

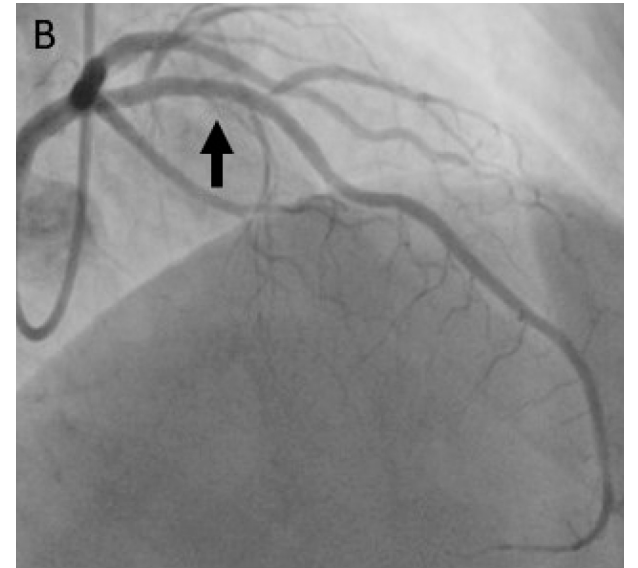
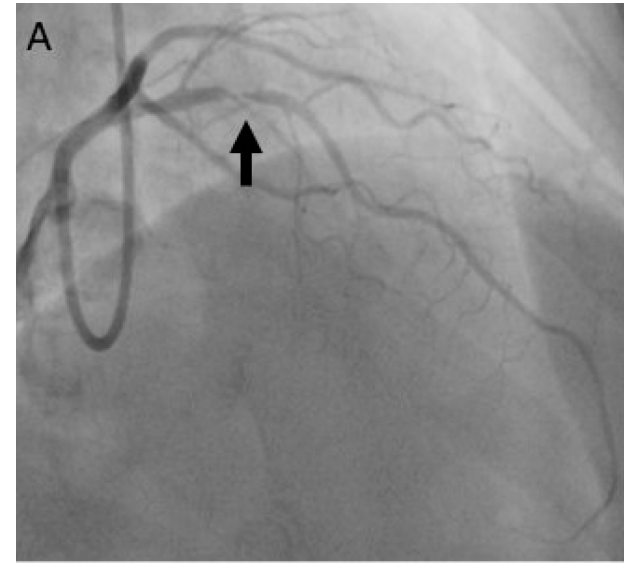
Anticoagulation soup – demystifying anticoagulation in patients with CAD +/- AF

Drs Jith Somaratne and Andrew Martin

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Case 1

- 76yo Māori woman with a history of AF
- Background of coronary heart disease, hypertension, and dyslipidaemia
- On dabigatran 110mg bd, bisoprolol 10mg od, ISMN 120 mg od, candesartan 8mg od, and rosuvastatin 20mg od
- PCI to LAD six weeks ago for ongoing angina despite medical therapy
- Bisoprolol and isosorbide mononitrate discontinued
- Commenced on aspirin and clopidogrel



Time from PCI	Default strategy	Patients at high ischemic/thrombotic and low bleeding risk	Patients at low ischemic/thrombotic or high bleeding risk
Peri-PCI	Triple Therapy (OAC + DAPT)	Triple Therapy (OAC + DAPT)	Triple Therapy (OAC + DAPT)
1 month	Double Therapy up to 12 months (OAC + P2Y ₁₂ inhibitor)	Triple Therapy up to 1 month (OAC + DAPT)	Double Therapy up to 6 months (OAC + P2Y ₁₂ inhibitor)
3 months		Double Therapy up to 12 months (OAC + P2Y ₁₂ inhibitor)	
6 months			OAC alone
12 months			
>12 months	OAC alone	OAC alone	OAC alone

Angiolillo DJ, et al. Circulation
2021;143:583–596

Case 2

- 78 year old woman
- Background of hypertension
- Usual medications: Felodipine ER 5mg daily, Candesartan 16mg daily
- BP 132/75, HR 85bpm, euvolaemic
- ECG: AF, normal QRS duration and axis
- Echo: Normal ventricular size/function, no valvular heart disease, moderate bi-atrial enlargement
- Bloods: HB 130, Creatinine 135 (eGFR 32), HBA1c 42, Urine ACR 2, normal LFTs, normal TSH

	Dabigatran	Apixaban	Edoxaban	Rivaroxaban
Dosing recommendation	CrCl \geq 50 mL/min: 150 mg BID)	SCr \geq 1.5 mg/dL: 5 mg BID)	Not available	CrCl \geq 50 mL/min: 20 mg QD)
Dosing if CKD	When CrCl 30-49 mL/min, 150 mg BID is possible (SmPC) but 110 mg BID if “high risk of bleeding”	CrCL 15-29 mL/min: 2.5 mg BID SCr > 1.5 mg/dL in combination with age \geq 80 years or weight \leq 60 kg (SmPC) or with other cautionary factors: 2.5 mg BID	Not available	15 mg QD when CrCl 15-49 mL/min

Heidbuchel H, et al. *Eur Heart J*. 2013;34:2094-2106.

Table 3: Oral anticoagulation in stroke patients with atrial fibrillation for all patients and for Māori and by non-urban hospitals.

	All	Māori	Non-Māori	P-value	Urban	Non-Urban	P-value
Overall AF prevalence*	807/2,352 (34.9)	107/273 (39.2)	700/2,079 (33.7)	0.07	478/1,420 (33.7)	329/932 (35.3)	0.42
Known AF on anticoagulant	442/666 (66.3)	66/88 (75.0)	376/578 (65.0)	0.06	251/388 (64.7)	191/278 (68.7)	0.28
New AF diagnosis at time of stroke (not on anticoagulant at time of stroke)	141/807 (17.5)	19/107 (17.8)	122/700 (17.4)	0.92	90/478 (18.8)	51/329 (15.5)	0.23
Known AF but not on anticoagulant	224/666 (33.6)	22/88 (25.0)	202/578 (34.9)	0.07	137/388 (35.3)	87/278 (31.3)	0.28
Reason for no anticoagulation							
Falls	14/205 (6.8)	2/20 (10)	12/185 (6.5)	0.56	9/123 (7.3)	5/82 (6.1)	0.74
ICH, GI bleed, other bleed	42/205 (20.5)	3/20 (15)	39/185 (21.1)	0.52	26/123 (21.1)	16/82 (19.5)	0.78
Frailty, comorbidities, side effects	27/205 (13.2)	1/20 (5)	26/185 (14.1)	0.25	12/123 (9.8)	15/82 (18.3)	0.79
Pre-/peri-procedure	7/205 (3.4)	1/20 (5)	6/185 (3.2)	0.67	6/123 (4.9)	1/82 (1.2)	0.15
Patient preference/non-compliant	37/205 (18.1)	5/20 (25)	32/185 (17.3)	0.40	22/123 (17.0)	15/82 (18.3)	0.81
Stopped for procedure and never restarted	7/205 (3.4)	1/20 (5)	6/185 (3.2)	0.67	3/123 (2.4)	4/82 (4.9)	0.33
AF duration felt not significant	5/205 (2.4)	0/20 (0)	5/185 (2.7)	0.46	4/123 (3.3)	1/82 (1.2)	0.34
Unknown	66/205 (32.2)	7/20 (35)	59/185 (31.9)	0.78	41/123 (33.3)	25/82 (20.5)	0.05

Data are number of patients (% of group)

*Missing values/unknown: n=63

#Missing values=19