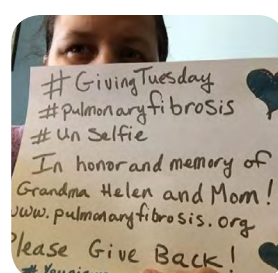
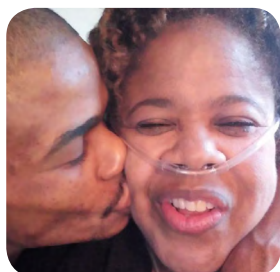


Pulmonary Fibrosis FOUNDATION

ANNUAL REPORT
FISCAL YEAR ENDING JUNE 30, 2016

we imagine a world without
pulmonary fibrosis



LETTER FROM OUR INTERIM CEO AND CHIEF MEDICAL OFFICER

Friends,

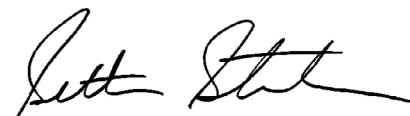
The past year has been one of the most exciting yet for the Pulmonary Fibrosis Foundation (PFF).

In March of 2016, the first patient was enrolled into the PFF Patient Registry, a valuable research tool available in the fight against pulmonary fibrosis (PF). The Registry is a collection of information about PF patients, including their diagnosis and how it was made, lab and test results, and medical outcomes. You can read more about the Registry and the role it is playing in the search for a cure on page 4.

In this report you will also read about the *PFF Summit 2015*, our largest Summit yet (page 10), *Broadway Belts for PFF! 2016* (page 14), the recipients of the research awards the Foundation sponsored in 2015 (page 18), and much more.

We are deeply grateful to the PFF's donors, sponsors, partners, and volunteers. Your contributions lay the groundwork for everything the PFF accomplishes, and help bring us closer to realizing our shared vision of a world without pulmonary fibrosis.

Sincerely,



SCOTT STASZAK
INTERIM CHIEF EXECUTIVE OFFICER



GREGORY P. COSGROVE, MD, FCCP
CHIEF MEDICAL OFFICER



OUR MISSION

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.

WHAT WE DO

By actively engaging the PF community, the Pulmonary Fibrosis Foundation has developed essential programs available to those living and working with pulmonary fibrosis.

Our signature programs include:

- PFF Care Center Network
- PFF Patient Registry
- PFF Patient Communication Center
- PFF Ambassador Program
- An international network of support groups and online communities
- *PFF Summit*
- A comprehensive suite of disease education materials

Our expert Medical Advisory Board and the biennial *PFF Summit* allow us to maintain an ongoing dialogue with physicians, researchers, industry representatives, and the patient community. The Foundation fosters a collaborative environment between these groups. Our peer-reviewed research program supports projects that improve understanding of pulmonary fibrosis and can lead to successful therapies.

The Pulmonary Fibrosis Foundation has a four-star rating from Charity Navigator and is a Better Business Bureau accredited charity.

OUR VISION

We imagine a world without pulmonary fibrosis.

WHAT IS PULMONARY FIBROSIS AND WHAT ARE THE CAUSES?

Pulmonary fibrosis is a condition in which the walls of the air sacs of the lungs become thickened and stiff due to a build up of scar tissue. This scar tissue makes it more difficult for the lungs to transfer oxygen into the bloodstream.

Pulmonary fibrosis can result from a number of causes: certain environmental agents (molds, birds, fibers, dusts), medications (antibiotics, chemotherapy), radiation therapy, autoimmune diseases (scleroderma, rheumatoid arthritis), or a genetic predisposition. However, in most situations the cause is unknown.

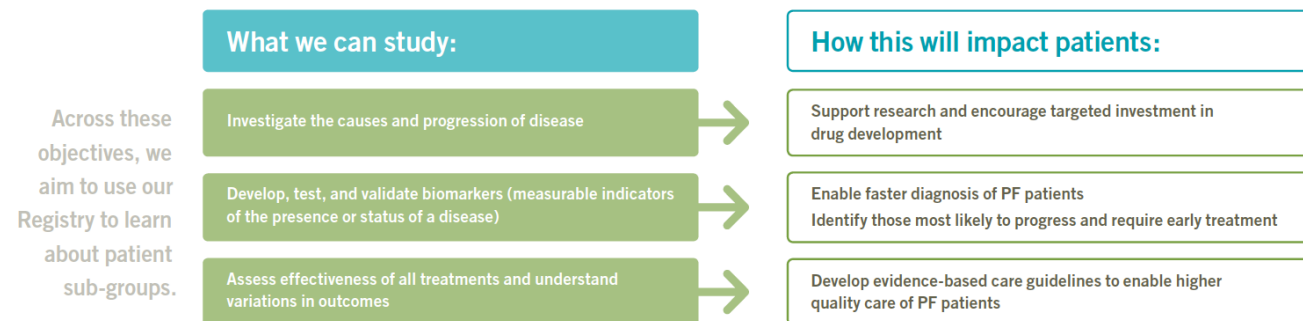
There are many types of PF for which we don't know the cause. One specific type of PF of unknown cause is called "idiopathic pulmonary fibrosis" or IPF. IPF is diagnosed when either a computerized axial tomography scan or a lung biopsy show certain findings in the absence of an identifiable cause. IPF is a progressive, fatal lung disease, for which there is no known cure.

