

# Phase 1b/2a Randomized Double-blind Study to Investigate Safety Tolerability PK PD Preliminary Efficacy of Oral Administration of SNIPR001 in Patients With Hematologic Malignancy Scheduled for Allogeneic Hematopoietic Stem-Cell Transplant Receiving FQ Prophylaxis & Harboring FQR Ecoli Pre-Transplant

NCT06938867

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Status	RECRUITING
Phase	Phase 1, Phase 2
Sponsor	SNIPR Biome Aps.
Enrollment	24 participants

## Key Eligibility Criteria

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### Inclusion (9)

- Male or female ≥18 years of age at the time of consent.
- Patient is able and willing to provide written informed consent prior to any study-related procedure.
- Confirmed diagnosis of any hematologic malignancy.
- Planned to undergo an allogeneic hematopoietic stem cell transplant.
- Patient is scheduled to receive fluoroquinolone (levofloxacin) prophylaxis.

... and 4 more (see full listing online)

### Exclusion (8)

- Use of any treatment (approved or investigational product) considered to interact with the study drug, or which might impact the outcome of the study within 14 days (or 5 half-lives of the approved or investigational product, whichever is greater) prior to the first administration of study drug, as judged by the Investigator.
- Use or planned use of any antibiotics with intrinsic activity against E. coli in the gut (e.g., beta-lactam antibiotics) between Pre-Screening and until the end of the SNIPR001/placebo treatment period, with the exception of TMP-SMX and levofloxacin.
- Have known hypersensitivity or allergy to any component of SNIPR001, levofloxacin and/or Alka-Seltzer Gold treatment.
- Unwilling or unable to comply with the requirements of this Protocol, including providing stool samples.
- Female patients who are pregnant or lactating.

... and 3 more (see full listing online)

## Locations (8 total)

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City of Hope, Duarte, California, United States  
University of California, San Francisco, San Francisco, California, United States  
John Hopkins University, Baltimore, Maryland, United States

... and 5 more locations

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<https://clinicaltrials.gov/study/NCT06938867>

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