

## **HEALTHCARE PROFESSIONALS: BE PART OF THE SOLUTION FOR PULMONARY FIBROSIS (PF) AND INTERSTITIAL LUNG DISEASE (ILD)**

The PFF Registry™ now offers the PFF Community Registry, which provides an opportunity for patients, caregivers, and family to self-report information to help us move toward improved patient outcomes and a better understanding of PF and ILD.

### **Help recruit for the PFF Community Registry to drive improvements in care**

The simple act of promoting awareness of the PFF Community Registry among patients and caregivers will lead to increased participation and a greater chance of finding answers to key questions about PF and ILD.

Data from the PFF Patient Registry has been used in **almost 40 studies** and counting.

### **Encourage patients, caregivers and family members to join the PFF Community Registry**

#### **Participants can:**

- Enroll via the online portal at [pffregistry.org](https://pffregistry.org).
- Complete surveys online at their own pace.
- Choose what information to share.
- Receive periodic research updates via the PFF Registry newsletter.
- Trust that the Registry is highly secure, with technology utilized by hospitals worldwide.

*The PFF Community Registry is open to patients with PF or ILD, lung transplant recipients who have had PF or ILD, and family members and caregivers of patients with PF or ILD.*

### **SHARE THE BENEFITS OF JOINING THE PFF COMMUNITY REGISTRY TODAY!**

Eligible patients, caregivers, or family members can visit [pffregistry.org](https://pffregistry.org) to enroll.



**pffregistry.org**