Comparison of the efficacy and safety between ticagrelor and clopidogrel in patients with acute coronary syndrome undergoing percutaneous coronary intervention



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Background and Importance

The European Society of Cardiology (ESC) and American Heart
Association guidelines (AHA) recommend ticagrelor and prasugrel over
clopidogrel in dual antiplatelet therapy (DAPT) following percutaneous
coronary intervention (PCI) in patients with acute coronary syndrome (ACS).¹⁻²⁾

- Clopidogrel: Slower onset, variable efficacy due to genetic polymorphisms
- Ticagrelor & Prasugrel: Faster, more potent antiplatelet effects
 - ⇒ Superior clinical outcomes in ACS
 - Limitations: Higher bleeding risk
 - Asian population: Limited studies;

thrombotic and bleeding tendencies differ from non-Asians

Aim and Objectives

Compare the efficacy and safety of ticagrelor and clopidogrel in patients with ACS undergoing PCI, elucidating optimal antiplatelet therapy tailored to individual patient needs.

⇒ Need for domestic studies to optimize antiplatelet therapy in Asian patients

Materials and Methods

1. Research design

- Retrospective observational study
- ACS patients undergoing PCI at a tertiary hospital (Jan 2022 Jun 2023)
- Exclusion criteria: patients under 18 years of age, cases with insufficient medical records
- Groups: ticagrelor, clopidogrel
- * Review of electronic medical records over 1 year post-PCI
- * Analysis was divided into ticagrelor group and clopidogrel group based on the first P2Y12 inhibitor prescribed after PCI.

2. Outcomes



Major adverse cardiovascular events (MACEs)

: Death, MI, stroke, and Revascularization



Major bleeding events

Results

1. Distribution of Prescriptions

- : 460 patients included,
- Group: Ticagrelor (n=129, 28.0%), Clopidogrel (n=331, 72.0%)

2. Patient Characteristics

- Clopidogrel group (compared to Ticagrelor group)
- : Age ↑, male ↓, HTN/DM ↑,
 - Length of hospital stay ↑, Myocardial infarction (ACS type) ↓

3. Efficacy (Table1)

- Before PSM: No significant difference (23.8% vs. 18.1%, p=0.756)
- After PSM: No significant difference (24.5% vs. 18.4%, p=0.460)

4. Safety (Table2)

- Before PSM: No significant difference (3.1% vs. 3.9%, p=0.789)
- After PSM: No significant difference (0.0% vs. 4.1%, p=0.495)

Table 1 Efficacy outcomes in patients treated with ticagrelor or clopidogrel

Efficacy outcomes	Before PSM (n=460)		After PSM (n=98)	
	Tica (n=129)	Clo (n=331)	Tica (n=49)	Clo (n=49)
MACEs, no. (%)	25 (23.8)	60 (18.1)	12 (24.5)	9 (18.4)
Death	4 (3.1)	23 (6.9)	3 (6.1)	2 (4.1)
MI	11 (8.5)	22 (6.6)	4 (8.2)	3 (6.1)
Stroke	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)
Revascularization*	15 (11.6)	32 (9.7)	7 (14.3)	6 (12.2)

^{*}Revascularization: CABG, PCI

Table 2 Safety outcomes in patients treated with ticagrelor or clopidogrel

Safety outcomes	Before PSM (n=460)		After PSM (n=98)	
	Tica (n=129)	Clo (n=331)	Tica (n=49)	Clo (n=49)
Major bleeding, no. (%)	4 (3.1)	13 (3.9)	0 (0.0)	2 (4.1)

All values are P>0.05

- To minimize bias between the two groups, propensity score matching (PSM) was performed.
- Patients who switched P2Y12 inhibitors within 1 year after PCI (n=78) were excluded from the matching process.

Conclusion and Relevance

- **Guideline Recommendation**: Ticagrelor is recommended over clopidogrel for DAPT post-PCI per ESC and ACC/AHA/SCAI guidelines.¹⁻²⁾
- Clinical Insights: Clopidogrel was preferred in older, high-risk patients with comorbidities and poor renal function, reflecting real-world decisionmaking based on patient characteristics.
- **Study Outcomes**: PSM analysis showed no significant differences in MACEs or major bleeding rates between ticagrelor and clopidogrel groups, suggesting similar efficacy and safety in this population.
- Meta-analysis Findings (Wu et al., 2020)³⁾: Regional Differences
- ✓ Efficacy of ticagrelor:
 - Europe/US: Significant reduction in MACEs (OR=0.82, p<0.001)
- Asia: More pronounced effect (OR=0.66, P<0.001)
- ✓ Bleeding Events of ticagrelor:
 - Overall population: No significant difference (OR=1.19, p=0.21)
 - Asian patients: Higher risk with ticagrelor (OR=1.52, p<0.001)



Further multicenter, prospective studies are needed to provide stronger evidence for the optimal selection of antiplatelet agents.

References

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All values are P>0.05