Bicentric Clinical Investigation to Assess Safety and Performance of LuxBoost IOL

NCT06258707

Status	RECRUITING
Phase	Not Applicable
Sponsor	Cutting Edge SAS
Enrollment	58 participants

Key Eligibility Criteria

Inclusion (11)

- Subject aged 50 or over on the day of inclusion, presenting a
- bilateral cataract for which posterior chamber IOL implantation
- has been planned.
- Fit within the available IOL diopter range.
- Have had no previous refractive surgery.

... and 6 more (see full listing online)

Exclusion (10)

- Ocular surface disease potentially affecting study results
- Subjects suffering from diagnosed degenerative visual disorders
- Pre-existing ocular pathology
- Acute or chronic disease or illness that would increase risk or confound study results
- Axial lengths and keratometry such as the IOL spherical power is

... and 5 more (see full listing online)

Locations (2 total)

Clinique Honore Cave, Montauban, France
West Ophta, Rennes, France