I was diagnosed with Young-Onset Parkinson’s disease in October 2018. I was only 33 years old. My two daughters were 2 and 6 years old. I had symptoms for more than a decade, starting with an obvious tremor in my right foot that developed into a severe limp. I slowly started not being able to do some basic things without difficulty, like zipping up my daughters’ jackets. An online search of Parkinson’s symptoms cannot explain how unpredictable this disease can be and why some days you feel like a superhero, and why others you struggle to get your shoes on. A search won’t tell you that after being diagnosed with this disease, you will find a community, like the Parkinson’s Foundation; nor will it tell you that you will find new friends who will zip up your daughters’ jackets when they know your hands can’t do it themselves.

The Parkinson’s Foundation has been reliable, encouraging and incredible. They love us like family. They embrace not just me, but my husband and two daughters. I’m thankful and feel that if anyone is struggling with their recent diagnosis or life with Parkinson’s, rely on the Foundation because you will find a home there.

Christina Korines is living with Parkinson’s. She gave the keynote speech at the Parkinson’s Foundation 2019 New York Gala. This excerpt is part of her speech that inspires us to do more.
MESSAGE FROM THE CEO

Together, we expanded our reach in 2019, making life better for our Parkinson’s disease (PD) community by developing our care, research and education efforts. We impacted more people than ever before — launching exciting new research initiatives, reaching the underserved and closing the information gap for the 60,000 Americans newly diagnosed this year.

In 2019, we launched two new initiatives that will significantly impact the lives of people living with Parkinson’s:

• **PD GENEration: Mapping the Future of Parkinson’s Disease** is a national initiative that offers free genetic testing for clinically relevant PD-related genes and free genetic counseling. By contributing genetic testing results to this flagship program, we have the opportunity to help accelerate scientific research to improve our understanding of PD and potentially identify new and/or better treatment options.

• **Newly Diagnosed: Building a Better Life with Parkinson’s Disease** is a program that connects those newly diagnosed with the right information and resources to live better with PD. We want them to know that they are not alone in this journey and that we are here to help.

Last year, we honored former Pennsylvania Governor Edward G. Rendell for his contributions to the Parkinson’s community. His words inspire us to further our reach. “Can I say that I am going to defeat Parkinson’s? Not yet,” Governor Rendell said. “Someday soon maybe people will be able to say that. Can I say I’m going to fight this as long as I can? Absolutely.”

Our 2020 goals are designed to inspire optimism as we continue to support our international community. The individual donors, foundations and corporations in this Annual Report help us change lives every day and get us closer to the promise of a cure. Thank you for your incredible support in 2019 and in the new decade. Together, we are a powerful force in the fight against Parkinson’s disease!

Thank you,

John L. Lehr  
President and Chief Executive Officer

Howard D. Morgan  
Chairman of the Board
Expert care is critical to living better with Parkinson’s. It can improve quality of life and lower the risk of complications, which is why we are committed to make expert care more accessible. We identify, support and connect people living with Parkinson’s with premier treatment and research centers worldwide.

We designate Parkinson’s Foundation Centers of Excellence — the largest network of Parkinson’s clinics. These centers house teams specialized with deep expertise in the latest PD medications, therapies and research that altogether treat more than 193,500 people with Parkinson’s.

In the last two years, we added six Centers of Excellence to our global network of 47 leading medical centers. Each recently designated center brings us to new, underserved PD communities in: Colorado, Indiana, Pennsylvania, Ohio, South Carolina and Iowa.

As the prevalence of Parkinson’s is expected to increase with our aging population, we work to ensure healthcare professionals are better prepared to deliver expert PD care. Together, we are improving specialized professional training — from in-person sessions to online courses — created for nurses, physical therapists and other professionals devoted to delivering the latest in proven PD care techniques. In 2019, we launched a new online nurse course that provides essential educational tools for nurses to deliver optimal care through all stages of PD.

