On the Cover

The photo on the cover of this year’s Annual Report was taken by Brian Baker and features Charis Hill. Charis is, amongst many things, a professional model and an advocate. In 2013, she was diagnosed with ankylosing spondylitis, and she now runs the Spondylitis Association of America support group chapter in Sacramento, California. You can learn more about Charis here. She is just one of the almost 3 million faces of spondylitis in the United States, and we thank her for her time and effort, as well as the use of her photo.

The Faces of Ankylosing Spondylitis

Throughout the pages of this year’s Annual Report, you will see photos that were originally posted on the website The Faces of Ankylosing Spondylitis. The site was started by Cookie Hopper in 2011 for people diagnosed with spondylitis to share their images and stories. To date, the site has over 1,500 profiles. The photos are used here with the permission of Ms. Hopper, and those pictured do not necessarily endorse the Spondylitis Association of America in any way. To learn more about this agreement, please see here. To learn more about this fantastic project, please visit thefacesofankylosingspondylitis.com.

About the Spondylitis Association of America

The Spondylitis Association of America (SAA) was founded in 1983 by people affected by spondylitis. Prior to that time, there was no support network, no resource for educational materials, not even a single pamphlet for people diagnosed with the disease. SAA was the first and remains the largest resource in the US for people seeking information on anklylosing spondylitis and its family of related diseases. Our mission is to be a leader in the quest to cure anklylosing spondylitis and its family of diseases, and to empower those affected to live life to the fullest.

SAA Staff

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Executive Director

Chris Miller  
Programs Director

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Richard A. Howard, MBA  
Associate Executive Director

Elin Aslanyan  
Programs Manager

Linda Powell  
Administrative Associate

Diann Peterson, CFRE  
Director of Annual Giving

Robin Kindrick  
Executive Associate

Inna Mednikov  
Consulting Bookkeeper
Dear friends of SAA,

With your ongoing support, the Spondylitis Association of America continues to be the “go to” patient organization serving spondylitis and its family of related diseases in North America. FY2015 has been a year of expanded communications and outreach for SAA. We significantly ramped up our efforts to educate specialty doctors, including primary and family care physicians. In addition, we have supported groundbreaking research in HLA-B27 and the microbiome this past year that potentially will lead to a better understanding of the cause of this disease spectrum and lead to prevention and cure.

Moreover, we have expanded our partnerships and collaborative efforts with consumer, commercial, public and private-sector health organizations that aid us in pursuing our mission. Our social media environment continues to support and to educate those in need.

Indeed, our deepened commitment to research, with a newly focused collaborative approach, includes partnering with the American College of Rheumatology to develop treatment guidelines in spondyloarthritis. These first ever, and much needed, guidelines were completed and distributed to the rheumatology community and beyond and received a robust positive response from within the medical and patient community.

On the advocacy front, we have been working closely with other mission driven organizations, which has resulted in our lawmakers voting to extend funding for arthritis and joint disease research.

The Spondylitis Association of America is the only organization that solely supports individuals affected by spondylitis and its family of related diseases, who turn to us for information, support, and opportunities to be heard. We listen, and we act upon what we hear.

Our goal is to be bold, to work hard, and to make a difference. We are here to serve. We are proud of our successes, but our challenges — reaching those who are still suffering — are what drive us to work harder and do more every day. That said, while it is necessary that we remain steadily focused on the mission, we continue to be a flexible operation that periodically steps back from the day to day to check its surroundings.

It is our responsibility to remain fiscally robust in order to achieve these goals. That said, now is the time to not only continue to seize opportunities but to continue to drive them. More than ever, we are grateful for your ongoing encouragement and support.

Thank you.

Sincerely,

Laurie M. Savage MS FLE
Executive Director, CEO

Charlotte K. Howard
Chair, Board of Directors
SAA Programs:
Dedicated to You and Your Needs
We offer...

High quality in-person spondylitis educational seminars and information-packed webinars

One of a kind content presented by leading spondylitis researchers and clinicians, professionally developed with you in mind, and always free to attend.