A MAJOR MILESTONE FOR THE PFF PATIENT REGISTRY

In the spring of 2016, the Foundation reached a much-anticipated milestone: the first patient was enrolled into the PFF Patient Registry, the most comprehensive research tool available to help us in the fight against pulmonary fibrosis.

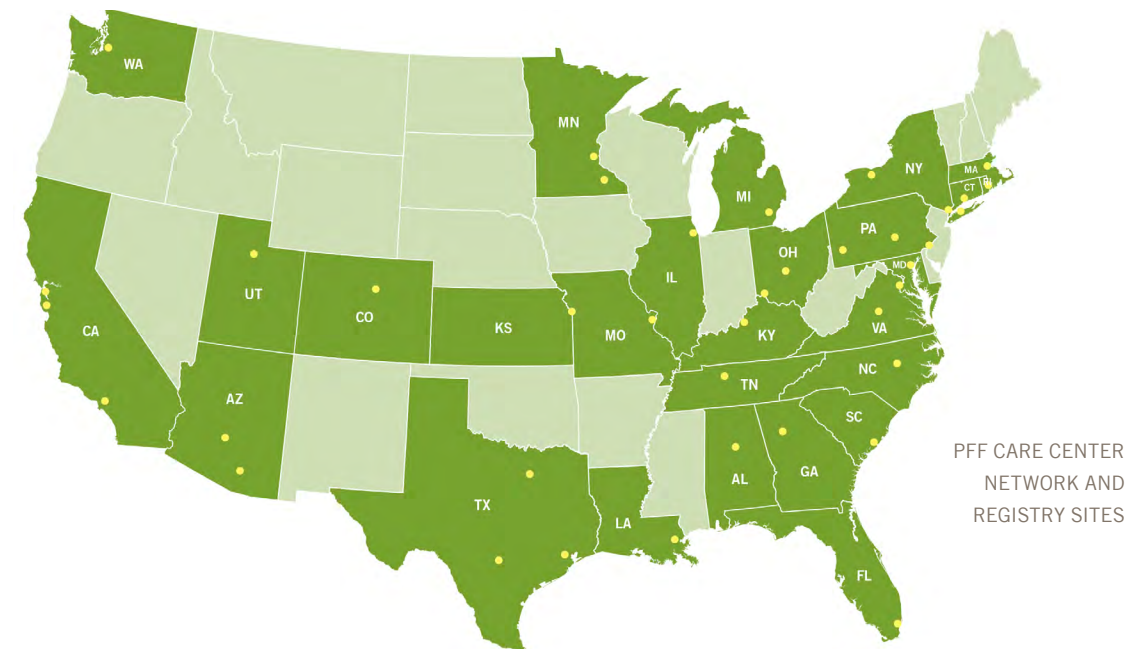
The PFF Patient Registry will permit researchers, through the generous contribution of medical data by patients, to gain a better understanding of how different forms of PF progress, who is affected by PF and how patients respond to different treatments. Patient enrollment takes place at sites in the PFF Care Center Network (CCN), a national network of 40 sites in 27 states across the country. The Registry includes a biorepository, where patients can choose to also donate blood samples. Data from the Registry will help inform best practices in care and identify potential treatment targets. The Registry will also assist in identifying patients who may wish to participate in clinical trials.

The Registry was made possible, first and foremost, by the patients who agreed to participate. We can't thank them enough for their contributions. The initiative was also made possible thanks to the hard work of our PFF Care Center Network colleagues, Registry funders, dedicated volunteers, and partners. Funding for the Registry was generously provided by founding partner Genentech, visionary partner Boehringer Ingelheim, The Cowlin Family Fund at the Chicago Community Trust, the Rattner Family Foundation, the McQuaid Family Foundation, InterMune, Biogen and the Jenny H. Krauss and Otto F. Krauss Charitable Foundation Trust, in memory of Stephen N. Dirks. Further donations will provide expansion opportunities that will enhance this research tool.