We can all play a role as Parkinson’s advocates to change the way people with Parkinson’s receive care in hospitals, which is why the Parkinson’s Foundation created the Aware in Care Ambassador Program. This volunteer group distributes hospital kits that bolster best practices in treating people with Parkinson’s to patients and healthcare providers. Armed with resources and the latest information, 81 ambassadors across 30 states actively help us educate hospital staff and community members nationwide to make life better for people with Parkinson’s.

Find out how we are expanding our reach in care at Parkinson.org/ExpertCare.
REACHING THE NEWLY DIAGNOSED

To help provide better outcomes for the 60,000 people who are newly diagnosed each year in the U.S., the Parkinson’s Foundation recently launched Newly Diagnosed: Building a Better Life with Parkinson’s Disease. This new campaign is designed to close the gap between diagnosis and utilizing the right information and resources to live better with PD.

Learn more at Parkinson.org/NewlyDiagnosed.

“I began doing advocacy work for Parkinson’s disease two years after I was diagnosed. I have been a Research Advocate with the Parkinson’s Foundation for 11 years, served on the advisory committee for three years and now also serve as an Aware in Care Ambassador. Though PD ended my career as a scientist, my association with the Parkinson’s Foundation opened a lot of new doors for me.”

— Girija Muralidhar, living with Parkinson’s

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Dan Chihos
This is the place for answers about Parkinson’s. Have you ordered your free Aware In Care Kit from the Foundation? It is free of charge and will make a huge difference if and when you have to stay in the hospital. Easy thing to do!
The Parkinson’s Foundation knows that a research breakthrough can happen at any time and in any lab, which is why we invested $12.2 million this year to explore what causes Parkinson’s, how to treat it and how to find a cure.

In 2019, we funded four institutions that will each receive $2 million to design and launch Parkinson’s-specific research studies over the next four years. These Parkinson’s Foundation Research Centers aim to drive innovative developments and advance Parkinson’s research at an even faster rate. The four prestigious institutions designated as Research Centers include: Columbia University Medical Center; University of Florida in collaboration with Emory University; University of Michigan in collaboration with The University of Texas Southwestern Medical Center; and Yale School of Medicine.

We have supported more than 550 scientists through grants that keep researchers and doctors in the Parkinson’s research field. In 2019, we funded $4.2 million across 46 research grants to advance promising research in labs across the world. We support early-career and established investigators as they uncover new biological and clinical insights to deliver improved Parkinson’s therapies.

Our knowledge of Parkinson’s has changed significantly, so has our understanding of how many people have this disease through our Parkinson’s Prevalence Project. This study established the most accurate number of people with Parkinson’s in the U.S., showing that 1.2 million people will be newly diagnosed with Parkinson’s by 2030.
REACHING MORE PEOPLE THROUGH GENETICS

Our understanding of the role genetics plays in Parkinson’s has dramatically changed in the past 20 years. As we progress towards precision medicine in Parkinson’s, the Foundation launched a new study called **PD GENEration: Mapping the Future of Parkinson’s Disease**. The first-of-its-kind national initiative offers free genetic testing and counseling to accelerate enrollment in clinical trials. Between 10 and 15 percent of participants can expect to find out if they have a genetic risk factor for Parkinson’s. Results will also be accessible in the future to help us develop improved treatments.

The Parkinson’s Foundation supports promising research to help drive change and make life better for people with Parkinson’s. Since 1957, we have funded more than $353 million in groundbreaking research, transforming the way we diagnose and treat Parkinson’s.

“*I’m planning to become part of the PD GENEration study to get a better understanding of my disease, and to contribute to research that will improve future care.*”

— Anne Hall, living with Parkinson’s

**Yulan Xiong, PhD**, Parkinson’s Foundation Stanley Fahn Junior Faculty Awardee, studies one of the primary genetic mutations tied to Parkinson’s at Kansas State University.

Natasha Ratcliffe
You guys are doing such great things for #parkinsonsresearch. Grateful to be working closely together and learning from you! 😊
Looking forward to seeing this work develop #patientinvolvement.

