

Which Cardiac Patients Require Antiplatelet(s) in Addition to Anticoagulants?

Starting anticoagulation in patients already prescribed single or dual antiplatelet therapy

In many cases when a patient is started on an anticoagulant, existing antiplatelet treatment can be stopped or reduced in intensity. This should always be reviewed at the start of treatment and a plan clearly documented.

- If a patient has had previous cardiac surgery e.g. coronary artery bypass graft (CABG) surgery or valve replacement/repair: **check with the cardio-thoracic team before stopping any treatment.**

- If the patient has ever had percutaneous coronary intervention (PCI) or cardiac stenting: **check with the cardiology consultant or SpR before stopping any treatment.**

Single antiplatelet and anticoagulation

If there is an indication for single antiplatelet therapy and anticoagulation, always document the reason and the intended duration of combined therapy.

Dual anti-platelet and anticoagulation ('triple therapy')

There is limited data on the safety of combining dual antiplatelet treatment with anticoagulation. However, there is a significantly increased bleeding risk and the choice of regime will depend on a balance of risks (major bleeding) and benefits (reduction in risk of stroke, CV events and stent thrombosis) for the individual patient. The most **common** scenarios are likely to be:

- Patients with new atrial fibrillation (AF) and a CHA₂DS₂VASc ≥2 who have had emergency or elective PCI.

- Patients with pre-existing prescription for anticoagulation (e.g AF,VTE) who have had emergency or elective PCI.

Version	Lead Authors	MGG Approval
1	Alison Warren, Consultant Pharmacist Cardiology Dr James Cockburn Consultant Cardiologist	September 2017
2	Updated by Lizzie Malcolm, Specialist Rotational Pharmacist	November 2022
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General Rules:

It is not possible to cover every scenario but the following apply if the patient is prescribed **triple therapy**:

- The decision for triple therapy should **always** be made by a cardiology consultant or cardiology SpR.
- Treatment plans should be **clearly documented** - which includes in the hospital discharge summary (TTO) or any clinical letters to primary care.
- Patients with AF and treated with coronary revascularisation(CABG) surgery, triple therapy should be **avoided**. In very rare circumstances, the risk of thromboembolism may be higher than the risk of bleeding post surgery and therefore, whenever this situation arises the decision for triple therapy **must** be made by the consultant and involve the patient throughout the process.
- As part of triple therapy, the dual antiplatelet regime is usually **aspirin + clopidogrel**. Due to a higher risk of bleeding, prasugrel or ticagrelor should be avoided unless there is a clear need for these agents (e.g. stent thrombosis on aspirin plus clopidogrel, or major and unavoidable drug interactions).
- The **duration** of triple therapy should **always** be stated by the cardiology consultant/SpR and should be kept **as short as possible**, weighing up the risk of thrombotic events and bleeding (using HAS-BLED or ORBIT scores and addressing modifiable bleeding risk factors).
- For triple therapy after uncomplicated PCI in patients with AF, **early cessation of aspirin after 1 week** may be considered if the risk of stent thrombosis is low or if bleeding risk is high (HAS-BLED ≥ 3). In most scenarios (including after ACS) aspirin can be stopped after **1 month** with the patient continuing on **clopidogrel plus anticoagulation**.
- If there is an on-going need for anticoagulation, clopidogrel should be **stopped after 12 months** (and anticoagulant continued) unless deemed to be high thrombotic risk by the cardiology consultant. **If any uncertainty discuss with cardiology consultant before stopping the antiplatelet.**

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Choice of anticoagulant in ‘triple therapy’:

- In eligible patients with AF, a **DOAC should be used in preference to warfarin** (as studies have shown a higher bleeding risk with warfarin compared to DOACs when used in triple therapy). **There is currently no evidence to suggest using one DOAC over another, however apixaban or rivaroxaban may be preferable based on clinician experience.** If there any doubts, consult the cardiology consultant.

- The **normal dose of DOAC** should be prescribed with the licensed dose for the indication and any criteria for dose reduction. However for AF patients at higher bleeding risk (HASBLED \geq 3 or ORBIT \geq 4) a lower dose of rivaroxaban (15mg once a day) may be preferred regardless of the renal function. The lower doses will need to be reviewed when the patient is changed to single antiplatelet + anticoagulant (and may need to be increased in line with dosing recommendations).

- If the anticoagulant is **warfarin** (e.g. for other high risk indications i.e. valve replacements or LV thrombus), close monitoring of the INR is required and the **INR target** must be clearly documented. Use of DOACs for LV thrombus is increasing and remains **unlicensed**, therefore is a decision to be made by the cardiology consultant.