“Information collected through the PFF Patient Registry will help facilitate ongoing research necessary to better understand how we can help people living with multiple types of pulmonary fibrosis, including other diseases that often include PF, such as rheumatoid-arthritis associated pulmonary fibrosis,” said Kevin Flaherty, MD, MS, Steering Committee Chairman, PFF Care Center Network and PFF Patient Registry. “This project provides an incredible opportunity to advance progress toward a cure for pulmonary fibrosis.”

The Foundation is grateful to its CCN partners across the country. To learn more about the CCN and to view a full list of the sites, visit pulmonaryfibrosis.org/medical-community/pff-care-center-network.



To contribute to this important program, please donate now.

For more information, call 844.TalkPFF (844.825.5733) or email pcc@pulmonaryfibrosis.org.

HIGHLIGHTS

2015

September THE PF COMMUNITY LET THE WORLD KNOW ABOUT PF DURING **PULMONARY FIBROSIS AWARENESS MONTH**

US CONGRESS FORMALLY RECOGNIZED PULMONARY FIBROSIS AWARENESS MONTH

November LARGEST-EVER **PFF SUMMIT** TOOK PLACE IN WASHINGTON, DC

THE PFF ACQUIRED THE RIGHTS TO THE DAUGHTERS OF PF PROGRAM FROM THE COALITION FOR PULMONARY FIBROSIS. THE FOUNDATION RELAUNCHED THE PROGRAM IN APRIL OF 2016 UNDER THE NAME **PFF DAUGHTERS**

December ON DECEMBER 1, **GIVING TUESDAY**, PFF SUPPORTERS RAISED OVER \$31,000



PFF Senior Medical Advisor, Research Development, Harold R. Collard, MD, speaking at the *PFF Summit 2015*, the largest *Summit* conference to date.

2016

February THE 6TH ANNUAL **BROADWAY BELTS FOR PFF!** RAISED \$180,000 TO BENEFIT THE PF COMMUNITY

March FIRST PATIENT WAS ENROLLED IN THE **PFF PATIENT REGISTRY**

April **PFF DAUGHTERS** RELAUNCHED



Michele Peters, PFF Ambassador, thanking the PF community for a successful Giving Tuesday.



The performing cast of *Broadway Belts for PFF!*

PFF DAUGHTERS members use their collective voice to increase awareness about pulmonary fibrosis, advocate for improved care, and raise support for continued research efforts. Membership in the PFF Daughters program is open to anyone affected by pulmonary fibrosis—past, present, and future. The inaugural application cycle opened in April of 2016.

PFF Daughters

A NEW PFF SIGNATURE PROGRAM

The PFF Daughters began with the Coalition for Pulmonary Fibrosis as a project focusing on women and girls whose parents had been diagnosed with PF. Now part of the Foundation's suite of signature programs, PFF Daughters brings members together to increase awareness about PF, advocate for improved care for people with PF, and raise support for continued research efforts. Membership in the PFF Daughters program is open to anyone affected by pulmonary fibrosis — past, present, and future.

“This is a group that’s really passionate about spreading awareness of PF,” says Kate Gates, PFF Director of Programs. “From hosting support groups and fundraisers, to talking to legislators about PF, to getting the word out on social media, PFF Daughters work really hard to find creative ways of reaching new segments of

the population and engaging them in the fight against PF.”

“We’re here to fully support the PFF’s drive and direction through advocacy, awareness and fundraising,” says Lisa Boyd, PFF Daughters Steering Committee Chair and a Team PFF leader. “All three are absolutely vital aspects of keeping a foundation healthy and strong. We take all our direction from the Foundation and are there to support them.”



PFF PATIENT COMMUNICATION CENTER (PCC)

staff is available to listen and lend support by personally answering every call and email and offering educational resources to patients, caregivers, and health care providers.



need info?
call (toll free)
844-TalkPFF

The PFF’s Information Hub

PROVIDING ANSWERS AND SUPPORT

For those living with pulmonary fibrosis, obtaining the most accurate and current information can be a challenging and frustrating task. The PFF Patient Communication Center (PCC) provides patients, caregivers, and health care providers with the most up-to-date medical information, communicates the availability of support services, and provides information about other essential resources.

“People find a lot of information online and some of it may be helpful,” says David J. Lederer, MD, MS, Senior Medical Advisor, Education and Awareness. “But a lot of information may be misleading or frightening and needs to be put into context. Reliable information about the disease, its expected course, available therapies, and local resources are critical to managing this disease.”