Learn about how we are reaching the next level of research at Parkinson.org/Research.
To reach more people in our global community, we believe in evolving the way we connect and address the unmet needs for those who are underserved.

Women with Parkinson’s experience disparities in care and treatment when compared to men, which is why we created the first national initiative for women with Parkinson’s. Together, we worked with more than 50 women across the country, including those living with PD and neurologists, to develop the first “Women and PD” research and care agenda.

Strengthening our local impact furthers our reach. In 2019, we expanded our community grants program across the country. Putting our mission in action, we awarded $1.5 million that supported 118 life-changing programs and classes — from music therapy and non-contact boxing classes to education symposiums, support groups and research studies — across local Parkinson’s communities.
We also expanded our bold reach on a local level by holding 65 educational events across the nation. With the help of our chapters and Centers of Excellence, we hosted interactive sessions led by a Parkinson’s Foundation representative and PD experts that addressed issues such as: New Frontiers in Research and Care, Women and PD: Closing the Gender Gap, On the Menu: Nutrition in Parkinson’s and more. Participants leave each event with valuable resources and knowledge they can share with their community.

We believe in adapting our resources, tools and educational events to better reach everyone, regardless of where they are in their PD journey — from our newly diagnosed kit to our volunteer and care partner summits. We look forward to connecting with our community, be it a sponsored event, funded class or an online webinar.

“Wow, I was truly blown away by the Helpline specialist I spoke to. She was totally professional, yet warm and personable. I immediately felt a comfort level and sense of trust. I have to say after dealing with the challenges of PD for almost 12 years, I wish I had called years ago and availed myself of this incredible resource.”

— Andrée Jannette, Helpline caller

Help us reach new levels of Parkinson’s awareness at Parkinson.org/GetInvolved.
2019 IN NUMBERS

$353 MILLION+
invested in Parkinson’s research and clinical care since 1957

550+
scientists received Parkinson’s Foundation funding since 2010

$12.8 MILLION
supported Parkinson’s Research Centers and early-career scientists in 2019

74,000
health professionals completed professional education courses

193,500
people with Parkinson’s treated in Centers of Excellence — the largest global network of Parkinson’s clinics
14,550 people joined an Expert Briefings educational webinar

79,000 podcast episodes downloaded

4.1 MILLION people visited Parkinson.org, the go-to online Parkinson’s resource

$1.5 MILLION awarded to 118 community grants across 37 states

212,338 resources mailed through the Helpline in 2019

91,413 free, life-saving Aware in Care kits distributed through our Helpline and local chapters to help people get better care in a hospital

127,000 people received vital counseling from specialists on our toll-free Helpline, 1-800-4PD-INFO
The Parkinson’s Foundation gratefully acknowledges all our supporters who donated $2,500 or more between July 1, 2018 and June 30, 2019. Our generous donors allow us to further our mission to make lives better for people with Parkinson’s and advance research toward a cure. We thank them for their dedicated support.

### Honor Roll 2019

#### $250,000 AND ABOVE
- ACADIA Pharmaceuticals, Inc.
- Estate of Edwin F. Armstrong
- Estate of Carolyn Flowers Brantley
- Estate of William J. DeVirgilio
- Edmond J. Safra Philanthropic Foundation
- Estate of Catherine Hinterbuchner
- Johnson & Johnson Pharmaceutical Research & Development
- Mertens Family Revocable Living Trust
- Jean Myers
- Dominick and Mary Rutigliano Revocable Trust
- Sharron and Joseph Ashby Hubert Fund of the Community Foundation of Broward
- Estate of Nette M. Swinney
- US WorldMeds, LLC