The latest spondylitis research and breaking news right in your inbox

We bring you spondylitis news updates via our monthly e-newsletter, eSUN.

Connections to your community

In-person support through our national support group network, SAA-hosted online message boards, and thriving social media communities on Facebook and Twitter.

The only news magazine in the us dedicated entirely to spondylitis: SAA’s advertising-free Spondylitis Plus

Four issues a year offering in depth exploration of issues affecting those with spondylitis. The magazine delivers original content by SAA’s expert panel of medical professionals and SAA staff, as well as community voices.

“Thank you for all the support you provide. When I have a question about my condition, I go directly to your website and find the answer. I also love the message boards. It’s comforting to hear from others who share the challenges of this disease.”

— Theresa

“I have learned more from this seminar than I have the last 16 months with my rheumatologist. Thank you so much!”

— Jennifer
Reliable information and accurate educational materials on all aspects of living with spondylitis

Our website, spondylitis.org.

Our teen site, Spondylitis Web Info For Teens (SWIFT) teens.spondylitis.org.

Educational brochures, downloadable practical materials, and valuable resources - all provided free of charge to make sure you have all you need to understand and effectively treat spondylitis.

“I learn something new with every issue of Spondylitis Plus and every presentation like today’s webinar. I am especially grateful for people like today’s presenter, Dr. Reveille, who has devoted much of his career to helping spondylitis patients live better lives!”

— Debra

“This group is an environment to ask questions. It’s great to know that there is a support community here.”

— Joseph

Patient advocacy and physician education, along with our ongoing focus on raising spondylitis awareness to ensure your voice is heard

Attendance and exhibition at prestigious medical conferences with the goal of raising awareness of spondylitis and the issues impacting our community. Reaching out to the general public through awareness campaigns and our social media channels.
“Thank you for all you do to raise awareness!”
— Jacquie

“Thank you SAA for all your hard work. I love receiving your publications! It seems like every time there is something I want to read more about—women and AS, Sjogrens, new therapies and research, low starch and other diets, etc. You write about it all.”
— Elana

“It’s so nice to feel a part of this {Los Angeles area} seminar. I’m live streaming your program from Anchorage, Alaska. Thank you for doing this for us!”
— Donna

“Thank you for all you do to raise awareness!”
— Jacquie

Austin
SAA/Bruckel Early Investigator Award in Axial Spondyloarthritis (AxSpA) 2014

Awardee

In 2011, in honor of SAA Co-founder, Jane Bruckel, the Spondylitis Association of America established an award as part of its stated mission to encourage young research clinicians to consider a career in rheumatology and specifically to care for the spondylitis patient and to support research in this field. Thus, an award in the amount of $20,000 is now made available annually to early career investigators who already have a demonstrated body of work in spondylitis and its related family of diseases.

Pamela Weiss, MD, is an attending physician and the Rheumatology Clinical Research Director at the Children’s Hospital of Philadelphia (CHOP.) Her current research efforts are focused on defining the clinical and imaging features of children with early juvenile spondyloarthritis. Long-term studies evaluating critical disease outcomes in children with spondyloarthritis are sparse. Outcomes critical to study include the rate and proportion of children who experience progression of arthritis of the lower back and spine, the likelihood of maintaining remission while on medication or if medications are discontinued, and the impact of therapy on outcomes that matter to children and their families (such as anxiety, peer relationships, fatigue, pain). Studying these critical outcomes through the creation of a longitudinal registry of children with spondyloarthritis has the potential to transform the way we approach and care for these patients long-term.
“Support from SAA has allowed me to begin registry development and enhance the scope of my research. Ideally, data collection will also eventually include blood samples and imaging. This project has the potential to vastly improve our understanding of the natural history of the disease in children, contributing factors to disease onset and progression, as well as the current therapy’s impact on the quality of life for patients.”

— Pamela Weiss, MD. 2014 Award Winner
Advancing Science

The SAA mission states that we are committed to being on the forefront of research toward prevention and a cure. In other words, the Spondylitis Association of America seeks to improve outcomes for patient on all fronts. In light of that commitment, in FY2015, SAA funded research in the following areas which we believe hold a robust potential to improve patient outcomes.

