

# The Efficacy of a Smart Phone-based Test on Measuring Pupillary Light Reflex Alterations Following Cannabis Use Healthy in Adults

NCT06967051

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<b>Status</b>	RECRUITING
<b>Phase</b>	Not Applicable
<b>Sponsor</b>	Sobereye Inc.
<b>Enrollment</b>	20 participants

## Key Eligibility Criteria

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### Inclusion (14)

- Males and females 21 years of age or older
  - Females not of child-bearing potential, defined as those who have undergone a sterilization procedure (e.g. hysterectomy, bilateral oophorectomy, bilateral tubal ligation, complete endometrial ablation) or have been post-menopausal for at least 1 year prior to screening Or,
  - Individuals of child-bearing potential must have a negative baseline urine pregnancy test and agree to use a medically approved method of birth control for the duration of the study. All hormonal birth control must have been in use for a minimum of three months. Acceptable methods of birth control include:
    - Hormonal contraceptives including oral contraceptives, hormone birth control patch (Ortho Evra), vaginal contraceptive ring (NuvaRing), injectable contraceptives (Depo-Provera, Lunelle), or hormone implant (Norplant System)
    - Double-barrier method
- ... and 9 more (see full listing online)

### Exclusion (23)

- Individuals who are pregnant, breast feeding or planning to become pregnant during the study
  - Allergy, sensitivity, intolerance, or dietary restriction preventing consumption study products
  - Current and ongoing neurological or ophthalmological issue that could affect the retina (blindness, glaucoma, dry eyes, retinal diseases, pupil abnormalities, cataracts, sensitivity to bright lights)
  - History of surgery on eyes or retinas except for laser corneal surgery
  - Current or history of psychological disorders (e.g., schizophrenia and psychosis)
- ... and 18 more (see full listing online)

## Locations (1 total)

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KGK Science Inc., London, Ontario, Canada

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<https://clinicaltrials.gov/study/NCT06967051>

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