

Anticoagulation in Patients with Acute Myocardial Infarction Who Are Taking Dual Antiplatelet Agents

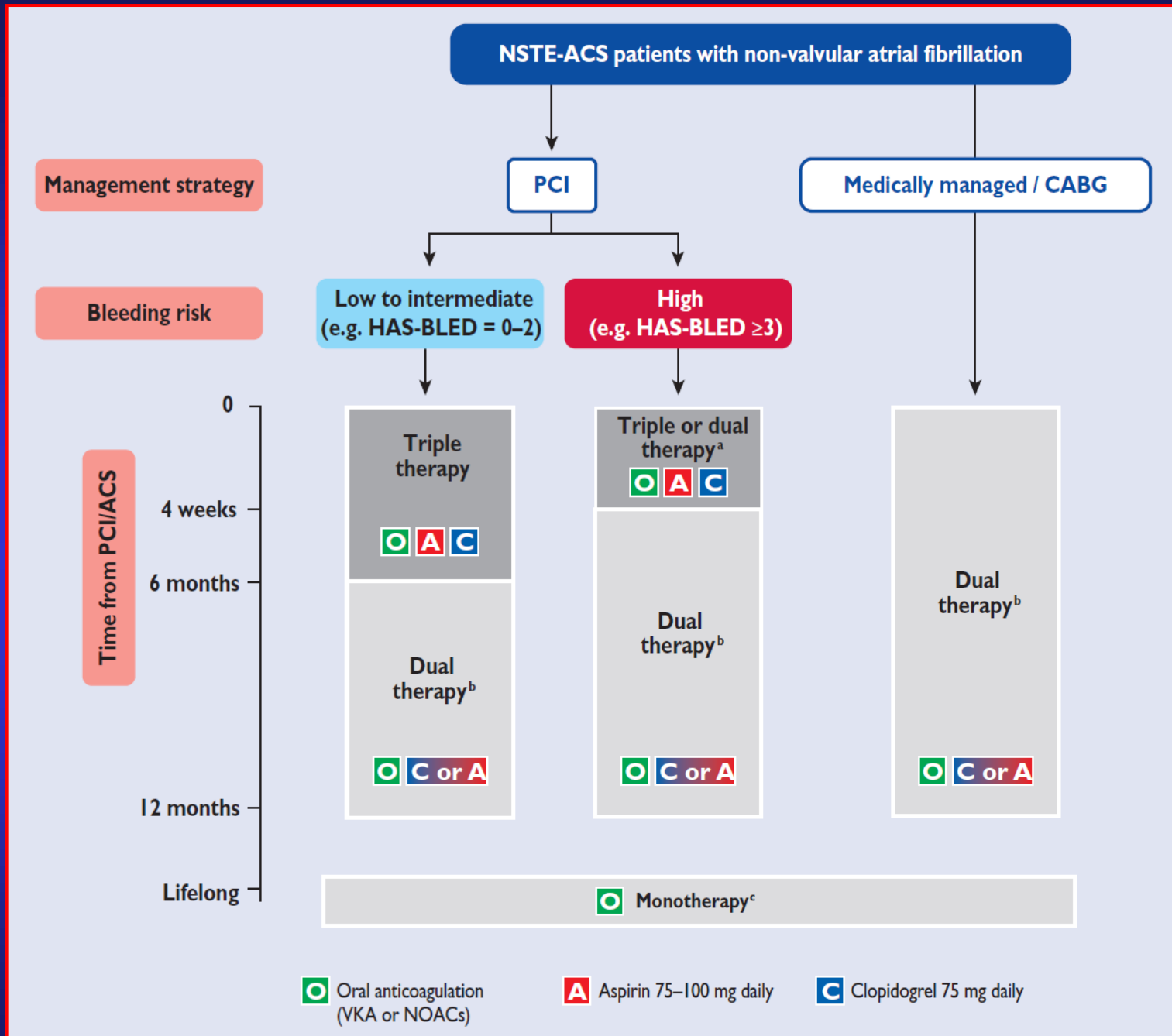
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Background

- Patients with atrial fibrillation and high thromboembolic risk need oral anticoagulant (OAC) to prevent the embolic events.
- OAC increases the bleeding risk in patients with acute myocardial infarction who are taking dual antiplatelet agents.

