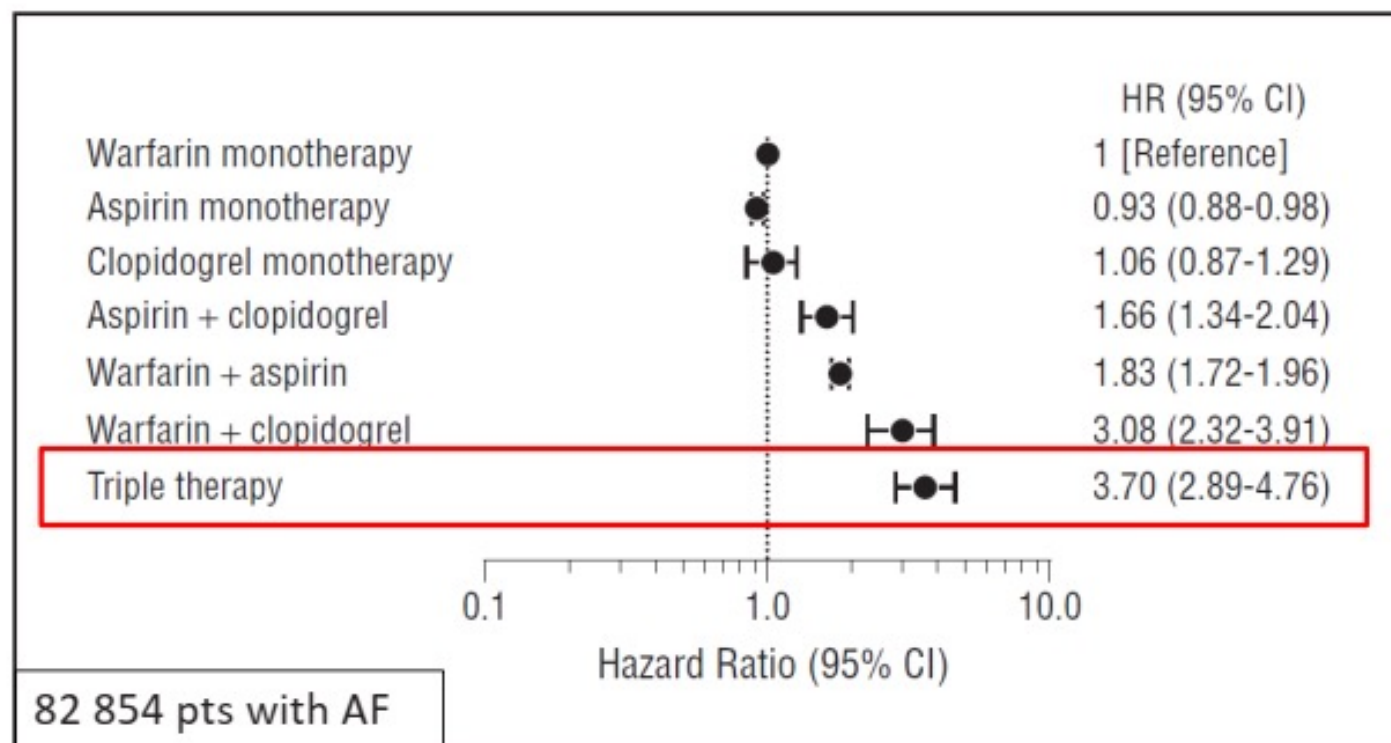
- Paragonata alla sola terapia anticoagulante orale, l'aggiunta della DAPT a questa **augmenta le complicanze emorragiche di circa 3 volte**
- Pertanto questi pz vanno considerati ad alto rischio di sanguinamento
- Le indicazioni alla terapia anticoagulante orale vanno attentamente rivalutate
- La triplice terapia va continuata solo fino a quando viene valutata assolutamente indispensabile