#### $100,000 TO $249,999
- Amodor Therapeutics, Inc.
- Amneal Pharmaceuticals, LLC
- Tom and Marcia Mondavi Borger Family
- Boston Scientific Corporation
- Estate of Thomas Brown
- Estate of Roseann B. Comstock
- Alberto and Lourdes Dosal Fairchild Martindale Foundation
- Richard and Schuyler Field
- Richard and Shelley Friedland Global Kinetics Corporation
- Estate of Michael D. Hirsch
- Michael Inden Living Trust
- Estate of Maurice Laboz
- Esther D. Lewis Living Trust
- Light of Day Foundation, Inc.
- Lundbeck, LLC
- Delia McCullough
- Medtronic, Inc.
- Sandra Salka Milken
- Paul and Marcy Nathan
- Parkinson’s Unity Walk, Inc.
- Partners Healthcare System
- The Patient Centered Outcomes Research Institute (PCORI)
- Estate of Ann M. Reynolds Right at Home, LLC
- Estate of Michael J. Rinaldi, Jr.
- Sunovion Pharmaceuticals, Inc.
- Roselle Taylor
- Estate of Virginia Thompson
- UCB, Inc.
- Vertical Pharmaceuticals, LLC
- Estate of Grace Westcott
- Estate of Lena Willis
- Steve and Cindy Wolfe Family
- World Parkinson Coalition, Inc.
- The Wrobel Family Foundation

#### $50,000 TO $99,999
- Abbott
- Adamas Pharmaceuticals, Inc.
- The Batchelor Foundation, Inc.
- Estate of Lulu J. Bates
- Estate of Dr. Bearce Batson
- Mr. and Mrs. J. Gordon and Sully Beckham
- The Beneficly Community Impact Fund
- Estate of Frances Blank
- James Bobo
- CIGNA Foundation
- Community Health Charities
- Estate of Mary Lou Donley
- Estate of Peter J. Dorn
- Estate of Dorothy June Emig
- The Don A. Hunziker Memorial Foundation, Inc.
- The Eleanor M. and Herbert D. Katz Family Foundation, Inc.
- Trust of James Kolling
- Estate of Arnold Levin
- Margaret D. Lienemann Charitable Trust
- Estate of Louise Mazza
- Estate of Eileen Messina
- Charlotte and Jim Monhart
- Estate of Shirley June Muise
- Estate of Harriet M. Pennekamp
- Estate of Carol C. Price
- Estate of Barbara B. Riggi
- Roche TCRC, Inc.
- Estate of Elaine K. Shelton
- Estate of Royce E. Shetson
- Dr. Lou Sherfesee, IV
- The Smith Family Foundation
- The GMK Family Fund
- Cynthia Vannec
- The Wasyly Family Foundation

#### $25,000 TO $49,999
- Abbott Laboratories
- AbbVie, Inc.
- Trust of Suzanne Aganian
- Mona and Andy Albert
- The Applebaum Foundation, Inc.
- Dr. Janet W. Bay
- The Jane H. Berktold Parkinson Fund
- Terri Springer
- Lester W. Stevenson, Jr. Trust
- Takeda Pharmaceuticals USA
- The David Tepper Charitable Foundation
- Joan H. and Preston R. Tisch Fund
- Leon E. and Patricia M. Westbrook Family Charitable Trust
- Linda and Robert P. Fetch
- Fondazione Fresco Institute Italia
- Gerald Z. Gibian
- Mrs. Stephanie Goldman-Rosen and Mr. Steven Rosen
- Estate of Willard L. Groenewoud
- Michael and Gita Kinney
- LendingTree Foundation
- Arlene and Jerome Levine
- Loeb & Loeb, LLP
- Mindy Mclroy
- Gail and Robert Milhous
- Moelis & Company
- Howard and Ginger Morgan
- Network For Good
- Neurocrine Biosciences, Inc.
- OhioHealth
- The Oreffice Foundation
- Parker Hannifin Corporation
- Bill and Marlene Perdan
- The Donna Prendergast Trust
- Patricia Schreer
- Anne J. Sherr
- Solomon Family Foundation
- Sherwin and Sheri Zuckerman