Recommendations	Class	Level
In patients with a firm indication for OAC (e.g. atrial fibrillation with a CHA2DS2-VASc score ≥ 2 , recent venous thromboembolism, LV thrombus or mechanical valve prosthesis), OAC is recommended in addition to antiplatelet therapy.	I	C
If at low bleeding risk (HAS-BLED ≤ 2), triple therapy with OAC, aspirin and clopidogrel should be considered for 6 months, followed by OAC and aspirin or clopidogrel continued up to 12 months.	IIa	C
If at high bleeding risk (HAS-BLED ≥ 3), triple therapy with OAC, aspirin and clopidogrel should be considered for a duration of 1 month, followed by OAC and aspirin or clopidogrel continued up to 12 months irrespective of the stent type (BMS or new-generation DES).	IIa	C



Methods

- KAMIR-NIH Registry; 13,031 patients with AMI
- Atrial fibrillation on the initial ECG and finished 1-year follow-up visit; 322 patients
 - Group 1; 73 patients with OAC
 - Group 2; 249 patients without OAC

Results

Baseline Characteristics of Patients

	Group 1 (N = 73)	Group 2 (N = 249)	p value
Age (years)	70.0 ± 10.1	67.0 ± 12.3	0.033
Male	40 (54.8%)	183 (73.5%)	0.004
Smoker	11 (15.1%)	92 (36.9%)	0.001
Hypertension	48 (65.8%)	144 (57.8%)	0.465
Diabetes mellitus	29 (39.7%)	64 (25.7%)	0.066
Dyslipidemia	11 (15.1%)	17 (6.8%)	0.086
Myocardial infarction	6 (8.2%)	24 (9.6%)	0.728
Angina	12 (16.4%)	27 (10.8%)	0.406
Congestive heart failure	9 (12.3%)	10 (4.0%)	0.014
Hemorrhagic stroke	0 (0%)	2 (0.8%)	0.442
Ischemic stroke	15 (20.5%)	13 (5.2%)	0.002
TIA	1 (1.4%)	1 (0.4%)	0.354
Family history of CAD	4 (5.5%)	14 (5.6%)	0.988
Killip class ≥ II	42 (57.5%)	173 (69.5%)	0.120
ST elevation MI	28 (38.4%)	122 (49.0%)	0.031
PCI	53 (72.6%)	214 (85.9%)	0.012
IV Thrombolysis	1 (1.4%)	3 (1.2%)	1.000

Results

Medications

	Group 1 (N = 73)	Group 2 (N = 249)	p value
Aspirin	71 (97.3%)	248 (99.6%)	0.130
Clopidogrel	64 (87.7%)	221 (88.8%)	0.836
Prasugrel	4 (5.5%)	29 (11.6%)	0.186
Ticagrelor	3 (4.1%)	15 (6.0%)	0.774
Cilostazol	8 (11.0%)	40 (16.1%)	0.352
CCB	11 (15.1%)	25 (10.0%)	0.289
Beta-blockers	56 (76.7%)	208 (83.5%)	0.225
ACE inhibitors	22 (30.1%)	114 (45.8%)	0.022
ARB	34 (46.6%)	91 (36.5%)	0.134
Statins	58 (79.5%)	224 (90.0%)	0.025

ACE; angiotensin-converting enzyme, ARB; angiotension receptor blockers, CCB; calcium channel blockers