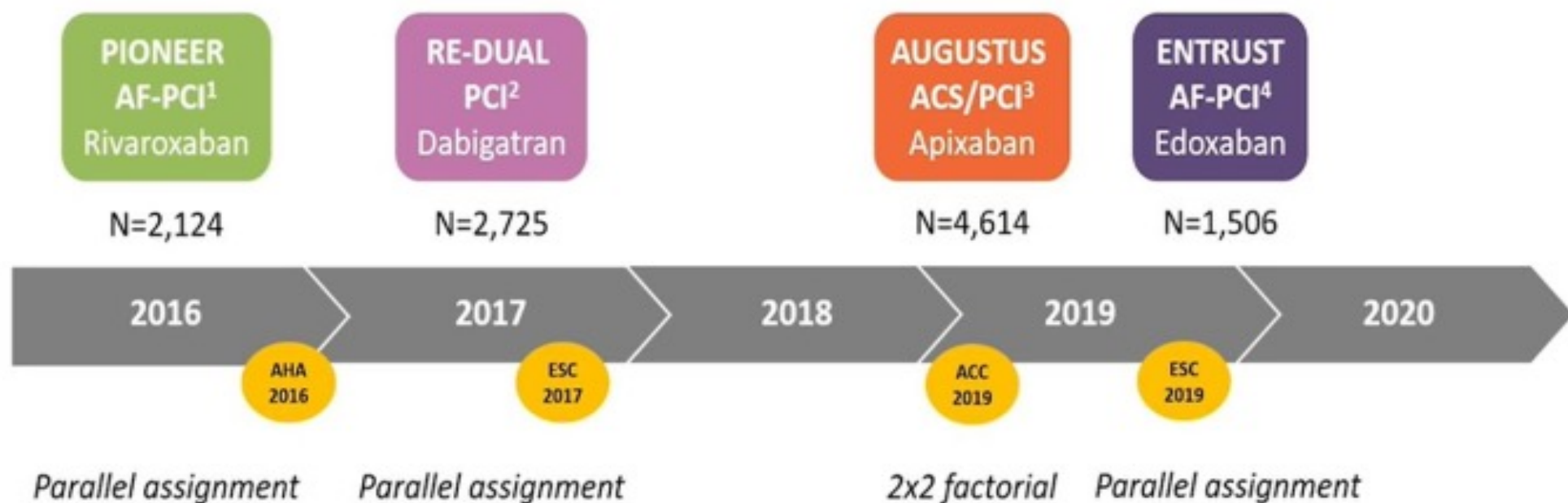
LESS IS MORE

Risk of Bleeding With Single, Dual, or Triple Therapy With Warfarin, Aspirin, and Clopidogrel in Patients With Atrial Fibrillation

Morten L. Hansen, MD, PhD; Rikke Sørensen, MD; Mette T. Clausen, MSc Pharm;
 Marie Louise Fog-Petersen, MSc Pharm; Jakob Raunso, MD; Niels Gadsbøll, MD, DMSc; Gunnar H. Gislason, MD, PhD;
 Fredrik Folke, MD; Søren S. Andersen, MD; Tina K. Schramm, MD; Steen Z. Abildstrøm, MD, PhD;
 Henrik E. Poulsen, MD, DMSc; Lars Køber, MD, DMSc; Christian Torp-Pedersen, MD, DMSc



NOACs nella FA + ACS/PCI



All 4 trials are powered for safety (bleeding) and underpowered for efficacy (ischaemic endpoints)

Sospensione della terapia antiaggregante

- I dati al riguardo non sono univoci
- Nei pz stabili e liberi da eventi, viene in genere consigliata la sospensione dei farmaci antiaggreganti a 1 anno dopo la procedura di stent
- Studi dimostrano che dopo tale periodo la terapia anticoagulante da sola è superiore all'aspirina, e che la TAO + aspirina non dà maggiore protezione ma è associata con un maggiore rischio emorragico

**2020 ESC Guidelines for the diagnosis
and management of atrial fibrillation
developed in collaboration with the
European Association for
Cardio-Thoracic Surgery (EACTS)**

What is new in the 2020 Guidelines? New recommendations (12)

Recommendations	Class
Recommendations for patients with AF and an ACS, PCI, or CCS	
<i>Recommendations for AF patients with ACS</i>	ACUTI
In AF patients with ACS undergoing an uncomplicated PCI, early cessation (≤ 1 week) of aspirin and continuation of dual therapy with an OAC and a P2Y ₁₂ inhibitor (preferably clopidogrel) for up to 12 months is recommended if the risk of stent thrombosis is low or if concerns about bleeding risk prevail over concerns about risk of stent thrombosis, irrespective of the type of stent used.	I
<i>Recommendations in AF patients with a CCS undergoing PCI</i>	CRONICI
After uncomplicated PCI, early cessation (≤ 1 week) of aspirin and continuation of dual therapy with OAC for up to 6 months and clopidogrel is recommended if the risk of stent thrombosis is low or if concerns about bleeding risk prevail over concerns about risk of stent thrombosis, irrespective of the type of stent used.	I