#### $10,000 TO $24,999
- 23andMe
- Jeni E. Abernathy
- Stephen and Marsha Ackerman
- Craig and Darla Albert
- Kathy Albert
- Mr. and Mrs. Thomas S. Alexander
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- AR Global Investments, LLC
- The Avery Family Charitable Fund
- The Beckham Family Charitable Fund
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- Arthur W. Bine
- Biogen
- Blue Cross and Blue Shield of Kansas City
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- Bonner Family Private Foundation, Inc.
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- Dr. Karen E. Burke and Mr. Peter John Goulardis
- Constatte Foundation
- Robert Baker Carlson Trust
- CGL Financial Fund
- Ex Investments, Inc.
- Robert Baker Carlson Trust
- CGL Financial Fund
- EX Investments, Inc.
- Louis & Virginia Clemente Foundation, Inc.
- John P. Clulow Fund
- Robert and Joanne Crown Income Charitable Fund
- Sherwin and Sheri Zuckerman

### World Parkinson Congress in Japan

In June 2019, the Parkinson’s Foundation attended and sponsored the 5th World Parkinson Congress in Kyoto, Japan. The Foundation shared eight research studies with our global Parkinson’s community that highlighted our women and PD initiative, the importance of patient engagement, genetics and the importance of Spanish-language programming.
Beverly Adams is a proud grandmother who lives in Dunwoody, GA, and was diagnosed with Parkinson’s in 2019. She began exercising in the “Living Well with Parkinson’s” program at the Marcus Jewish Community Center of Atlanta, a Parkinson’s Foundation community grant recipient.
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The Erickson Family Charitable Fund
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Revocable Living Trust
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John Mead
Medical Professional Clinical Research Center, Inc.
Susan and Bill Medlin
Estate of Frederick H. Meeder, Jr.
Louis Messer & Helen Messer Charitable Foundation
PARKINSON’S FOUNDATION HOSTS
NIGHT AT THE MARKET
Celebrity Chefs take a bow at Night at the Market on October 23, 2019, at the Lincoln Eatery in Miami, FL. From left to right: John Mooney, Aarón Sánchez, James Tahhan, Sue Torres, Stephen Pyles, Michael Schwartz, Richard Holes.
Find an event near you at Parkinson.org/Events.
Foundation Hosts First-Ever Medical Marijuana and Parkinson’s Conference

The Parkinson’s Foundation hosted its first-ever conference focused on medical marijuana and Parkinson’s in Denver, CO, in March 2019. The Foundation will publish suggested practices and areas for further research on using medical marijuana as a treatment in 2020.

Learn more at Parkinson.org/Marijuana.
Family, fitness and finding a cure for Parkinson’s fuel Kathy Jordon. Newly diagnosed, with children in high school and college, the mother of four immediately immersed herself in her new PD community.

First, she first learned about the disease, reaching out to the Parkinson’s Foundation and attending local support groups and exercise classes. She then joined the Moving Day Southeastern Pennsylvania Planning Committee, contributing her past fundraising and event planning expertise.

Kathy’s first Moving Day was a success, which she credits to her strong support system. Though family and friends had only recently learned of her diagnosis, they were at Moving Day manning tables, volunteering, serving as an event photographer and walking. “They were unbelievable, not just my husband, John, and four children, but my siblings and neighbors,” Kathy said. “They all told me, ‘We’re doing it again next year.’"

Kathy’s team raised more than $5,000. "People just want to help. Moving Day is a way for them to jump in," Kathy said.