ERAP1 in HLA-B27 Transgenic Rats
Joel D. Taurog, MD

Since 1980, Dr. Taurog’s goal has been to identify the molecular basis for the association of the molecule HLA-B27 in spondylitis. For that purpose, HLA-B27 transgenic rats that develop spondyloarthritis were developed as an animal model; the purpose, to investigate the role of B27. As a result, it has been learned that ERAP1, a gene implicated in spondylitis, provides a critical functional interaction with HLA-B27. More importantly, research clinicians have identified that ERAP1 was not associated with spondylitis in HLA-B27-negative patients. Dr. Taurog, in collaboration with researchers at the National Institutes of Health, has been able to make additional seminal observations with these rats. The following hypotheses support this effort.

Dr. Taurog’s work has been able to uncover that decreasing the expression of HLA-B27 may potentially prevent HLA-B27-associated inflammatory disease. Two basic strategies are being explored. One is based on gene encoding a specific RNA, and the other is based on repeatedly administering a small interfering RNA. In the upcoming years, Dr. Taurog is anticipating expanded support of his work based on the outcomes made possible by the support of SAA.
Does HLA-B27 Shape the Gut Microbiome?
James T. Rosenbaum, MD

The intestinal microbiota has been firmly identified not only in the causation of bowel disease but also in the development of inflammatory diseases in other parts of the body. Researchers have been able to firmly correlate significant clinical, genetic, immunological, and microbiological overlap between many of these conditions including spondyloarthritis, specifically in conjunction with inflammatory bowel disease. Dr. Rosenbaum’s work supports a hypothesis that manipulation of the gut microbiota itself may be a promising future target for therapy in spondylitis.

The Spondylitis Association of America helped fund this novel hypothesis, and we will soon be inviting and enrolling patients to actively participate in this research, which is being conducted at the Oregon Health Sciences University in Portland, OR.

“Keep doing what you are doing. It’s good to not feel alone. I had never heard of spondylitis before last year.”
— Sydney
Research and Physician Education

Magnetic Resonance Imaging Training for Rheumatologists and Radiologists

This cutting edge series of hands-on workshops sponsored by SAA is geared to train physicians to identify spondylitis using 3D imaging (DICOM) on individual case study preloaded laptops. The objective of this accredited program is to accelerate diagnosis by recognizing early clinical signs of axial spondyloarthritis as well as being able to track potential disease progression in more advanced, already diagnosed cases. This program has been presented in Los Angeles, Denver, and Chicago.

“I’m very impressed with how SAA treats us members. I’m grateful to be a member of such a well-run, caring organization, and will continue to seek ways to give back.”

— Susan
Treatment Recommendations

ACR/SAA/SPARTAN Treatment Guidelines in Ankylosing Spondylitis and Non-radiographic Axial Spondyloarthritis Consultant: Michael M Ward, MD, MPH Intramural Research Program NIAMS/NIH, Bethesda, MD

*To provide evidence-based recommendations on the treatment of patients with ankylosing spondylitis (AS) and non-radiographic axial spondyloarthritis (nr-axSpA).*

The Spondylitis Association of America collaborated with the American College of Rheumatology in the funding and development of the very first spondylitis guidelines established in the US. These recommendations are intended to provide doctors, physical therapists, and other clinicians who care for patients with spondylitis and its family of related diseases with guidance on approaches to treatment that are based on current and best-available medical research. Given the wide diversity of medical conditions and the breadth of medical knowledge, it is sometimes difficult even for subspecialists, such as rheumatologists, to be fully up-to-date on the treatment options and alternatives to consider in all types of situations.

There are three important points that patients should keep in mind regarding these recommendations. First, these are not a set of rules that doctors must follow. There are many reasons why a particular recommendation may not be followed for an individual patient, including their prior medical history, other co-existing illnesses or medications, the stage of disease, and considerations of the trade-offs between the potential benefits and harms of a given treatment choice. Think of these more as a resource guide that doctors can use to help their decision-making in recommending (or not recommending) particular tests or treatments. All treatment decisions must be individualized to fit the patient and his or her particular circumstances.

