The Effects of Ripasudil in Patients With FED Undergoing Femtosecond Laser Assisted Cataract Surgery

NCT06048380

Status	RECRUITING
Phase	Phase 3
Sponsor	Singapore Eye Research Institute
Enrollment	120 participants

Key Eligibility Criteria

Inclusion (4)

- Patients with FED who will be listed for cataract surgery as part of standard clinical practice (A cataract will be defined as clouding of the lens that interferes with normal vision).
- Only patients older than 50 years will be included, as cataracts develops with increasing age. No gender criteria are applied.
- Ability to provide informed consent, and are willing and able to sign a written Informed Consent Form prior to any study-specific procedures.
- Patients must be willing and able to return for scheduled follow-up examinations for up to 12 months after the surgery.

Exclusion (5)

- Patients who are unable to give consent.
- An only-functioning eye in a patient who has lost visual potential in the contralateral eye.
- Diagnosis of clinically relevant eye diseases that may interfere with the aim of the study.
- History of ripasudil use, as well as other systemic disease such as cardiovascular or renal disease.
- Patients who have previous allergic reactions to the contents present in Glanatec Ophthalmic solution 0.4%

Locations (1 total)

Singapore National Eye Centre, Singapore, Singapore, Singapore