DeciPHer-ILD: A Real-world Patient Registry in Group 3 Pulmonary Hypertension Associated With Interstitial Lung Disease (PH-ILD)

Status: RECRUITING

Eligibility Criteria

Age: 18 years and over

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

1. Adults aged 18 years or older 2. Diagnosis of ILD by traditional or HRCT determined by the site/institution that conducted the HRCT 3. Patients with connective tissue disease must have a baseline forced vital capacity of \<70% 4. RHC confirmed PH (mean pulmonary artery pressure \>20 mmHg, pulmonary arterial wedge pressure ≤15 mmHg, pulmonary vascular resistance \>2 WU). 5. For patients to be eligible for Cohort 1, they must not be receiving inhaled treprostinil at Baseline. 6. For patients to be eligible for Cohort 2, they must have initiated Tyvaso/Tyvaso DPI at 1 of the following time points: 1. Baseline 2. ≤60 days prior to Baseline 7. For patients to be eligible fo Cohort 3, they must be receiving Tyvaso/Tyvaso DPI at Baseline and for \>60 days prior to Baseline 8. Co-enrollment in other observational or interventional studies is permitted 9. Patient is willing and able to provide informed consent and complete surveys/questionnaires in English or Spanish

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Exclusion Criteria:

1. Confirmed diagnosis of Group 1, 2, 4, or 5 PH 2. Confirmed diagnosis of Group 3 PH related to chronic obstructive pulmonary disease or conditions that cause hypoxemia, such as untreated or inadequately treated obstructive sleep apnea and alveolar hypoventilation disorders 3. Patients receiving Yutrepia (inhaled treprostinil) at Baseline.

Conditions & Interventions

Interventions:
OTHER: Prospective study assessments
Conditions:
Pulmonary Hypertension Due to Lung Diseases and Hypoxia, Pulmonary Hypertension, Interstitial Lung Disease
Keywords:
pulmonary hypertension, interstitial lung disease, pulmonary hypertension associated with interstitial lung disease

More Information

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