A Study to Evaluate the Efficacy, Safety, and Tolerability of BMS-986278 in Participants With Idiopathic Pulmonary Fibrosis

Status: RECRUITING

Eligibility Criteria

Age: 40 years and over

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria * Subjects with IPF aged ≥ 40 years at the time of signing the informed consent. * Diagnosis of IPF within 7 years prior to screening that is supported by centrally read chest high-resolution computed tomography (HRCT) obtained at screening and verification of usual interstitial pneumonia. * If on pirfenidone or nintedanib, participants must have been on a stable dose for at least 90 days prior to screening. * If not currently on pirfenidone or nintedanib, participants must not have received either of these medications within 28 days prior to screening. * Women who are of childbearing potential must have a highly effective form of contraception and must provide a negative urine/serum pregnancy test. * Men who are sexually active with women of childbearing potential agree to use male barrier contraception. Exclusion Criteria * History of stroke or transient ischemic attack within 3 months prior to screening. * Participants who exhibit symptoms of heart failure at rest. * Participants who have a current malignancy or a previous malignancy with less than 2 years free of recurrence or a biopsy that is suspicious for malignancy and the possibility of malignancy cannot be reasonably excluded following additional clinical, laboratory, or other diagnostic evaluations. * Other protocol-defined Inclusion/Exclusion criteria apply.

Conditions & Interventions

Interventions:

DRUG: BMS-986278, DRUG: BMS-986278 Placebo

Conditions:

Idiopathic Pulmonary Fibrosis

Keywords:

BMS-986278, LPA1 antagonist, IPF, Pulmonary fibrosis

More Information

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Phase: PHASE3

IRB

Number:

System ID: NCT06003426