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*Photos for this Annual Report are courtesy of Barbara Alper of Barbara Alper Photography.*
The Spondylitis Association of America is dedicated to improving the lives of people impacted by the disease. In 2017, we made significant strides to live out our mission and purpose. Below are some successes we celebrated in raising awareness of spondylitis, supporting and empowering SpA patients, and leading the fight for a cure.

- 2017 – SAA conducted the third Life Impact Survey to understand the effect the disease has on quality of life and to compare those results to ones collected in 2002 and 2011.

- 2017 – SAA held the Spondyloarthritis Unmet Needs Conference III on the NIH campus to continue driving innovative and creative approaches to research.

- 2017 – SAA launched our new website with additional features and ease in navigation. The redesigned website continues its legacy of being a trusted source of information for patients and professionals for spondyloarthritis.

- 2017 – SAA launched major awareness initiatives that reached more than 500 million people. Elements included white board animation videos, electronic wallboards in physician waiting rooms, and the This AS Life Live! video series with pop-rock celebrity Dan Reynolds.
The Spondylitis Association of America (SAA) was founded in 1983 by a group of volunteers to address the needs of people affected by spondylitis. Our journey arose from the recognition that prior to our founding, there was no support network, resources for educational materials, or even a single pamphlet for people diagnosed with the disease. SAA was the first and continues to be the largest resource in the U.S. for people seeking information on Spondyloarthritis. Our mission is to be a leader in the quest to cure ankylosing spondylitis and related diseases, and to empower those affected to live their lives to the fullest. SAA receives no government funding and relies on generous donations from individuals to create and maintain programs and services aimed at improving the futures of the more than 2.7 million American adults affected by spondylitis.

Meet the SAA Staff

Laurie M. Savage, MS FLE
Executive Director

Richard Howard, MBA
Associate Executive Director

Diann Peterson, CFRE
Associate Executive Director

Elin Aslanyan
Programs Director / Editor-in-Chief, Spondylitis Plus

Sean Ewert
Development & Annual Giving Manager

Robin Kindrick
Executive Associate

Helene Hart
Membership Associate

Linda Powell
Administrative Associate

Inna Mednikov
Consulting Bookkeeper
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Harry Bruckel, CPA

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Joel Taurog, MD
Ruben Burgos-Vargas, MD
Robert W. Warren, MD, PhD, MPH
Michael H. Weisman, MD
SAA uses its resources wisely to assure both stability and innovation in programs for our members. Through the generosity of our members and support from our corporate sponsors, we drive awareness, support funding for research, advocate on behalf of our community for changes in public policy, and offer programs to address the needs of spondylitis patients and caregivers. We are pleased to share our Revenue and Expense Statements, Statement of Activities, Statement of Financial Position, and Statement of Financial Activities for the year ending June 30, 2017. SAA’s volunteer Board of Directors provides proactive protection of our financial resources, ensuring that we operate as good stewards of the public’s trust and use our resources efficiently to advance our mission.

The accounts and financial statements of SAA 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 Armanino, LLP and our nonprofit tax return (IRS form 990), please visit spondylitis.org or call 800-777-8189.
# Statement of Activities

**Spondylitis Association of America**  
Statement of Activities  
for the Year Ended June 30, 2017

<table>
<thead>
<tr>
<th>Revenues, gains, and other support</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Corporate and foundation contributions</td>
<td>$516,865</td>
<td>$516,865</td>
<td></td>
</tr>
<tr>
<td>Program revenue</td>
<td>$6,118</td>
<td>$6,118</td>
<td></td>
</tr>
<tr>
<td>Interest &amp; dividends</td>
<td>$25,145</td>
<td>$25,145</td>
<td></td>
</tr>
<tr>
<td>Realized and unrealized gains on investments</td>
<td>$4,665</td>
<td>$4,665</td>
<td></td>
</tr>
<tr>
<td>Net assets released from restriction</td>
<td>$220,833</td>
<td>$(220,833)</td>
<td></td>
</tr>
</tbody>
</table>

Total revenues, gains, and other support: $1,578,782

![Revenues, gains, and other support table]

<table>
<thead>
<tr>
<th>Functional expenses</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Program services</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Biomarker research</td>
<td>$50,000</td>
<td>$50,000</td>
<td></td>
</tr>
<tr>
<td>NIH Collaboration Unmet Needs</td>
<td>$22,660</td>
<td>$22,660</td>
<td></td>
</tr>
<tr>
<td>Life Impact Study</td>
<td>$60,475</td>
<td>$60,475</td>
<td></td>
</tr>
<tr>
<td>Physicians’ education (MRI, Rheumatology)</td>
<td>$78,644</td>
<td>$78,644</td>
<td></td>
</tr>
<tr>
<td>Patient conferences, education &amp; support</td>
<td>$777,141</td>
<td>$777,141</td>
<td></td>
</tr>
</tbody>
</table>

