

WHAT IS THE GOAL OF THE STUDY?

To evaluate the clinical performance of an investigational soft contact lens compared to a commercially available lens.

WHO MAY PARTICIPATE IN THE STUDY?

Those who meet the following criteria may be eligible to participate:

- Age 18 and over
- Prescription between -1.00 and -6.00 D (inclusive), astigmatism ≤ 0.75 D in each eye
- Current spherical soft contact lens wearer in both eyes who has worn contact lenses for a minimum of 5 days per week and 12 hours per day during the past 3 months
- Willing to wear study lenses for at least 12 hours per day, over an 8-day period per lens, and for at least 18 hours on the day prior to each follow-up visit (Visits 3 & 5)
- Willing to use own digital device to access select study questionnaires
- Must have spectacles and be willing to wear them when study lenses are not worn
- Have normal, healthy eyes with good corrected vision

Further screening questions will be asked prior to scheduling an appointment.

WHAT DOES THE STUDY INVOLVE?

Participants will first have a screening visit to verify that they meet the requirements of the study.

The study involves 5 in-office visits over a 19 to 27-day period. Subjects will be asked to wear two different types of soft contact lenses. The lenses will be provided at no cost.

WILL I BENEFIT FROM PARTICIPATING?

There is no expected direct benefit, however your participation may benefit future patients.

WHAT ELSE SHOULD I KNOW?

Participants will be compensated up to \$470 for their time.

WHO CAN I CONTACT TO LEARN MORE ABOUT THE STUDY?

You may contact the Clinical Vision Research Center at the SUNY College of Optometry

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