- For patients with AF already on warfarin, efforts should be made to switch to a DOAC if appropriate. A target INR of 2-2.5 may be preferred rather than the usual 2-3.

- **If the plan is unclear always check with the cardiology consultant /SpR**

Gastro-protection

For patients without a pre-existing prescription, a prophylactic dose of proton pump inhibitor (e.g. lansoprazole 15mg once a day) should be offered for gastro-protection to all patients on combination therapy with single or dual antiplatelet and an anticoagulant. Avoid omeprazole and esomeprazole as they may reduce the antiplatelet effect of clopidogrel.

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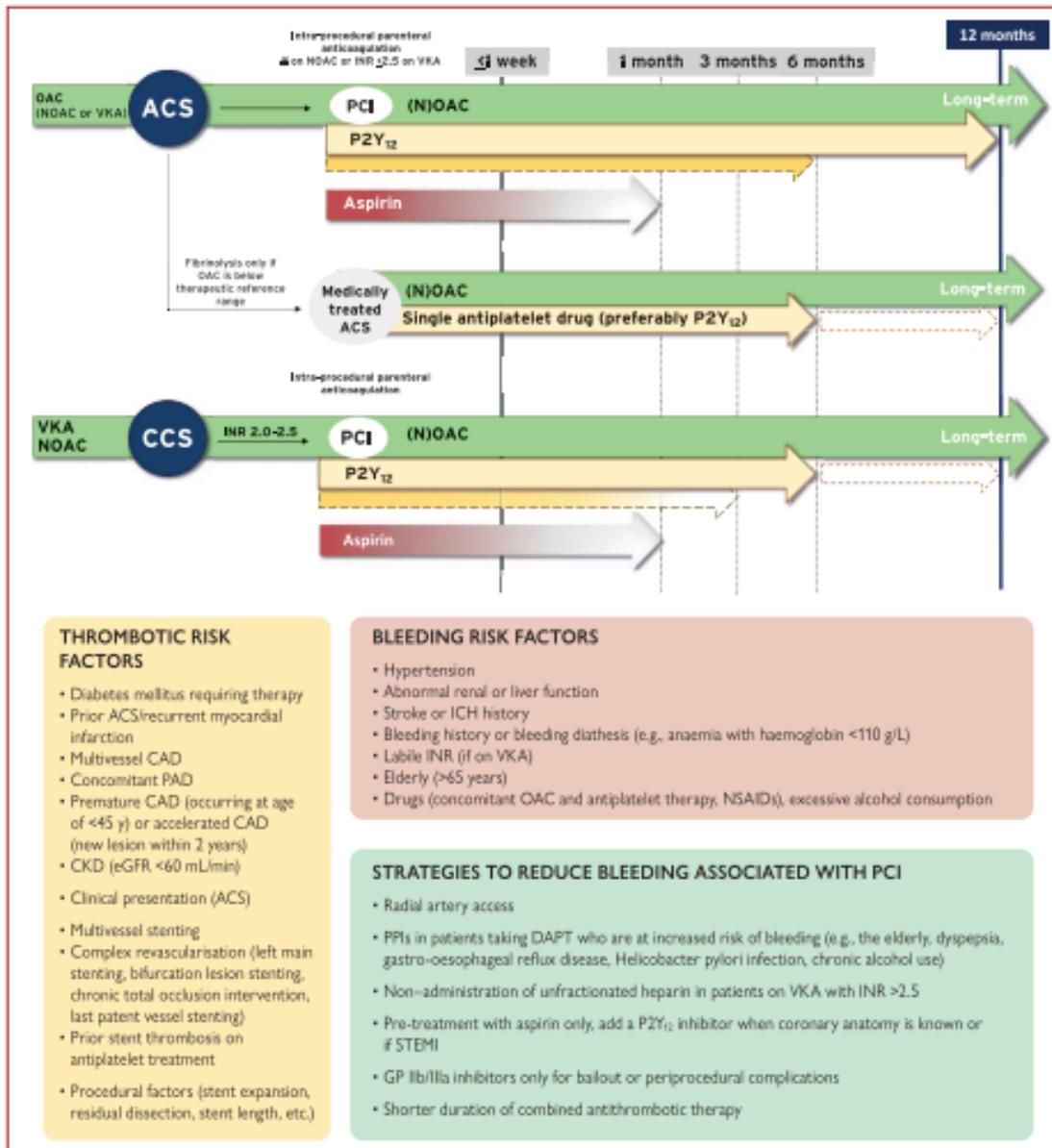


Figure 20 Post-procedural management of patients with AF and ACS/PCI (full-outlined arrows represent a default strategy; graded/dashed arrows show treatment modifications depending on individual patient's ischaemic and bleeding risks).

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