Results

In-hospital Clinical Events

	Group 1 (N = 73)	Group 2 (N = 249)	p value
Cardiogenic shock	11 (15.1%)	26 (10.4%)	0.529
New-onset HF	6 (8.2%)	10 (4.0%)	0.327
Recur ischemia	2 (2.7%)	5 (2.0%)	0.724
Re-MI	0 (0%)	3 (1.2%)	0.430
Stent thrombosis	0 (0%)	3 (1.2%)	0.346
Ischemic stroke	2 (2.7%)	1 (0.4%)	0.162
Hemorrhagic stroke	1 (1.4%)	0 (0%)	0.148
Decrease of Hb >5 g/dL	1 (1.4%)	1 (0.4%)	0.518
Decreased of Hct >15%	1 (1.4%)	3 (1.2%)	0.744
Minor bleeding	2 (2.7%)	12 (4.8%)	0.443

Hb; hemoglobin, Hct; hematocrit, HF; heart failure, Re-MI; re-attack of myocardial infarction

Results

Medications at 1 Year

	Group 1 (N = 73)	Group 2 (N = 249)	p value
Aspirin	49 (67.1%)	210 (84.3%)	0.006
Clopidogrel	33 (45.2%)	165 (33.3%)	0.011
Prasugrel	1 (1.4%)	12 (4.8%)	0.447
Ticagrelor	0 (0%)	1 (0.4%)	1.0
Cilostazaol	1 (1.4%)	9 (3.6%)	0.334
CCB	10 (13.4%)	43 (17.3%)	0.840
Beta-blockers	49 (67.1%)	200 (80.3%)	0.707
ACE inhibitors	7 (9.6%)	50 (20.1%)	0.326
ARB	37 (50.7%)	114 (45.8%)	0.687
Statins	58 (79.4%)	215 (86.3%)	0.097
Oral anticoagulant	43 (58.9%)	18 (7.2%)	<0.001

ACE; angiotensin-converting enzyme, ARB; angiotension receptor blockers, CCB; calcium channel blockers