Second, the recommendations don’t consider all possible issues or questions that can occur in patients with spondylitis. The recommendations focused on the most common questions, but many others remain. We hope that these recommendations can be expanded to include more questions in the future.
Third, these recommendations are not unchangeable. While they are based on the best evidence available to date, new research may uncover new findings that would cause us to change a recommendation. This would be unlikely for recommendations that were designated as “strong,” because the evidence for these was considered so solid that any new research was thought very unlikely to change the conclusion. However, for “conditional” recommendations, new research may change the recommendation as more facts are uncovered. Also, new treatments may be developed that would change our thinking about the way patients should be treated in particular circumstances. Or new information about side effects of treatment may be produced to change our ideas about the safety of a given treatment. It is important to note that most of the recommendations were conditional ones. To make sure the recommendations stay up-to-date, we plan to review them every few years and issue updates earlier if important breakthroughs occur that make a current recommendation no longer valid.
We are pleased to present the Spondylitis Association of America’s Financials and Statement of Financial Position for the year ended June 30, 2015. Our financial position and diversified fundraising activities support our ability to fund the most promising medical research, influence public policy for prevention of spondylitis and quality spondylitis care, and offer programs supporting the needs of spondylitis patients and caregivers. SAA’s volunteer board of directors is committed to proactive stewardship of the organization’s financial resources.

The accounts and financial statements of the Spondylitis Association of America are maintained in accordance with generally accepted accounting principles for nonprofit organizations as established by the American Institute of Certified Public Accountants. For a complete copy of our annual financial audit report, conducted by RBZ, LLP and our nonprofit tax return (IRS form 990), please visit spondylitis.org or call 800-777-8189.

“I wanted to thank you all so much for everything you do, have done, and will continue to do for all of us. It is very doubtful that you will ever truly understand the amount of gratitude and appreciation we have for all of your hard work, dedication, and sacrifice.”

— Cookie
Financials

FY2015 Revenue
- Bequests 24%
- Corporate and Foundation Grants 43%
- Individual Contributions 30%
- Program Revenue 1%
- Interest Income 1%
- Gain on Investment <1%

FY2015 Expenses
- Management & General 15%
- Physicians Education 17%
- Patient Conferences, Education & Support 45%
- ERAP1 Genetic Research 8%
- Scientific Study (MRI, Registry, Guidelines) 6%
- Fundraising 7%
- Microbiome Research 2%

Gain on Investment <1%
Program Revenue 1%
Interest Income 1%
Statement of Financial Position

Year ended June 30, 2015

Assets
Cash and Cash Equivalents ................................................................. 489,148
Investments, at market value ............................................................ 1,714,216
Prepaid Expenses .............................................................................. 9,979
Property and Equipment
   Furniture and equipment ............................................................... 57,466
   Less: Accumulated depreciation .................................................. (42,064)
   Total .......................................................................................... 15,402
Deposits ............................................................................................ 5,441
Total Assets ...................................................................................... 2,234,186

Liabilities and Net Assets
Accounts Payable ............................................................................ 4,411
Accrued Vacation ............................................................................ 28,485
Total Liabilities ................................................................................ 32,896
Net assets
   Unrestricted net assets ................................................................ 1,388,669
   Board designated fund ................................................................. 183,632
   Temporarily restricted net assets .................................................. 628,989
Total Net Assets ................................................................................ 2,201,290
Total Liabilities and Net Assets ........................................................ 2,234,186
Statement of Activities

Year ended June 30, 2015

Changes in unrestricted net assets

Revenue and Support
Corporate and foundation grants ............................................................ 680,283
Individual Contributions ........................................................................... 477,461
Bequests .................................................................................................. 381,348
Interest income ......................................................................................... 15,420
Program revenue ....................................................................................... 17,201
Realized & Unrealized Gain on investment ................................................. 4,660
Total unrestricted revenue ........................................................................ 1,576,373

Net assets released from restrictions
Satisfaction of program restrictions ......................................................... 237,720
Total unrestricted revenue and other support ......................................... 1,814,093