Figure 20 (1) Post-procedural management of patients with AF and ACS/PCI (full-outlined arrows represent a default strategy; graded/dashed arrows show treatment modifications depending on individual patient's ischaemic and bleeding risks)

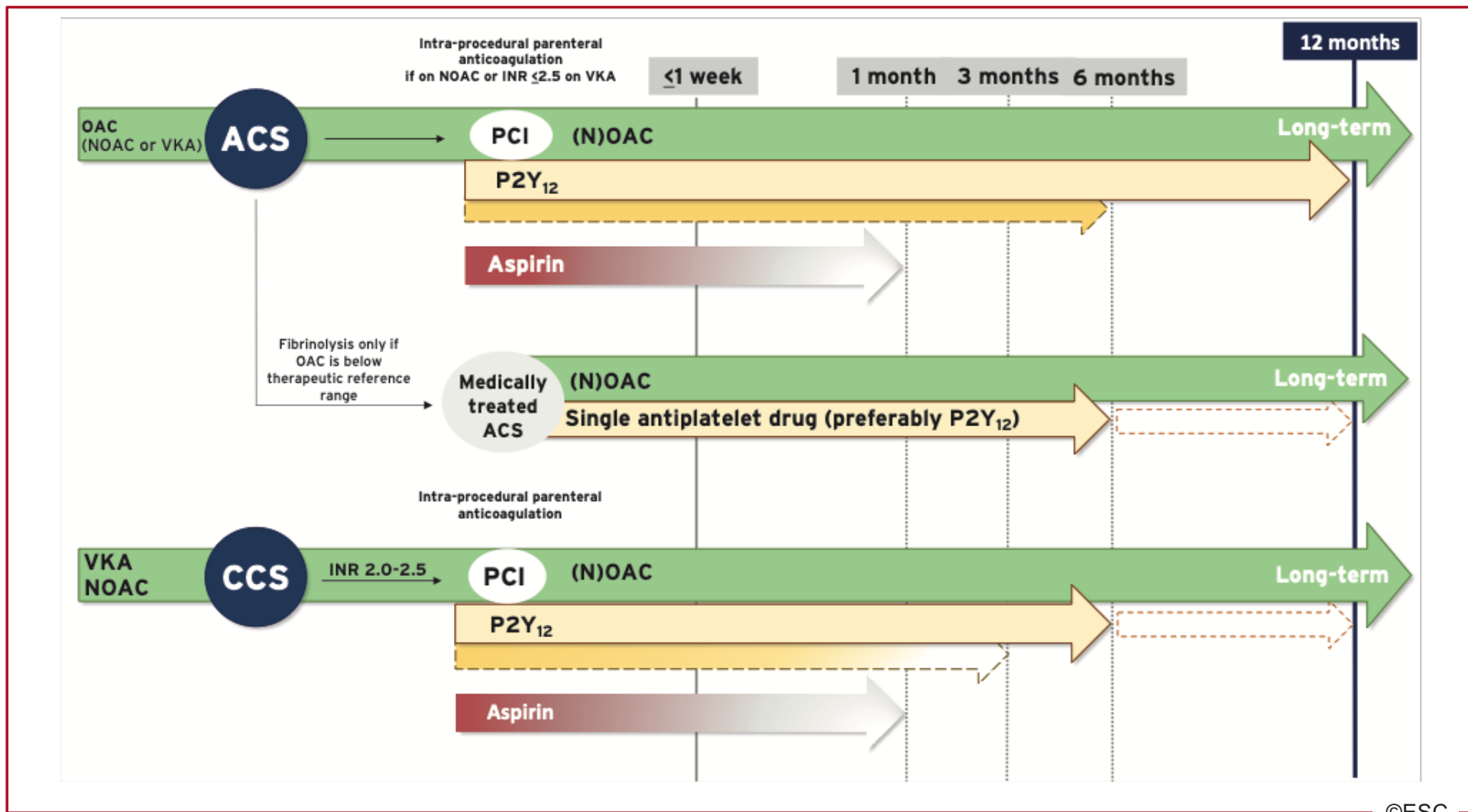


Figure 20 (2) Post-procedural management of patients with AF and ACS/PCI (full-outlined arrows represent a default strategy; graded/dashed arrows show treatment modifications depending on individual patient's ischaemic and bleeding risks)

