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# Proof-of-concept Study to Evaluate the Safety, Tolerability, Pharmacodynamics, and Pharmacokinetics of SION-719 When Added to Trikafta

**Code:** NCT07108153    **Year:** 2025    **Date:** 2025

**Author:**

## Study design (if review, criteria of inclusion for studies)

Allocation: RANDOMIZED|Intervention Model: CROSSOVER|Masking: QUADRUPLE (PARTICIPANT, CARE\_PROVIDER, INVESTIGATOR, OUTCOMES\_ASSESSOR)|Primary Purpose: TREATMENT

## Participants

ADULT, OLDER\_ADULT with CF

## Interventions

DRUG: SION-719|DRUG: Placebo-to-match SION-719

## Outcome measures

Adverse events [Safety and Tolerability] when SION-719 is administered to people with cystic fibrosis (CF) who are taking a standard stable dose of physician-prescribed Trikafta

<https://clinicaltrials.gov/study/NCT07108153>

## Keywords

CFTR Modulators; Genetic Predisposition to Disease; pharmacological\_intervention; placebo; VX-770; VX-661; ivacaftor; Aminophenols; tezacaftor; VX-445; elexacaftor; Trikafta; SION-719; CFTR NBD1 inhibitors;