You can make a difference

You have the power to make an impact and create hope for PF patients everywhere. Start by making a donation to support the PFF in its mission to help find a cure.

PFF Government Affairs: Advocate and engage with lawmakers regarding the needs of the PF community.

PFF Walk: Build a team, raise funds, and lead the way toward a world without PF by joining one of our national walks. You can participate in-person or virtually.

Fundraise for Team PFF: Create and host an event that is personally meaningful to you. Organize an online fundraiser, hold a bake sale, or host a community walk.

PF Awareness Month: Throughout September, get involved by participating in disease education days, support group events, fundraisers, or with #BlueUp4PF.

Shop PFF: Support the cause when you wear wristbands, t-shirts, caps, and more. Visit Shop-PFF.com.

Our research

The Foundation places enormous importance on creating an environment that will assist in the development of effective treatments for pulmonary fibrosis. The PFF is committed to funding research to identify new therapies and one day, a cure.

PFF Patient Registry: A resource of data gathered from more than 2,000 patient volunteers across the country to help researchers further understand the disease and develop treatments. The data collected can be used to better diagnose, treat, and potentially cure pulmonary fibrosis.

PFF Scholars: A new grant program to support and provide career development opportunities for early stage investigators.

PFF Clinical Trial Finder: A tool to help patients navigate feasible and relevant clinical trials. This searchable platform filters the type of trial, patient characteristics, and proximity to home thereby accelerating the development of new treatment options for patients.

Make a Gift

You can support the PFF’s important programs by making a donation today. Visit pulmonaryfibrosis.org to learn about the many ways you can give.
The Pulmonary Fibrosis Foundation (PFF) mobilizes people and resources to provide access to high quality care and leads research for a cure so people with pulmonary fibrosis (PF) will live longer, healthier lives.

The Trusted Resource
The PFF is the nation’s leading pulmonary fibrosis patient education and advocacy organization. From providing disease education materials to creating the largest patient registry focused on all-cause PF and establishing a nationwide care center network, the PFF is dedicated to improving all aspects of a patient’s quality of life.

We work with leading experts in the field of pulmonary fibrosis. The physicians on our world-class medical team, investigators at each CCN site, and members of the Medical Advisory Board champion collaborative efforts in the care of patients and research.

The PFF is at the forefront of research for different lung diseases that make up the family of pulmonary fibrosis diseases, including idiopathic pulmonary fibrosis (IPF).

We hope you will join us in the fight.

“As the leading advocate for the pulmonary fibrosis community, we are dedicated to advancing the care of people living with these devastating diseases.”

Gregory Cosgrove, MD
PFF Chief Medical Officer