

The PFF Registry™ is collecting data from patients with pulmonary fibrosis (PF) or interstitial lung disease (ILD), including those who have had lung transplants, as well as their, caregivers and family members nationwide to support researchers working to diagnose, treat, and potentially prevent PF and ILD.

The PFF Registry has two distinct data sets:

- **The PFF Patient Registry**, established in 2016, collects medical data from more than 2,000 patients and is administered at PFF Care Centers across the country.
- **The PFF Community Registry**, which relies on ongoing data that is provided directly by the participant.



IT'S ABOUT FINDING ANSWERS

The PFF Registry offers a wealth of longitudinal data on this rare disease, providing unparalleled insight into the experience of patients with PF or ILD, lung transplant recipients, caregivers, and family members.

Leverage the PFF Registry data for your studies now.

- Clinical and translational investigators are invited to utilize PFF Registry data by visiting pffregistry.org or inquiring at registry@pulmonaryfibrosis.org.
- Both academic and industry researchers are welcome to apply.

ENCOURAGE YOUR PATIENTS TO JOIN

Prospective participants can visit pffregistry.org to sign up for the PFF Community Registry.

