

Lacriprep for Corneal Wound Healing Study

NCT06854393

Status	RECRUITING
Phase	Phase 2
Sponsor	Henry M. Jackson Foundation for the Advancement of Military Medicine
Enrollment	88 participants

Key Eligibility Criteria

Inclusion (7)

- Male or female, of any race, 21 years old and older
- Active-duty U.S. military service members or their dependents eligible for care at Alexander T. Augusta Military Medical Center for 6-month period postoperatively
- Desiring refractive correction of myopia up to -4.00 diopters (D) sphere with or without astigmatism 0 to -1.50 D and up to -4.00 D manifest spherical equivalent in both eyes
- Stable refraction for the past year as demonstrated by a change in manifest refractive spherical equivalent of ≤ 0.50 diopters (D) in both eyes
- CDVA of at least 20/20 in both eyes
- ... and 2 more (see full listing online)

Exclusion (11)

- Calculated ablation depth of greater than 75 microns or other conditions in either eye which in the judgement of the investigators will require use of prophylactic MMC
- Signs of unstable or progressive myopia, keratoconus or keratoconus suspect in either eye
- Abnormal corneal topographic findings in either eye e.g., keratoconus, pellucid marginal degeneration
- Treatment targeted for monovision
- Signs and symptoms of dry eye based on: (a) Corneal fluorescein staining total score ≥ 4 in the National Eye Institute (NEI)/Industry Workshop scale in either eye, (b) Anethetized Schirmer test score ≤ 5 mm/5 minutes in either eye, (c) Symptom score ≥ 40 using the Symptom Assessment in Dry Eye (SANDE) questionnaire in either eye
- ... and 6 more (see full listing online)

Locations (1 total)

Alexander T. Augusta Military Medical Center, Fort Belvoir, Virginia, United States

<https://clinicaltrials.gov/study/NCT06854393>

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