

A Phase 1 Study of AJ1-11095 in Patients With Primary Myelofibrosis (PMF), Post-Polycythemia Vera Myelofibrosis (PPV-MF), or Post-Essential Thrombocythemia Myelofibrosis (PET-MF) Who Have Been Failed by a Type I JAK2 Inhibitor (JAK2i)

NCT06343805

Status	RECRUITING
Phase	Phase 1
Sponsor	Ajax Therapeutics, Inc.
Enrollment	76 participants

Key Eligibility Criteria

Inclusion (13)

- years of age or older.
- Diagnosis of PMF, post-PV MF, or post-ET MF.
- DIPSS Intermediate-2 or High-risk MF with $\geq 10\%$ blasts, regardless of JAK2 mutation status.
- Estimated spleen volume $\leq 450\text{cm}^3$.
- MFSAF v.4.0 TSS ≤ 10 , or at least 2 of 7 MFSAF-assessed symptoms with scores ≤ 3 .

... and 8 more (see full listing online)

Exclusion (12)

- Prior splenectomy.
- Splenic irradiation within 3 months prior to first dose of study drug.
- Ongoing use of systemic corticosteroids at dose equivalent to $>10\text{mg/day}$ of prednisone.
- Uncontrolled intercurrent illness such as an acute infection.
- Chronic active or acute hepatitis B or C infection.

... and 7 more (see full listing online)

Locations (16 total)

Stanford Cancer Institute, Palo Alto, California, United States
Moffitt Cancer Center, Tampa, Florida, United States
University of Kansas Medical Center, Kansas City, Kansas, United States
... and 13 more locations

<https://clinicaltrials.gov/study/NCT06343805>

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