THROMBOTIC RISK FACTORS

- Diabetes mellitus requiring therapy
- Prior ACS/recurrent myocardial infarction
- Multivessel CAD
- Concomitant PAD
- Premature CAD (occurring at age of <45 y) or accelerated CAD (new lesion within 2 years)
- CKD (eGFR <60 mL/min)
- Clinical presentation (ACS)
- Multivessel stenting
- Complex revascularisation (left main stenting, bifurcation lesion stenting, chronic total occlusion intervention, last patent vessel stenting)
- Prior stent thrombosis on antiplatelet treatment
- Procedural factors (stent expansion, residual dissection, stent length, etc.)

BLEEDING RISK FACTORS

- Hypertension
- Abnormal renal or liver function
- Stroke or ICH history
- Bleeding history or bleeding diathesis (e.g., anaemia with haemoglobin <110 g/L)
- Labile INR (if on VKA)
- Elderly (>65 years)
- Drugs (concomitant OAC and antiplatelet therapy, NSAIDs), excessive alcohol consumption

STRATEGIES TO REDUCE BLEEDING ASSOCIATED WITH PCI

- Radial artery access
- PPIs in patients taking DAPT who are at increased risk of bleeding (e.g., the elderly, dyspepsia, gastro-oesophageal reflux disease, Helicobacter pylori infection, chronic alcohol use)
- Non-administration of unfractionated heparin in patients on VKA with INR >2.5
- Pre-treatment with aspirin only, add a P2Y₁₂ inhibitor when coronary anatomy is known or if STEMI
- GP IIb/IIIa inhibitors only for bailout or periprocedural complications
- Shorter duration of combined antithrombotic therapy

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Recommendations for patients with AF and an ACS, PCI, or CCS (1)

Recommendations	Class	Level
General recommendations for patients with AF and an indication for concomitant antiplatelet therapy		
In AF patients eligible for NOACs, it is recommended to use a NOAC ^a in preference to a VKA in combination with antiplatelet therapy.	I	A
In patients at high bleeding risk (HAS-BLED ≥ 3), rivaroxaban 15 mg o.d. should be considered in preference to rivaroxaban 20 mg o.d. for the duration of concomitant single or DAPT, to mitigate bleeding risk.	IIa	B

^aSee summary of product characteristics for reduced doses or contraindications for each NOAC in patients with CKD, body weight <60 kg, age >75–80 years, and/or drug interactions.

Recommendations for patients with AF and an ACS, PCI, or CCS (2)

Recommendations	Class	Level
General recommendations for patients with AF and an indication for concomitant antiplatelet therapy (continued)		
In patients at high bleeding risk (HAS-BLED ≥ 3), dabigatran 110 mg b.i.d. should be considered in preference to dabigatran 150 mg b.i.d. for the duration of concomitant single or DAPT, to mitigate bleeding risk.	Ila	B
In AF patients with an indication for a VKA in combination with antiplatelet therapy, the VKA dosing should be carefully regulated with a target INR of 2.0–2.5 and TTR >70%.	Ila	B

Recommendations for patients with AF and an ACS, PCI, or CCS (3)

Recommendations	Class	Level
Recommendations for AF patients with ACS		
In AF patients with ACS undergoing an uncomplicated PCI, early cessation (≤ 1 week) of aspirin and continuation of dual therapy with an OAC and a P2Y ₁₂ inhibitor (preferably clopidogrel) for up to 12 months is recommended if the risk of stent thrombosis ^b is low or if concerns about bleeding risk ^c prevail over concerns about risk of stent thrombosis ^b , irrespective of the type of stent used.	I	A
Triple therapy with aspirin, clopidogrel, and an OAC ^d for longer than 1 week after an ACS should be considered when risk of stent thrombosis ^b outweighs the bleeding risk, ^c with the total duration (≤ 1 month) decided according to assessment of these risks, and the treatment plan should be clearly specified at hospital discharge.	IIa	C

