Feasibility Study of an Accommodating IOL Design

NCT07147192

Status	RECRUITING
Phase	Not Applicable
Sponsor	Alcon Research
Enrollment	85 participants

Key Eligibility Criteria

Inclusion (4)

- Able to understand and sign an Informed Consent Form.
- Willing and able to attend all scheduled study visits required per protocol.
- Diagnosed with bilateral cataracts requiring removal by phacoemulsification.
- Preoperative corneal astigmatism equal to or less than 1.50 diopter (D) in both eyes.

Exclusion (4)

- Women of childbearing potential who are pregnant, intend to become pregnant during the study, or are breastfeeding.
- Taking medications that could increase risk or may affect accommodation.
- Eye conditions as specified in the protocol, including glaucoma or ocular hypertension.
- Medical conditions that could increase operative risk as specified in the protocol.

Locations (5 total)

Contact Alcon Call Center for Trial Locations, Fort Worth, Texas, United States
Clinica 20/20, San José, Costa Rica
Instituto Espaillat Cabral, Santo Domingo, Dominican Republic
... and 2 more locations