



Patient powered. Data driven. Research results.

CAREGIVERS AND FAMILY MEMBERS:

WE NEED YOUR HELP TO UNDERSTAND THE IMPACT OF PULMONARY FIBROSIS (PF).

PF and interstitial lung disease (ILD) affects more than a person's lungs. They affect their loved ones and caregivers, who play a critical role in the progression and treatment of a patient with PF or ILD. As a participant in the PFF Community Registry, your story will provide vital information to help us better understand the impact of PF on patients, caregivers and family members.



Share your story to help find

Researchers are working hard to understand the genetic and lifestyle effects of PF and ILD. With your participation, we can fast forward medical research and improve the lives of patients with PF or ILD and their families, for today's and future generations.

Joining the PFF Community Registry is easy

- Visit the PFF Community Registry's easy-to-use online portal to sign up.
- Surveys can be completed at your own pace, with minimal effort and in the comfort of your own home.
- · How much information you share is up to you.
- The PFF Community Registry database is highly secure, using the same technology trusted by doctor's offices, hospitals, and clinical trials worldwide.
- Receive periodic research updates via the PFF Registry newsletter.

SIGN UP TODAY

Caregivers and family members: visit **pffregistry.org** to enroll in the PFF Community Registry now!