Safety and Efficacy of Paclitaxel Coated PTCA Balloon Catheter With a Shellac Plus Vitamin E Excipient (GENOSS® DCB) in Patients With Coronary In-stent Restenosis (ISR): A Prospective, Multi-center, Observational Study

NCT06104007

Status	RECRUITING
Sponsor	Seoul National University Hospital
Enrollment	1,000 participants

Key Eligibility Criteria

Inclusion (2)

- Patients with coronary in-stent restenosis (ISR) who underwent percutaneous coronary intervention using the Genoss® DCB.
- Participants who have agreed to the clinical trial protocol and the clinical follow-up study plan, voluntarily decided to participate in this clinical research, and have provided written consent on the research participant agreement form.

Exclusion (6)

- Women of childbearing age who plan to become pregnant during the study duration.
- Patients scheduled for a surgery within 12 months of enrollment that requires discontinuation of antiplatelet agents.
- Patients for whom the expected remaining life span is less than one year.
- Patients who presented with cardiogenic shock at the time of their visit and, based on medical assessment, are predicted to have a low likelihood of survival.
- Patients currently involved in a randomized medical device study.

... and 1 more (see full listing online)

Locations (1 total)

Bon-Kwon Koo, Seoul, South Korea