The PCC helps callers find in-person and online support groups where they can reach out to other people with this condition. Dr. Lederer also notes, “When you call the PCC, you can expect to talk to

someone who’s knowledgeable not only about the Foundation and the resources we offer but is also well-informed about the disease itself.”

The PCC received over 3,000 inquiries between July 1, 2015 and June 30, 2016. The top requests were for general disease information and education materials. Questions came in from 35 countries around the world, including Australia, Argentina, Canada, India, Mexico, Italy, Puerto Rico and the United Kingdom. Callers contacted the PCC from 45 US states. The top five states were California, Florida, New York, Texas and Illinois.

The PFF Patient Communication Center is staffed Monday through Friday, 9:00 a.m.–5:00 p.m. CT. Contact the PCC at 844.TalkPFF (844.825.5733) or pcc@pulmonaryfibrosis.org.

PFF SUMMIT 2015 brings community together

Held November 12-14, the *PFF Summit 2015* provided a collaborative environment aimed at improving education and awareness of pulmonary fibrosis (PF) and identifying new approaches to treat, and ultimately cure, the disease.

RECORD ATTENDANCE

A record number of more than 700 researchers, health care professionals, pulmonary fibrosis patients, caregivers and industry leaders from 18 countries gathered in Washington, DC for the third biennial *PFF Summit 2015*, an international scientific conference presented by the PFF.

The *PFF Summit 2015* featured new plenary sessions for the entire PF community and two keynote addresses. “The Future of Drug Development in Pulmonary Fibrosis — A Regulatory Perspective” was delivered by Janet Woodcock, MD, Director, Center for Drug Evaluation and Research at the Food and Drug Administration on “Collaborative Networks as Tools for Improving Clinical Care” was presented by Pat Furlong, president and chief executive officer, Parent Project Muscular Dystrophy, on November 14.



POSTER AWARDEES

Robert Guzy Washington University School of Medicine, “Fibroblast-Specific FGF Signaling in Bleomycin-Induced Pulmonary Fibrosis”

Justin Oldham the University of Chicago, “TOLLIP, MUC5B and the Response to N-acetylcysteine Among Individuals with Idiopathic Pulmonary Fibrosis”

Rachel Scheraga Cleveland Clinic Foundation, “The Mechanosensitive Transient Receptor Potential Vanilloid 4 (TRPV4) Ion Channel Mediates the Pro-resolution/Anti-fibrotic Macrophage Response to Endotoxin (LPS)”

Louise Hecker University of Arizona, “Impaired Myofibroblast Dedifferentiation Contributes to Non-Resolving Fibrosis in Aging”

Renea Jabolonski Northwestern University Feinberg School of Medicine, “SIRT3, the Anti-Aging Major Mitochondrial Deacetylase, is Important for Preventing Pulmonary Fibrosis.”



DONATIONS power all of the Pulmonary Fibrosis Foundation's (PFF) accomplishments. Every dollar given to the PFF increases awareness of pulmonary fibrosis (PF), provides resources and support to those living with the disease, and expands the research that improves outcomes for patients.



TERENCE HALES

NYC RUN-WALK-HIKE FOR PULMONARY FIBROSIS

The 9th Annual NYC Run-Walk-Hike for Pulmonary Fibrosis was held in June in New York City's Central Park.

"As the son of someone who was diagnosed with PF, and who knows the horrible shock and pain of the disease, I am committed to helping find a cure," said event leader Terence Hales.

The Annual NYC Run-Walk-Hike for Pulmonary Fibrosis has raised more than half a million dollars for PF research since its inception.

Terence is a member of the PFF's board of directors. His father, Thomas Hales, is an emeritus member.



EMILY

ONE PAGE AT A TIME

Nine-year-old Emily decided to join the fight for a future without PF by creating her own fundraiser. Emily chose to read for her event because it's something she struggles with and it reminds her of what her grandma, who has PF, experiences every day: struggling to breathe.

"You just gotta stay strong and have hope," Emily said. "There're so many people out there losing lives because of this. Give back. You have to give back."

WAYS TO GIVE

Please consider supporting the mission of the PFF. Your donations to the PFF enhance patient support services, increase research funding, and broaden disease education outreach.

MAKE A DONATION

Online Visit pulmonaryfibrosis.org/donate

By Mail Use the donation envelope in this issue

By Phone Call us at 888.733.6741

PLAN A TEAM PFF FUNDRAISING EVENT

Team PFF Event Leaders are individuals who have created fundraising events to honor a friend or a loved one. To learn more about Team PFF, visit pulmonaryfibrosis.org/teampff.

