

# Highest Efficacy of Dual Antiplatelet Therapy After Carotid Artery Stenting in High Bleeding Risk Patients

NCT06276374

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Status	RECRUITING
Phase	Not Applicable
Sponsor	Woo-Keun Seo
Enrollment	1,556 participants

## Key Eligibility Criteria

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### Inclusion (18)

- Patients ≥19 years
  - Symptomatic patients with carotid artery stenosis ≥50% and asymptomatic patients with carotid artery stenosis ≥70% who are scheduled to undergo or who have undergone carotid artery stenting
  - High bleeding risk is defined as a Bleeding Academic Research Consortium type 3 or 5 bleeding risk of ≥4% at 1 year or a risk of an intracranial hemorrhage (ICH) of ≥1% at 1 year, Patients who meet at least one of the criteria for high bleeding risk below
  - The degree of stenosis is determined using the method performed in the North American Symptomatic Carotid Endarterectomy Trial.
  - Criteria for high bleeding risk (see 1)
- ... and 13 more (see full listing online)

### Exclusion (8)

- Incidence of net clinical events, including cardiovascular and cerebrovascular accidents or major bleeding events, within 30 days following carotid artery stenting
  - Discontinuation of dual antiplatelet therapy within 30 days after carotid stent implantation (However, use of a single antiplatelet therapy within 7 days due to acute infection and trauma is allowed, but dual antiplatelet therapy must be administered at 28 to 30 days after carotid stent implantation)
  - Coronary artery stenting or other vascular stenting or vascular recanalization within 1 year (Revascularization surgery that requires CABG(Coronary Artery Bypass Graft) and other dual antiplatelet therapy)
  - Aspirin or clopidogrel hypersensitivity
  - Pregnant or breastfeeding women (Women of childbearing need to check for pregnancy using urine or blood tests before enrollment, and use appropriate contraception methods during the clinical trial period)
- ... and 3 more (see full listing online)

## Locations (1 total)

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Samsung Medical Center, Seoul, South Korea

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<https://clinicaltrials.gov/study/NCT06276374>

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