Join us at Moving Day. Sign up today at MovingDayWalk.org.
### TOP 10 MOVING DAY TEAMS

<table>
<thead>
<tr>
<th>Team Name</th>
<th>Event City</th>
<th>Amount Raised in 2019 Season</th>
</tr>
</thead>
<tbody>
<tr>
<td>Larry’s Lemons</td>
<td>Twin Cities</td>
<td>$60,255</td>
</tr>
<tr>
<td>Berkowitz Pollack Brant/Provenance Wealth Advisors</td>
<td>Miami</td>
<td>$58,562</td>
</tr>
<tr>
<td>Rock Steady Boxing NC Triangle</td>
<td>NC Triangle</td>
<td>$44,020</td>
</tr>
<tr>
<td>Team Beaufort #IWNQ</td>
<td>Charleston</td>
<td>$43,885</td>
</tr>
<tr>
<td>Team Zuckerman</td>
<td>Chicago</td>
<td>$42,397</td>
</tr>
<tr>
<td>Edward Jones Heartland</td>
<td>Kansas City</td>
<td>$40,355</td>
</tr>
<tr>
<td>Team Albert</td>
<td>Chicago</td>
<td>$39,602</td>
</tr>
<tr>
<td>Pops’ Peeps</td>
<td>Tampa Bay</td>
<td>$39,162</td>
</tr>
<tr>
<td>23andMe</td>
<td>San Francisco</td>
<td>$34,611</td>
</tr>
<tr>
<td>Rock Steady Boxing Boston</td>
<td>Boston</td>
<td>$27,145</td>
</tr>
</tbody>
</table>

From an off-road vehicle race to a golf tournament, a marathon to a Facebook fundraiser — Parkinson’s Champions have raised nearly $9.4 million to make life better for people with Parkinson’s.

We would like to thank Parkinson’s Champions community fundraisers who raised $2,000 or more between July 1, 2018 and June 30, 2019. We appreciate the energy and dedication shown by all our Champions in helping us beat Parkinson’s.

FACEBOOK FUNDRAISERS

Alison Bailin Handler
Doug Bell
Francine Birbragher-Rozencwaig
Stephen Bittel
Risa Davis
Bethany Dowell Layne
Sara Finnerty Anton
Don Foster
Steph Giardino
Kendall Arntz Green
Michael Johnson
David Kaplansky
Jacob Koressel
Richard Kramer
Shona MacKenzie
Karen Marie Madrigale
Rebecca O’Reilly
Nicole Rocklin
Michael Samuels
Janet Scheer
Inderbir Singh Atwal
Leslie Milsten Thornton
Elizabeth Travis
Barbra Turitz Horner
Jann Vlah Holzman
Richard Winkles
Donna Zinnershine Goldslager

Dahlia Abadir
2018 New York City Marathon
Maris Ackerman
2018 Chicago Marathon
Ruby Aldridge
2018 New York City Marathon
Stephanie Andrews
2018 New York City Marathon
Dom Arioli
Knock Out PD Challenge
Jamie Billings & Tom Wigglesworth
2018 New York City Marathon
Kim Bova
2018 New York City Marathon
Peggy Brug
2018 New York City Marathon
Molly Burkhardt
2018 New York City Marathon
Moira Christoudias
2018 New York City Marathon
Greg Cockrell
Mel Cockrell Tribute
Katie Ann Connors
2018 New York City Marathon
Evon Cutler
Push Ups for Parkinson’s
Brenna DeVincentis & Mark Zak
2019 Chicago Marathon
Anna Duewiger
2018 Phoenix Marathon
Drew Eichelberger
Stomping the Pavement for Steve
Kirsten Elizabeth
2018 Tough Mudder Team
Flat Creek Baptist Church
The Chris Daunt 5K Run/Walk Over Parkinson’s
Jeannine Fleegle
2018 Marine Corps Marathon
Aaron & Anissa Fleisher
2018 Marine Corps Marathon
Kingsley Fuller
2018 New York City Marathon
Greg Gale
2018 New York City Marathon
Adrianna Garrett
2019 Chicago Marathon
Erin Gerrity
2018 Philadelphia Marathon
Katie Harryman
Heartland Chapter
Cocktails for a Cure
Beatriz Hernandez
Bill Hernandez Tribute
Michael Horning
2018 New York City Marathon
Kristie Hutchinson
2019 New York City Marathon
Dawn Judson
Shake, Rattle & Crawl 2019
Kristen Kanner
2018 New York City Marathon
Kristen Kolombatovich
2018 New York City Marathon
Cassie Kornblau
2018 Chicago Marathon
Bernadette Lindquist
Annual Garage Sale
Kelsey Lorence
2018 New York City Marathon
Glenn Lucas
2019 New York City Marathon
Lisa MacGregor
Barefoot Open for Parkinson’s
Kris Machnick
8 for 80 Climb
Jim Manders & Tim Thurn
PD450
Michael Mano
2018 New York City Marathon
Danielle Marcus
Salon Acai Holiday Party
Tony Mayes
2018 New York City Marathon
Richard L. McFadden
Spaghetti Dinner - in honor of Dennis Harris
Matt McKee
Long Live Dance Fundraiser
Amanda Mitchell
2018 New York City Marathon
Lisa Pagan
2018 New York City Marathon
Jenna Palek
2018 Philadelphia Marathon
Melissa Parks
2018 Marine Corps Marathon
Victoria Pekarski
2019 Chicago Marathon
Anna Petkovsek
2019 Chicago Marathon
Nancy Redkey
Walk Over Water
David Schwartz
Dave’s Ties for PD
Erin Stack
2018 New York City Marathon
Ruth Stilwell
2018 Marine Corps Marathon
Joyce Tracy
Zero Mile Fun Run
Elizabeth Travis
Twin Cities Marathon
Anne Wray
2018 New York City Marathon
David Young
When I’m 64 Concert
Allyson Zoller
2018 New York City Marathon