^bRisk of stent thrombosis encompasses: (i) risk of thrombosis occurring and (ii) risk of death should stent thrombosis occur, both of which relate to anatomical, procedural, and clinical characteristics. Risk factors for CCS patients include stenting of left main stem or last remaining patent artery; suboptimal stent deployment; stent length >60 mm; diabetes mellitus; CKD; bifurcation with two stents implanted; treatment of chronic total occlusion; and previous stent thrombosis on adequate antithrombotic therapy. ^cBleeding risk in AF patients may be assessed using the HAS-BLED score, which draws attention to modifiable bleeding risk factors; those at high risk (score ≥ 3) can have more frequent or early review and follow-up. Bleeding risk is highly dynamic and does not remain static, and relying on modifiable bleeding risk factors alone is an inferior strategy to evaluate bleeding risk. ^dWhen dabigatran is used in triple therapy, dabigatran 110 mg b.i.d. may be used instead of 150 mg b.i.d., but the evidence is insufficient.

Egregio dott.ssa
Medico Curante _____ di anni 76 ,
Relazione clinica di ricovero dal 21/03/2023 al 29/03/2023

Diagnosi di Dimissione

RECENTE INFARTO MIOCARDICO ACUTO (STEMI) A SEDE INFERIORE SOTTOPOSTO AD ANGIOPLASTICA PRIMARIA SULLA CORONARIA DESTRA. FIBRILLAZIONE ATRIALE PAROSSISTICA RECIDIVANTE. LIEVE DISFUNZIONE VENTRICOLARE SINISTRA.

Da segnalare durante la degenza

- Persistente stabilità emodinamica e buon compenso circolatorio. I valori di troponina risultavano in progressivo calo rispetto alle precedenti misurazioni.
- In relazione agli episodi recidivanti di fibrillazione atriale parossistica ed all'elevato rischio tromboembolico (CHA2-DS2-VASc = 4), è stata impostata terapia anticoagulante orale con apixaban e contestualmente eseguito shift antiaggregante da Ticagrelor a Clopidogrel.

Conclusioni

Paziente affetto da cardiopatia ischemica post-infartuale (recente STEMI inferiore) con lieve disfunzione ventricolare sinistra ed episodi di fibrillazione atriale parossistica. Impostata terapia antitrombotica con Apixaban e Clopidogrel. Il Clopidogrel andrà proseguito per almeno 6 mesi ed auspicabilmente per 12 mesi, ma tale scelta terapeutica dovrà essere valutata nel follow-up, anche in base ad eventuali eventi emorragici ed all'andamento dei controlli dell'esame emocromocitometrico.

Programma Terapeutico

- CLOPIDOGREL 75 MGR (OS) 1 Cp ore 08:00 (da proseguire per almeno 6 mesi)
- ELIQUIS 5MG (OS) 1 Cpr ore 08:00 + 1 Cpr ore 20:00
- CORDARONE 200 MGR (OS) 1 Cpr ore 13:00 (da Lunedì a Venerdì)
- BISOPROLOLOLO 2,5MG (OS) 1 Cp ore 08:00 + 1 Cp ore 20:00

Conclusioni

- ✓ Non sono infrequenti pazienti con indicazione a terapia anticoagulante orale e che richiedano una duplice terapia antiaggregante per PCI in elezione o ACS, per ridurre occlusione dello stent e reinfarti
- ✓ L'impiego dei NOACs è preferibile al warfarin per il favorevole profilo di sicurezza ed efficacia
- ✓ Sono stati pubblicati vari lavori che impiegano solo una duplice terapia antitrombotica - NOACs + inibitore recettori piastrinici P2Y₁₂ (clopidogrel) - subito dopo la procedura o l'evento ischemico cardiaco
- ✓ Le recentissime LG ESC sulla FA hanno cambiato le indicazioni, riducendo il tempo di triplice terapia a 1 settimana e in certi casi fino al massimo di 1 mese, per continuare poi con la duplice terapia fino a 1 anno
- ✓ Oltre l'anno nella maggioranza dei casi è indicato l'uso del solo anticoagulante orale

Grazie per l'attenzione

