

Safety, Tolerability and Efficacy of Intravitreal KIO-104 in Patients With Macular Edema

NCT06825702

Status	RECRUITING
Phase	Phase 2
Sponsor	Kiora Pharmaceuticals, Inc.
Enrollment	28 participants

Key Eligibility Criteria

Inclusion (17)

- Participants must meet all the following criteria:
- Be aged 18 to 85 years inclusive at the time of consent.
- Provide informed consent prior to any study procedures, as stipulated by local laws, Ethics Committee (EC) and Regulatory Authority (RA) guidelines.
- Be willing and able to follow all study instructions, attend all study visits, and complete all study assessments.
- Have a clinical diagnosis of ME in the study eye secondary to non-infectious uveitis, retinal vein occlusion, diabetic retinopathy or cataract surgery.

... and 12 more (see full listing online)

Exclusion (27)

- Participants must not meet any of the following criteria:
- Have media opacities (cornea, anterior or posterior synechia, cataract, vitreous haze and others) of either eye that preclude investigation and documentation of the posterior pole and intravenous fluorescein angiography, or optical coherence tomography evaluation in the study eye.
- Receive local or systemic biologicals (i.e. tumour necrosis factor \[TNF\]-blockers, B-cell blockers, cytokines, cytokine-blockers, receptor antagonists) 90 days prior to Day 1 or planned during the study.
- Receive treatment with cyclophosphamide or chlorambucil during the study.
- Receive intravitreal injections (including but not limited to anti-vascular endothelial growth factors) 90 days prior to Day 1 or planned during the study.

... and 22 more (see full listing online)

Locations (4 total)

Sydney Eye Hospital, Sydney, New South Wales, Australia
Royal Adelaide Hospital, Adelaide, South Australia, Australia
Centre for Eye Research Australia, East Melbourne, Victoria, Australia
... and 1 more locations

<https://clinicaltrials.gov/study/NCT06825702>

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