

Evaluate the Safety and Effectiveness of the AccuraSee™ IOPCL for Secondary Implantation in the Capsular Bag to Improve Near and/or Intermediate Vision Following Previous Cataract Surgery

NCT06625749

Status	RECRUITING
Phase	Not Applicable
Sponsor	OnPoint Vision Inc
Enrollment	10 participants

Key Eligibility Criteria

Inclusion (19)

- All ocular eligibility criteria refer to the study (non-dominant) eye only unless otherwise noted.
 - Subjects aged 22 years and older.
 - Subjects who have had cataract surgery with an Alcon monofocal intraocular lens model SN60WF, SA60WF, or SA60AT (with a lens power from 10.0D to 30.0D), or Johnson and Johnson monofocal lens model ZCB00 (with a lens power from 10.0D to 26.0D), or Zeiss monofocal lens model CT LUCIA 602 (with a lens power from 10.0D to 19.0D) clearly evidenced by photographic documentation with one of the following: patient medical record, clinic chart with labeling attached, surgical record with labeling attached, or patient identification card with make, model, power and serial number.
 - Subjects who have had cataract surgery at least 6 months from the planned date of IOPCL surgery.
 - Subjects who require a reading add of +1.50 to +2.50 to achieve an BCNVA of 20/32 or better.
- ... and 14 more (see full listing online)

Exclusion (43)

- Subjects who have had cataract surgery with other than a monofocal posterior chamber intraocular lens (PCIOL).
 - Subjects who have had previous laser refractive surgery.
 - Subjects who have had cataract surgery with an Alcon monofocal intraocular lens model SN60WF, SA60WF, or SA60AT (with a lens power below 10.0D or greater than 30.0D), or Johnson and Johnson monofocal intraocular lens model ZCB00 (with a lens power below 10.0D or greater than 26.0D) or Zeiss monofocal lens model CT LUCIA 602 (with a lens power below 10.0D or greater than 19.0D).
 - Subjects who were treated with a PCIOL in a manner that is not consistent with the labeling, contraindications, or indications for use statement.
 - Subjects whose continuous curvilinear capsulorhexis is less than 4.5 mm or more than 6.0 mm in size at the time of the preoperative visit.
- ... and 38 more (see full listing online)

Locations (1 total)

Midwest Vision Partners, Cleveland, Ohio, United States

<https://clinicaltrials.gov/study/NCT06625749>

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