Functional Expenses

Program Services
Medical research Grants .......................................................................... 125,000
Scientific Study ......................................................................................... 80,318
Medical conferences presented ................................................................. 223,530
Patient Education Conferences & Patient Support .................................. 592,944
Total Program Services ........................................................................... 1,021,792

Management and General ...................................................................... 191,407

Fundraising Expenses ............................................................................. 93,715

Total expenses .......................................................................................... 1,306,914

Increase in unrestricted net assets .......................................................... 507,179

Net assets, beginning of year ................................................................. 1,694,111
Net assets, end of year ............................................................................ 2,201,290
Board Officers

Charlotte K. Howard  
*Board Chair*

Michael Pianin, Esq.  
*Vice Chair*

Leslie Kautz, CFA  
*Treasurer*

Eric Goldstein, CPA, MBA  
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*Craig Gimbel, DDS*  
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Toronto, ON

Muhammad Asim Khan, MD  
Cleveland, OH

Honorary Board Members

Harry Bruckel, CPA  
Jane Bruckel, BSN, RN
Deverell

Development
Raising the Funds to Lead the Fight
Advancing medical research and improving the lives of the spondylitis community everywhere means responsibly raising the funds needed to support scientific study and provide educational and emotional support to the community. This is a substantial commitment. SAA depends upon the generous contributions from individuals, foundations, and corporations to meet its annual operating budget and to fund research, as well as special programs and services.

The following individuals, companies, and foundations have made important and generous contributions to the Spondylitis Association of America in support of our shared mission. We are extremely thankful for the donors, partners and volunteers whose efforts give us the power to fund groundbreaking research, advocate for healthy public policy, provide information about spondyloarthritis, promote prevention and early diagnosis, and support those living with all forms of spondylitis.

We also gratefully acknowledge the support of the donors who contribute to SAA through the Combined Federal Campaign, State Employee Campaigns, the United Way, and all other federated campaigns in the workplace.

Every dollar given supports our passion to deliver on our mission to be a leader in the quest to prevent spondyloarthritis and empower those affected to live their lives to the fullest.

“For nearly 25 years, I have dealt with the effects of AS. through my legacy gift, I am making a lasting contribution so future generations will not have to experience the disease in the same way I have.”

— Brian Mackenzie, Quest Legacy Society Member
Corporate Partners

The Corporate Membership Program provides a way for pharmaceutical companies to positively impact the lives of those affected by spondylitis by contributing to our general operating budget. In addition, SAA receives corporate support for special educational programs.

Foundations

SAA foundation funders make a positive difference in the lives of all those in the spondylitis community by supporting the Spondylitis Association of America’s general operating budget as well as funding special projects. SAA would like to thank the following foundations for their generosity and support.

Cecile & Fred Bartman Foundation
Ellen & Marshall Cole Philanthropic Fund
Community Foundation Alliance
Fred & Mabel R. Parks Foundation
Higgins Family Charitable Foundation
Jean & E. Floyd Kvamme Foundation
Kautz Family Foundation
Koven Foundation
Murdy Foundation
Stupp Bros. Bridge & Iron Co. Foundation
Sydney T. Levenson, M.D. Foundation
Dalís Foundation
Thistle & Rose Foundation
Young Foundation
Leadership Circle

We are proud to acknowledge the members of the SAA Leadership Circle. These dedicated donors provide significant support that enables us to remain fiscally nimble and step up to meet opportunities as they arise. They are committed to being our partners in progress and have made possible many of the research and educational programs discussed in this Annual Report.