PLANNED GIVING

Please think about including the Pulmonary Fibrosis Foundation in your estate plan and leave a legacy that will live on in the PF community. To get started, contact your financial advisor.

MONTHLY GIVING

Our generous monthly donors provide stability and support for our important programs. To learn more, visit us at pulmonaryfibrosis.org/pledgefaqs.

EMPLOYER MATCHING GIFTS

Please consider maximizing your donation through an employer matching gift program. To find out whether your employer will match your gifts, visit pulmonaryfibrosis.org/employermatch.

SPONSORSHIPS

The PFF offers a number of opportunities to sponsor the Foundation and specific events. Custom opportunities are available. To learn more, visit pulmonaryfibrosis.org/ways-to-give/sponsor-an-event.

Your donations have a positive and lasting impact on the PF community. The Foundation thanks you for your generosity and support.

Broadway Belts for PFF!

Attendees spent an evening with the stars in New York City at this signature event showcasing Broadway's best performers.

A NIGHT OF STARS

Key members of the PF community and Broadway's finest united for a unique evening of exceptional entertainment during the sixth annual *Broadway Belts for PFF!* Grossing approximately \$180,000 in one night, proceeds benefit the Pulmonary Fibrosis Foundation's Michael Kuchwara Fund for Idiopathic Pulmonary Fibrosis Research, Education, and Advocacy.

Held at New York's historic Edison Theatre, award-winning actress and dedicated PFF advocate Julie Halston hosted the event for the sixth time with her incomparable humor and skillful narration. She most recently appeared in the hit Tony-nominated Broadway

show *You Can't Take it With You*.

Broadway stars showcased their belting talents during an evening of melody under the musical direction of Christopher McGovern and returning director Carl Andress, who has directed all six of the *Broadway Belts for PFF!* performances. The star-studded cast included Tony Award winner Randy Graff (*City of Angels*), Cady Huffman (*The Producers*), Judy Kaye (*Phantom of the Opera*, *Nice Work If You Can Get It*), Grammy Award winner Daveed Diggs (*Hamilton*), Telly Leung (*Allegiance*), Robert Creighton (*The Mystery of Edwin Drood*), Lisa Howard (*It Should Have Been You*), Margo Seibert (*Rocky*) and Hunter Ryan Herdlicka (*Once Upon a Mattress*).



Daveed Diggs performing his original rap titled "Breathe."

HAMILTON STAR'S PULMONARY FIBROSIS RAP

Actor and rapper Daveed Diggs, who won both a Grammy and Emmy for his roles as Thomas Jefferson and Marquis de Lafayette in the hit musical *Hamilton*, debuted a special rap during *Broadway Belts for PFF!* 2016. The rap, titled "Breathe," was inspired by the PF community and the mission of the PFF.



From the left: Lisa Howard, Zakiya Young, Telly Leung, and Ryan Herdlicka



Robert Creighton leading an energetic performance.

PULMONARY FIBROSIS AWARENESS MONTH

We let the world know! September is Pulmonary Fibrosis Awareness Month. Members of the global PF community celebrated and raised awareness in a variety of ways.

Letting the World Know

PFF SUPPORTERS GET CREATIVE

The PFF's awareness initiative, Pulmonary Fibrosis Awareness Month, provides members of the PF community with an opportunity to create their own unique grassroots campaigns and fundraising events. Participants in 2015 went on to golf, run, walk, bowl, cook, and much more to raise awareness and funds for the PFF. Visitors to pulmonaryfibrosis.org found links to information about physician and patient educational events, and tools to help them achieve their fundraising goals.

Alongside fundraising events, supporters participated in a variety of ways, including by

- Asking their elected officials to officially declare the month of September Pulmonary Fibrosis Awareness Month
- Starting fundraisers
- Joining the #BlueUp4PF social media campaign

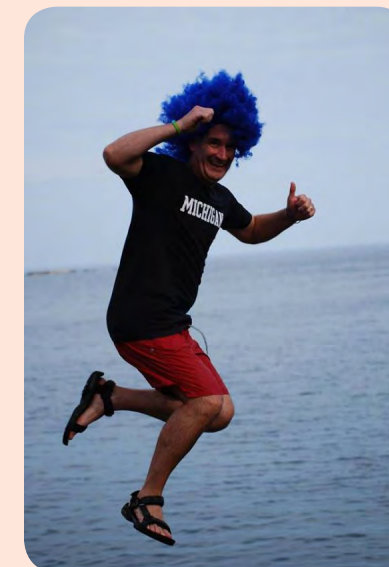
- Writing letters to the editor
- Personalizing their social media pages with custom PFF timeline photos and profile pictures



Nick Khan participated in #BlueUp4PF during Pulmonary Fibrosis Awareness Month in memory of his dad.



