

# An Investigational Drug (TPST-1495) in Patients With Familial Adenomatous Polyposis

NCT06557733

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<b>Status</b>	RECRUITING
<b>Phase</b>	Phase 2
<b>Sponsor</b>	National Cancer Institute (NCI)
<b>Enrollment</b>	38 participants

## Key Eligibility Criteria

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### Inclusion (24)

- Diagnosis of familial adenomatous polyposis (FAP), defined as at least one of the following:
  - Genetic diagnosis with confirmed APC mutation (clinical CLIA [clinical laboratory improvement amendments] certified lab or research testing)
  - Obligate carrier
  - Clinical diagnosis of classic FAP with e 100 colorectal adenomas status post colectomy or a sub-total colectomy and a family history of FAP
  - Clinical diagnosis of FAP, based on personal and family history. Note: This criterion requires documented review and agreement from either the study chair or the MW consortium lead investigator
- ... and 19 more (see full listing online)

### Exclusion (11)

- Use of any other investigational agents d 12 weeks prior to pre-registration
  - History of gastric or intestinal ulceration due to NSAID therapy
  - Uncontrolled intercurrent illness or recent surgical procedure that in the opinion of the investigative team would limit compliance with study requirements
  - History of invasive malignancy d 3 years prior to pre-registration (exception: adequately treated carcinoma of the cervix, carcinoma in situ, or basal or squamous cell carcinomas of the skin)
  - History of any upper GI surgery that does not permit access to or evaluation of a 10 cm segment of the duodenum that includes the duodenal bulb, i.e. Whipple procedure or similar
- ... and 6 more (see full listing online)

## Locations (4 total)

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Mayo Clinic Hospital in Arizona, Phoenix, Arizona, United States  
Mayo Clinic in Rochester, Rochester, Minnesota, United States  
Huntsman Cancer Institute/University of Utah, Salt Lake City, Utah, United States  
... and 1 more locations

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<https://clinicaltrials.gov/study/NCT06557733>

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