Become a Parkinson’s Champion at Parkinson.org/Champions.
The Parkinson’s Foundation recognizes and honors all Legacy Society members who support the Foundation in their wills, trusts, life income gifts, retirement plans and other planned gifts. Legacy Society members leave meaningful, lasting legacies for future generations.

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HOW SHIRLEY GORDON HELPED MOVE PARKINSON’S RESEARCH FORWARD

"If this gift will help people with Parkinson’s disease, I’m happy to help people not go through what I went through," Shirley Gordon told Karen Verrone, who went from Shirley’s financial advisor to loyal friend.

Shirley’s walking issues led her to the University of Rochester, a Parkinson’s Foundation Center of Excellence, where she was diagnosed and treated for Parkinson’s.

With the passing of her parents and later on her husband, Shirley became a savvy investor. Over time, she developed a giving plan. “100% of what she left behind went to charity, supporting the causes that left an impression on her life,” Karen said.

Above all, Shirley believed in research to help those dealing with health issues. She gave a generous gift to the Parkinson’s Foundation, along with the American Heart Association, cancer research and pediatric cardiac care. “Shirley was always very generous,” Karen said.

Over the last 10 years Shirley and Karen enjoyed weekly lunch together. “Every week, everyone in the restaurant would stop by to say hi to Shirley. It was the kind of person she was.”

Shirley’s memory will forever live on as a Parkinson’s Foundation Legacy Society member.

Learn more about the Legacy Society and planned giving options at Parkinson.org/PlannedGiving.
The Parkinson’s Foundation gratefully acknowledges those national corporate sponsors who actively support our mission critical programs and research initiatives.

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CORPORATE SPONSOR SUPPORTS CRITICAL FOUNDATION EVENTS

Jessica Daitch with Bill Keller, ACADIA Executive Director of Patient Advocacy & Industry Relations, and Becky Chow, ACADIA Senior Director of Consumer Marketing at the 2018 Caregiver Summit, an event presented by ACADIA. We thank ACADIA for their continued support in Moving Day, Centers of Excellence Leadership Conference and Care Partner Summit. With ACADIA’s support, the Parkinson’s Foundation can continue to raise Parkinson’s awareness and education efforts nationwide.

PARKINSON’S FOUNDATION HIGHLIGHTS

Neurologists attend the first national forum for women and Parkinson’s on October 19, 2018, in Houston, TX. More than 50 women worked to develop the first-ever patient-centered “Women and PD” research and care agenda.