PFF Ambassadors went to Washington, DC during Pulmonary Fibrosis Awareness Month to participate in the Rally for Medical Research and ask for increased NIH funding.



Kevin R. Flaherty, MD, MS, Steering Committee Chairman, PFF Care Center Network and PFF Patient Registry, wearing a blue wig and jumping into Lake Michigan to support PF awareness.



PF Warrior Linda and her family sporting blue during a homecoming event.

PULMONARY FIBROSIS
awareness month
SEPTEMBER

PRESENTED BY Pulmonary Fibrosis
FOUNDATION

2015 PFF RESEARCH FUND AWARDS

The Pulmonary Fibrosis Foundation (PFF) places enormous importance on creating an environment that will assist in the development of effective treatments for pulmonary fibrosis (PF). We are developing this environment by directly funding research, promoting advocacy efforts, encouraging collaborative relations between industry and academic researchers, delivering key communications to patients, and developing solutions to bridge existing gaps in PF research.

The I.M. Rosenzweig Junior Investigator Awards and the Albert Rose Established Investigator Awards support projects that offer a high likelihood of improving the understanding of pulmonary fibrosis (PF) in the following areas: basic science, translational research, clinical medicine/research, and social science/quality of life.

I.M. ROSENZWEIG JUNIOR INVESTIGATOR AWARDS

Jose D. Herazo-Maya, MD - Yale University

“Serum microRNA Expression Profiles as Biomarkers in Idiopathic Pulmonary Fibrosis”

Funded by Genentech

Jonathan Kropski, MD - Vanderbilt University

“RTEL1 and DNA Damage Signaling in Pulmonary Fibrosis”

Funded by the NYC Run-Walk-Hike for Pulmonary Fibrosis

ALBERT ROSE ESTABLISHED INVESTIGATOR AWARDS

Tracy Luckhardt, MD, MS - University of Alabama, Birmingham

“Frailty as an Outcome Measure in Idiopathic Pulmonary Fibrosis”

Funded by a grant from Boehringer Ingelheim Pharmaceuticals, Inc.

Hon Yuen, PhD, MS - University of Alabama, Birmingham

“Home-Based Pulmonary Rehab for Patients with Pulmonary Fibrosis”

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to our
donors

\$500,000 and above

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 Stephen & Mary Beth Verget
 Kathleen Vesey
 Veterans United
 Anna Viegas
 Viele, Solimano, Swagler & Chapman, CPA
 PC
 Christine L. Vigeland, MD
 Vineland Chapter #84 OES
 Vivian Virden
 Craig & Melanie Vitrano
 Andrew R. Vranic, MD

Robert & Debra Wacker
 Rajat Walia, MD
 Tarik D. Walker, MD
 Michael Walker
 The Walking Club and Mall Walkers
 Jon Walkley
 Marie Walsh
 Eugene Walter
 Robert Warchol
 Sarah H. Warren
 Wartner Family
 Sharon Waters
 Marla Watson
 Julie Wattenbarger
 Jeri Webb
 Sam Weigt, MD
 Peter N. Weil
 Sarah Wells
 Jane Wesner
 Lewis Wesselius, MD
 Mary Westberg
 John-Michael & Linnea Westerberg
 David White
 Leona White
 Wick and Brother, Inc.
 Kesh Narayanan & Annelie Wilde
 Terry & Kim Marie Wilkins
 Peter Williams
 Delbert Williamson
 Carolyn Wilson
 Tiffany Winstone, MD
 Whitney Winter, PhD
 Elizabeth Wiseman
 Kathleen Wittau
 Ed Wolski, MD
 Wong Family
 The Esther Wright-Alsup Family
 Xiaoping Wu, MD
 Wim A. Wuyts, MD, PhD
 Larry & Colette Young
 Guoying Yu, PhD
 Ana C. Zamora, MD
 Betty & Lisa Zayas Family
 Patty Zemke
 Yingze Zhang, PhD
 Karen Ziff
 Ken & Roxann Zorowski
 Michael Zucker
 Kate Zuckerman

thank you to our Team PFF volunteers

Dedicated Supporters

MAKING AN IMPACT

Team PFF event leaders are advocates who actively fundraise, promote, and host third-party events. Between July 1, 2015 and June 30, 2016, Team PFF event leaders raised more than \$450,000 in support of the PFF's mission.



Participants of a Team PFF event, Maureen's Memorial Mile Run, strike a pose.

