

A Registry of Subjects With Primary Indeterminate Lesions or Choroidal Melanoma

NCT03941379

Status	RECRUITING
Sponsor	Aura Biosciences
Enrollment	500 participants

Plain Language Summary

This is a long-term follow-up registry for patients who previously participated in a clinical trial testing a drug called AU-011 (bel-sar) for eye tumors — specifically indeterminate lesions (spots that may or may not become cancer) or choroidal melanoma (a type of eye cancer). The registry collects ongoing safety and disease status information from these patients over time.

****You may be eligible if...****

- You previously participated in an Aura Biosciences clinical trial for an eye lesion or choroidal melanoma
- You received AU-011 (bel-sar), a placebo, standard of care, or were in an observation group during that prior trial
- You are willing to provide ongoing health updates

****You may NOT be eligible if...****

- You did not participate in a prior Aura Biosciences sponsored clinical trial
- You are unwilling or unable to consent to ongoing follow-up

Talk to your doctor to see if this trial is right for you.

Key Eligibility Criteria

Inclusion (2)

- Have been clinically diagnosed with primary IL or CM at the time of entry to a previous Aura Biosciences sponsored clinical trial.
- Have received AU-011 bel-sar or assigned to a different cohort or treatment arm (sham, SoC, etc.) an observation cohort in a previous Aura sponsored clinical trial.

Exclusion (1)

- None

Locations (16 total)

University of Wisconsin, Madison, Wisconsin, United States
Columbia University Medical Center, New York, New York, United States
Wills Eye Hospital, Philadelphia, Pennsylvania, United States
... and 13 more locations

<https://clinicaltrials.gov/study/NCT03941379>

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