$50,000
Jean & E. Floyd Kvamme Foundation

$30,000
Higgins Family Charitable Foundation

$10,000 to $29,999
Cecile & Fred Bartman Foundation
John Eckermann
Fred & Mabel R. Parks Foundation
Kautz Family Foundation
Local Independent Charities of America
Herb & Barbara Shear

$2,500 to $4,999
Community Foundation Alliance
Margo & Stanley Itskowitch
Jonathan Lathrop
Eric Passmore
Milt B. Schulle
Thistle & Rose Foundation
Barbara Van Alstine

$1,000 to $2,499
AmazonSmile Foundation
Keith & Celia Arnaud
Jake Aronov
Ellen Ashby
Buck & Leslie Balkind
Brian Berman
Bradley & Marla Bockhorst
Bravelets LLC
Harry & Jane Bruckel

$1,000 to $2,499
Arnold Burgemeister
John Cella, III, MD
Kevin Cusick
Chris Deininger
Nero Deliwal
Kevin & Tina Elliott
Judy Fiskin
Andrew & Lauren Forbes
Fabio & Wei Garofolo
Craig Gimbel, DDS
Bill Giser
Susan & Alexander Goldberg
Eric & Tahl Goldstein
Philip Gully, III
David Hallegua, MD
Elmer & Arlene Hansen
Robert & Barbara Hasty
Todd & Kathleen Herzog
Shari & Craig Jankowsky
Marjorie B. Joseph
Shelby Jiulfs
Daniel Kautz
Rachel Knopoff
Kroger Grocery
Jacqueline & Richard Litchfield

$5,000 to $9,999
Charlotte K. Howard
Leslie Kautz
Deidra & Niki Krutop
Marshall Amplification, PLC
Murdy Foundation
Michael & Barbara Pianin

$1,000 to $2,499
Heather & Franklin Lopane
Robert & Mary Jo Lund
Brian MacKenzie
Mary Major
Network for Good
Maria & Martin Nyvall
Paula Olsiewski, PhD
Eileen & John Pacilio
Michael Patterson
Timothy & Suzanne Quinn
James Rosenbaum, MD
Laurie & Thomas Saylak
Alan Clifton Schaffer
Jeffrey & Diana Schmitt
The Dalis Foundation
Danielle Thomas
Barry & Christine Tobias
Alison Toth, MD
Robert W. Ulrich, PharmD
Laura Wagner
Tom & Barbie West
Joe & Cindy Wickwire
Jennifer Wilbur
Janet & David Williams
Hilary Wilson
Young Foundation
The Quest Legacy Society

The Quest Legacy Society recognizes those individuals who have taken the special step to include SAA in their estate plans through charitable bequests, trusts, insurance or other forms of deferred giving. Among our most valued supporters, members of the Quest Legacy Society are determined to ensure a future wherein SAA can continue to fund medical research, produce educational programs and advocate on behalf of the spondylitis community beyond their own lifetimes. The planned gift bestows upon all those affected by the disease, their families, and the future generations a reason for hope. Information about the Quest Legacy Society can be found at spondylitis.org/quest.

SAA is honored to recognize these Quest Legacy Society members:

Charles Anderson* (2014)
Morey & Lisa Anderson
Grace Becker and the late Stephen Becker (2014)
Jane & Harry Bruckel
Lillian Buckley* (2001)
Cecelia Bunch
Gary Cadle
Raymond & Mary Jean Cassidy* (2004)
Means Davis, Jr.
Richard and Deanna Day
Harvey R. Derscheid
Alan Fraser* (2015)
David Fulle
Kathleen Geier* (1994)
Craig B. Gimbel Family Trust
David S. Hallegua, MD
Robert and Barbara Hasty
Phil & Eileen Hipe
Charlotte Howard
Karin & Richard Howard
Marjorie Joseph
Marvin Kallison* (2009)
Nicholas Karatsoulis* (2002)
Leslie B. Kautz
Colin Kraft
Jennifer Layden
Batya Bilmes Lemler
Linda Lovello
Brian MacKenzie
Jane & Hugh McGonagle
Charles Mol
Barbara Murchison* (2008)
Christopher & Stephanie Oleksy
Paula J. Olsiewski
Barbara & Michael Pianin
Nicholas Pregibon* (2013)
Matthew J. Rathburn and William A. Schell
Nancy Regan
John D Reveille, MD
Dan Robinson* (2013)
James Rosenbaum, MD
Zoanne Sager and the late Stephan Howard Kramer
Randall Sapadin, The Maria Coppola Foundation
Laurie M. Savage
Rachel & Justin Shifrin
Katherine Sprouse
Shirley Symington* (2007)
Ira Thompson* (2002)
Bob & Janet Ulrich
Gregg Umek
Hilary Wilson
Darren B. Wolpert