2015/2016 TEAM PFF

- 3rd Annual James Lynam Memorial 5K
- 3rd Annual Kickin It Full Throttle for A Cure Classic Car Show Benefiting the PFF
- 3rd Irv Feldman Texas Hold' Em Tournament & Casino Night
- 5th Annual John F Tighe Walk for Pulmonary Fibrosis
- 5th Annual Vic Vittorino Walk for your Next Breath 5K
- 6th Annual Cruisin' Toward a Cure for PF
- 6th Annual Pete DeVito Memorial Golf Outing
- 7th Annual Garden State 5K
- 9th Annual Greg Chandler & Guy F. Solimano Memorial Golf Tournament
- Art Shirk Birthday
- BAF Memorial Bike Run/Picnic
- Bear Down for a Breath of Hope
- Bill Camp's 70th Birthday
- Bourbon & Barbeque
- Bowling for Breath
- Bowling for Pulmonary Fibrosis in Memory of Vincenzo Ciavaglia
- Bracelets for PFF by Jools Dunn
- Breath - Taking, Pulmonary Fibrosis Foundation Craft & Bake Sale
- Breathe and Live
- Breathe Easy Visual Arts Competition and Exhibition
- Breathe for Duffy
- Breathe Fundraiser
- Breathtaking Jewelry for a Cause for IPF!
- Katie Runs for Her Dad
- Caroline M Fell Memorial Walk
- Celebration of Sgt. Vern T Darlington's Life
- Change Makers Coin Jar Game
- Charity Golf Challenge
- Charlotte Saunders Transplaniversary
- Coffee/Donut Social for PFF
- Dance Performance
- Debbie Is Amazing
- Divy and Joanne's Wedding
- Donna Schwartz Memorial Walk
- Egg Toss in Memory of Stephen Epstein
- First Midwest Bank Jeans Day
- Flipping Pages for Pulmonary Fibrosis
- Free to Breathe
- Golf Scramble for Global PF Awareness Month
- Harold W. Elbert Memorial and Research Fund
- Hodgson Russ Dress Down Day
- Breathe Free, Live Free
- Hudson Golf Tournament
- IL Credit Union Jeans Day
- Innophos, Inc. Jean Day
- Ironman Chattanooga 140.6 Triathlon
- Jack up the WOD Against Pulmonary Fibrosis
- James Dylan and Amy's Blue It Up for Pulmonary Fibrosis Awareness
- Jennifer Wade Senior Project
- Jim Cormier Canal Walk
- John Deere Golf Classic
- Judges/Scheifele Wedding
- Keeping the Radiant Light of Jules Perez-Montiforte Alive (PartyLite)
- Kimberly Cole Afghan Fundraiser
- Knock Your Pulmonary Fibrosis Pants Off!
- Marree Paulson Memorial Fundraiser, Thirty One Party
- Mother Daughter Photo Shoot
- Mowery Clinic Jeans Day
- Nancy Bacon Walk
- Newtown Elementary School Dress Down Day
- NYC Run-Walk-Hike
- Octoberfest Half Marathon, 5K, 10K and 40K Bike
- Order of Eastern Star – Grand Chapter of Washington -Vineland #84 Coin Drill Donation
- PartyLite Fundraiser
- Pennies from Heaven
- Performance Matters Associates Fundraiser
- PFF Chipotle Fundraiser
- Philadelphia Flyers Pulmonary Fibrosis Awareness Night
- Pilot for A Cure
- Points for Pulmonary Fibrosis
- POP for PFF Walking Relay
- Psych Out Pulmonary Fibrosis
- Pulmonary Fibrosis Fundraiser with Jamberry!
- Pulmonary Fibrosis T-shirt Campaign
- Quad City Marathon
- Ready.Set.Breathe!
- Richmond Marathon
- Ride with Wave
- Robin Dellinger Yard Sale for PFF
- Rock for a Cure
- Seidensticker Cup
- Shop LaLuRoe for Pulmonary Fibrosis
- #ShowOut for Sharon
- Silver Falls Half Marathon in honor of Robert Lager
- Skate for Stephen
- Songs for Cindy at Shank's Tavern
- Spectrum Health Jeans Day
- Stephan Lively Fundraiser
- Stick it to IPF
- Strikeout Pulmonary Fibrosis
- T1 10 Mile and 10K
- Teal Spin Craft Sales

2015/2016 TEAM PFF (CONT'D)

- Team Jeffery Trivia Night
- Team Joyce Challenge
- The Breathe Easy Fundathon
- The Clarence Friar Memorial Ready-Set-Breathe 5k and Fun Run
- The Newport and Walt Disney World Marathons
- Thomas Charles Sanders Fundraiser
- Trails for Charities Day
- Walgreens CT Annual Golf Charity
- Wanda James Golf Event
- West Virginia University Bake Sale
- Westphal Golf Outing
- Wine Down for Pulmonary Fibrosis
- Zielke Wedding Anniversary