Jon and Katy Romero along with guests enjoy Celebrate Spring New York on April 9, 2019, at PHD Rooftop Lounge at Dream Downtown.
Through generous donations, the Parkinson’s Foundation can make life better for everyone in the Parkinson’s community. In 2019, these four generous donors and board members helped us expand our reach even further.

**PONDER HARRISON**

“I joined the Parkinson’s Foundation board to demonstrate solidarity of commitment with my wife and father who were diagnosed with PD,” said Ponder Harrison. “I wanted them, along with our children, to know I am in this fight, right beside them.” In 2020, the Parkinson’s Foundation will host the Power over Parkinson’s gala with Ponder as event co-chair, honoring Dr. Harrison and Judy Harrison. Ponder serves as Senior Advisor to Comvest Partners and Franklin Resources’ Industrial Technology Fund.

**MARCIA MONDAVI BORGER**

“Parkinson’s disease is personal to me and my family,” said Marcia Mondavi Borger, who is living with Parkinson’s. “I am determined to help the Foundation raise awareness and funds to support its urgent mission of making life better for the 10 million people worldwide living with the disease today.” Marcia helps the Foundation keep early-career scientists focused on Parkinson’s and does so as a Board Member and as Leaders In Research co-chair. She first became involved with the Foundation through Moving Day San Francisco. Marcia is the daughter of wine legend Robert Mondavi and Partner of Continuum Estates Winery.

**JAMES F.T. MONHART**

“I was determined to get involved with an organization that shared my passion for finding a cure for the disease and finding ways to slow the progression of Parkinson’s symptoms,” said Jim Monhart, who is living with Parkinson’s and has served as a Parkinson’s Foundation Board Member. Jim is currently a co-chair of the Foundation’s Leaders in Research, which brings together donors with a passion for investing in Parkinson’s research and advancing the Foundation’s scientific agenda.

**WROBEL FAMILY FOUNDATION**

The Wrobel Family Foundation is “driven by a single goal, to do our part in making the world a better place by supporting organizations that are working to find a cure for Cancer, Alzheimer’s and Parkinson’s disease.” Florence Wrobel, President and founder, along with her brother and sister-in-law, believed in supporting research initiatives. Through their generous donation, the Wrobel Family Foundation supports both the Parkinson’s Foundation National Volunteer Leadership Summit and the Leaders in Research Symposium.

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Participants salsa dance at the Parkinson’s Foundation Spanish-language conference, Hacia Adelante: Navegando el mar del Parkinson, in Los Angeles, CA, on October 19, 2018.

Edna Culp, nurse and caregiver of her husband with advanced Parkinson’s, pictured with the Parkinson’s Foundation hospitalization kit at the Aware in Care Ambassador Training in Rochester, NY, on June 29, 2019.

A family attends the Parkinson’s in the Light of Day event, hosted by the Parkinson’s Foundation and Light of Day Foundation, in Edison, NJ on April 13, 2019. The one-day event addressed the needs of those newly diagnosed with Parkinson’s.
As illustrated below, the Parkinson’s Foundation directed 83% of its 2018-2019 spending on mission-related activities and 17% on fundraising and management combined.*

**Includes Investment Income of $724,044**
The Parkinson’s Foundation is privileged to have an outstanding and dedicated group of volunteers who generously donate their time and talents by serving on the Board of Directors. Without their selfless efforts, the work of the Foundation would not be possible.

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The Parkinson’s Foundation makes life better for people with Parkinson’s disease by improving care and advancing research toward a cure. In everything we do, we build on the energy, experience and passion of our global Parkinson’s community.

WHAT WE BELIEVE

The Parkinson’s Foundation makes life better for people with Parkinson’s disease by improving care and advancing research toward a cure. In everything we do, we build on the energy, experience and passion of our global Parkinson’s community.

Engage with the Parkinson’s community and the Parkinson’s Foundation on social media.