* Remembered in perpetuity (year donation received)
Membership and Individual Donors

$500 to $999

Steven Abramson
Stan Amy
Andrea Barnett
Michael Barton
David Baxter, PhD, DO
Gerald & Estelle Becker
William Blaze
Rafael Bras
Todd Bruckel
Donald & Judy Bunin
Marshall & Hedy Butler
Joanne Chamberlin
Susan Copple
Nancy Davis
Tracey Dellaripa
John & Cathie Duniway
Deb & Kevin Elm
Jane & Dane Farnsworth
Britt-Louise Fletcher
Jesus & Gladys Garza
Robert & Rhonda Gendron
Alan Glaser
Doug Granat
Charles Grisemer
Greg Gulley
Buffa Hargett
Elizabeth Hogan
Jeff & Nathana Horn
David & Katharine Hutchison
Dan & Lynne Kelly
David King
Skaye & Albert Kirk
Koven Foundation
Damon F. Kvamme
Ivan & Fern Lefkowitz
Sandor Lehoczky
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Molly McLaughlin
Barry & Amy Misbin
James Moore
Jane Morrill
Amy Myers
Betty Opbroek
Bernadette Pahulje
Trisha & Joshua Palmer
Diann Peterson
Sheila Pohly
Katherine Poppe
Kalees Rayner
RBZ, LLP
David Reale
Steve Roethe
Ron & Christine Rogers
Kathleen Rose
David Rose
Michael Rothermund
Laurie M. Savage
Charles Sellers
Cynthia Shaghoian-Spizzirri
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Six Continents Hotels, Inc
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Markos I. Tambakeras
Terri Tate
Brian & Jill Tichenor
Jayne & Kelly Tien
David Towery
Norma & Donald Troester
John Trotter
United Way of Southeastern Pennsylvania
Peter & Katherine Van Schoonhoven
David & Christine Vernier
Kelly Vilaruel
Robert Warren, MD PhD
Donna & Jim Wisler
Orna & Keenan Wolens
Gayle Wyman
David T. Yu, MD

$250 to $499

Caroline & Rob Abrom
Allsup
Marika Apostolakis
Joanna Atkinson
Betty & George Baffa
Joanie Balach
Robert & Christine Baldoni
Laveesh Bansal
John Barnes, MD
James Barnett
Jennelle & Jesse Barton
Angela Bartz
Sarah Bates
Jeff Baumann
Larry Berkelhammer
John Bernier
Sarah Bittikofer
Carl & Sarah Blackmon
Robert Blauvelt
Kerry Block
Richard Bowie
Mike Braun
Jean Braxton
Michael R. Brown
Duncan Brown
William & Lauraine Buehler
Norman Buller
Howard Bulos
CeCelia Bunch
Susan Butler
Shelly Canale
Ann Cane
Ellen Carroll
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Jennifer Clark
Daniel O. Clegg, MD
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Michael A. Cohen
Marshall Cole
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Edward G. Conley
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Michael V. Criss
Holly Danneker
Edgardo V. De Jesus
Mary Jo Deichler
Kay & Joseph Demarte
John Digianni
Jennifer Ebbeler
Kim Evans Couch
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Kelly & Frederic Ewald
Rita Finley
Prentice Fish
Jason Fletcher
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Guy Glimpse
Kimberly Goldinger
Douglas Good
Melinda Greene
James & Laura Hale
Karen Hall
Jean Hansen
Machelle Haus
Thomas & Mary Hayden
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Robert Hecox
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Lisa Henderson
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Penny Honeycutt
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Rita M. Brady  
Ken Brasher  
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Steve Bricker  
Diane Brititz  
James Brinn  
Eleanor Brockington  
Louise & Joseph Brogna  
Joan Bronson  
Jack Brooks  
Wes Brooks  
Nancy Brothers  
Larry Brown  
Garry M. Brown  
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