TWO CAMPAIGNS, ONE GOAL

When Sharon Cirilli passed away from pulmonary fibrosis in July 2015, her daughter, Ashley M. Cirilli, and daughter-in-law, Ashley E. Cirilli, wanted to do something to honor her memory. Sharon was a prolific crocheter and an accomplished seamstress, and Ashley E., a needlework enthusiast herself, sells Sharon's and her own work at craft shows, with proceeds benefiting the Pulmonary Fibrosis Foundation. Ashley M. started a "Show Out for Sharon" campaign, which entailed mailing out hundreds of PFF bracelets and asking friends and family to share photos of themselves wearing the bracelets on social media. "We ended up getting 250 photos," Ashley M. says. "For each one, the Cirilli family donated \$5 and my company, Disney, matched the contributions. It was a great way to keep Mom's memory going and raise awareness of PF." The Ashleys have contributed more than \$12,000 to the PFF. "Even if it's only \$25 at a time, we know we're helping someone else and remembering an amazing person," Ashley E. says.



FINANCIAL SUMMARY: FY2015-2016

STATEMENT OF FINANCIAL POSITION

AUDITED NUMBERS

Total Assets	\$	6,743,740
Total Liabilities		2,404,432
<hr/>		
Total Net Assets	\$	4,339,308

**STATEMENT OF ACTIVITIES:
YEAR ENDED JUNE 30, 2016**

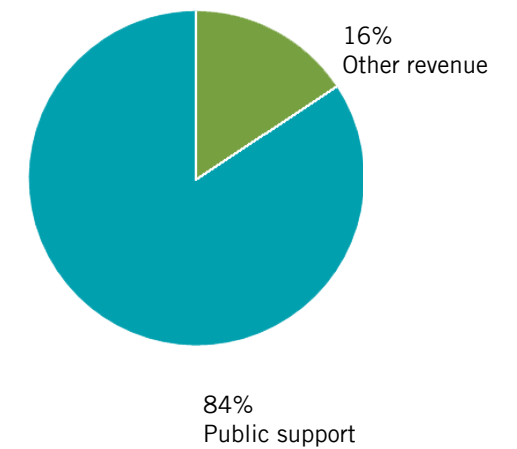
AUDITED NUMBERS

Total Revenue		
Unrestricted	\$	6,897,595
Temporarily Restricted		736,527
<hr/>		
Total Expenses	\$	7,634,122
<hr/>		
Change in Net Assets	\$	745,953

SOURCES OF REVENUE

Public Support	\$	6,434,748	84%
Other Revenue		1,199,374	16%
<hr/>			
Total Revenue	\$	7,634,122	100%

SOURCES OF REVENUE

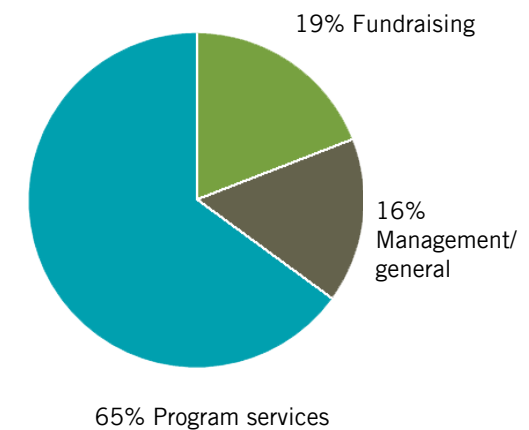


ALLOCATION OF EXPENSES

AUDITED NUMBERS

Program Services by Category			
PFF Patient Registry	\$	809,845	
Care Center Network		280,866	
Research		277,121	
Outreach and Awareness		269,585	
Program Support	1,101,641		
Patient Communication Center		38,790	
Physician and Patient Education		636,682	
PFF Summit		699,806	
Support Groups		345,110	
Legislative Advocacy		12,784	
<hr/>			
Program Services Total	\$	4,472,230	65%
Management and General		1,106,218	16%
Fundraising		1,309,721	19%
<hr/>			
Total FY2015-2016 Expenses	\$	6,888,169	100%

ALLOCATION OF EXPENSES



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The Pulmonary Fibrosis Foundation
has a four-star rating from Charity Navigator
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Sean Thinnis

230 EAST OHIO STREET, SUITE 500
CHICAGO, ILLINOIS 60611-33201

www.pulmonaryfibrosis.org