

Pulmonary Fibrosis Foundation Patient Registry (PFFR)

Status: Recruiting

Eligibility Criteria

Sex: All

Age: 18 Years to 99 Years old

This study is NOT accepting healthy volunteers

Inclusion Criteria:

1. 18 years old or older 2. Understand and sign the informed consent document 3. ILD Diagnosis must be made / confirmed at a participating Registry center. 1. The diagnostic evaluation must include, at a minimum, a medical history, physical examination, pulmonary function testing and a computerized tomography (CT) scan of the chest. 2. If patients exhibit another pulmonary disease (such as emphysema or asthma), the primary disease must be ILD. 4. Anticipated additional follow up at the Registry center within one year.

Exclusion Criteria:

1. Diagnosed with: 1. Sarcoid 2. Lymphangioleiomyomatosis (LAM) 3. Pulmonary alveolar proteinosis (PAP) 4. Cystic fibrosis (CF) 5. Amyloidosis

Conditions & Interventions

Conditions:

Interstitial Lung Disease (ILD), Idiopathic Pulmonary Fibrosis (IPF)

More Information

Description: The Pulmonary Fibrosis Foundation Patient Registry will collect data on at least 2,000 patients at approximately 40 clinical sites in the US. The Pulmonary Fibrosis Foundation Patient Registry will collect data on at least 2,000 patients at approximately 40 clinical sites in the US. Participants will be asked to complete patient reported outcome (PRO) surveys related to ILD symptoms and quality of life at the time of enrollment and during clinical follow-up visits (Appendix A - PRO Questionnaires). Each patient will donate approximately 30 mL of blood to the Biorepository, which will be separated into plasma, serum, RNA, and DNA.

Contact(s): Mandi DeGrote - carl1032@umn.edu

Principal Investigator: Hyun Kim

Phase: N/A

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