Results

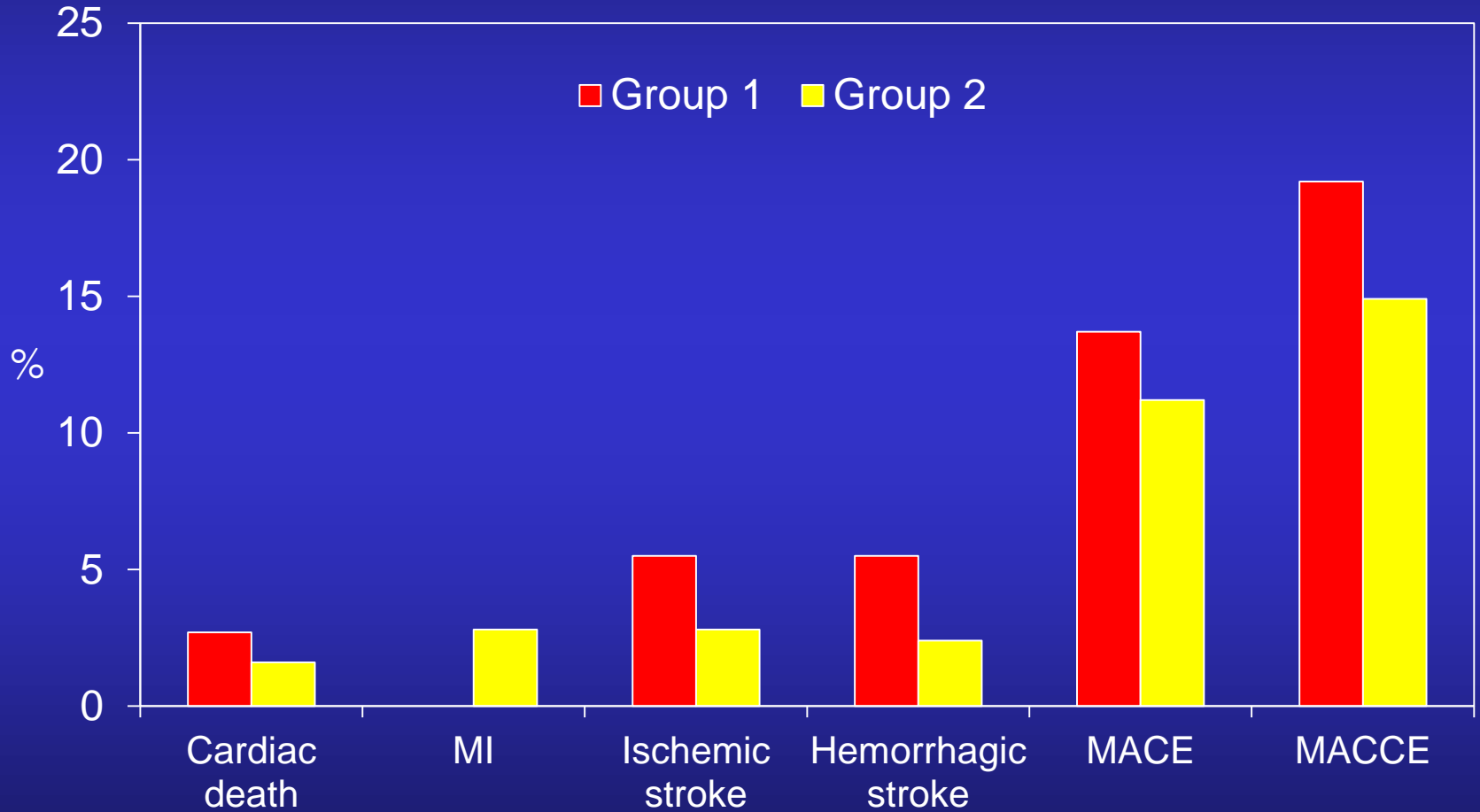
Clinical Events at 1 Year

	Group 1 (N = 73)	Group 2 (N = 249)	p value
All-cause death	3 (4.1%)	7 (2.8%)	0.574
Cardiac death	2 (2.7%)	4 (1.6%)	0.529
Myocardial infarction	0 (0%)	7 (2.8%)	0.148
Re-hospitalization due to heart failure	5 (6.8%)	9 (3.6%)	0.233
Stent thrombosis	0 (0%)	3 (1.2%)	0.346
Repeated PCI	3 (4.1%)	7 (2.8%)	0.574
Coronary a bypass graft	0 (0%)	1 (0.4%)	0.588
Ischemic Stroke	4 (5.5%)	7 (2.8%)	0.270
Cerebral infarction	4 (5.5%)	6 (2.4%)	0.184
Transient ischemic attack	0 (0%)	1 (0.4%)	0.588
Hemorrhagic Stroke	0 (0%)	2 (0.8%)	0.442
MACE	10 (13.7%)	28 (11.2%)	0.568
MACCE	14 (19.2%)	37 (14.9%)	0.374

MACE; major adverse cardiac events, MACCE; major adverse cardiocerebrovascular events, PCI; percutaneous coronary intervention

Results

Clinical Events at 1 Year



Summary

- Patients with OAC
 - Older (70.0 ± 10.1 vs. 67.0 ± 12.3 years)
 - More women (45% vs. 24%), and history of heart failure (12% vs. 4%) or ischemic stroke (22% vs. 6%)
- No differences of new ischemic stroke, major bleeding complications such as cerebral hemorrhage or decrease of hematocrit $>15\%$, and minor bleeding complications
- At 1 year, 59% of patients in group 1 and 7% of patients in group 2 were taking OAC. Aspirin was being taken in 67% and 84%, and P2Y12 inhibitors in 47% and 39% of patients in group 1 and 2, respectively.
- No difference of 1-year major adverse cardiac (13.7% vs. 11.2%).
- No difference of new ischemic stroke (5.5% vs. 2.8%) or hemorrhagic stroke (0% vs. 0.8%) at 1 year

Conclusion

Anticoagulation in selected patients with acute myocardial infarction who are taking antiplatelet agents may not increase the bleeding risk.

Thank You for Your Attention!