Total program services: $988,920

![Functional expenses table]

<table>
<thead>
<tr>
<th>Support services</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Management and general</td>
<td>$232,635</td>
<td>$232,635</td>
<td></td>
</tr>
<tr>
<td>Fundraising</td>
<td>$79,499</td>
<td>$79,499</td>
<td></td>
</tr>
</tbody>
</table>

Total support services: $312,134

![Support services table]

Total functional expenses: $1,301,054

<table>
<thead>
<tr>
<th>Change in net assets</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Net assets, beginning of year (Note 7)</td>
<td>$1,749,555</td>
<td>$521,306</td>
<td>$2,270,861</td>
</tr>
<tr>
<td>Net assets, end of year</td>
<td>$2,027,283</td>
<td>$644,723</td>
<td>$2,672,006</td>
</tr>
</tbody>
</table>

![Change in net assets table]
Statement of Financial Position

Spondylitis Association of America
Statement of Financial Position
June 30, 2017

ASSETS

Current assets
  Cash and cash equivalents $384,497
  Investments 2,129,789
  Pledges receivable 129,500
  Prepaid and other current assets 8,208
  Total current assets 2,651,994

Property and equipment, net 122,922

Deposits 5,441

Total assets $2,780,357

LIABILITIES AND NET ASSETS

Current liabilities
  Accounts payable $75,710
  Accrued vacation 32,641
  Total current liabilities 108,351

Commitment (Note 9)

Net assets
  Unrestricted
    General 1,684,457
    Board designated fund 342,826
    Total unrestricted 2,027,283
  Temporarily restricted 644,723
  Total net assets 2,672,006

Total liabilities and net assets $2,780,357
Dr. Lianne Gensler (center) is a highly-regarded rheumatologist at UC San Francisco Medical Center and the Director of the Ankylosing Spondylitis Clinic which specializes in treating patients who are impacted by the disease.

After earning her medical degree at UC Irvine, Dr. Gensler completed an internal medicine residency, chief residency and rheumatology fellowship at UCSF and subsequently joined the Rheumatology medical staff. Her primary research interests and expertise are in studying the disease progression of axial spondyloarthritis, including ankylosing spondylitis, and identifying the predictors of osteoporosis development in patients with systemic lupus erythematosus.

Also pictured: Michael Weisman, MD (left) and James Witter, MD, Ph.D. FACR (right)
Dr. Walter Maksymowych (below) was a guest speaker at the 2017 Spondyloarthritis Unmet Needs Conference III. He is a Professor in the Department of Medicine, Division of Rheumatology at the University of Alberta, Alberta, Canada. His primary research interests are the imaging, genetics, and treatment of spondyloarthritis, and the clinical validation of biomarker technologies for rheumatic diseases. In addition, he has published over 200 research articles and is a member of numerous international societies related to arthritis research.
Spondyloarthritis (SpA) is an umbrella term for inflammatory diseases that involve both the joints and the entheses (the sites where the ligaments and tendons attach to the bones). The most common of these diseases is ankylosing spondylitis. Others include reactive arthritis, psoriatic arthritis, and enteropathic arthritis, which is associated with inflammatory bowel disease.

In most cases, spondyloarthritis primarily affects the spine. Some forms can affect the peripheral joints – those in the hands, feet, arms and legs.

Work disability affects 10 percent to 20 percent of patients with ankylosing spondylitis, most often in those with physically demanding jobs. Lost income and lost productivity due to work disability represent major burdens to both families and society.

(Reveille et. al 2012)

In seven European and North American countries, almost a third of patients with psoriasis seen in dermatology centers had psoriatic arthritis (PsA) as determined by rheumatologists. Of the patients given the diagnosis of PsA in this study, 41 percent had not received a previous PsA diagnosis, suggesting under-diagnosis of patients in dermatologic practices of this potentially debilitating disorder.

(Mease 2013)
Corporate Sponsorship

The corporate sponsorship program provides a way for SAA’s pharmaceutical members to positively impact the lives of those affected by spondylitis by contributing to SAA’s general operating budget. SAA also receives additional corporate support for special – and costly – programs such as the live patient education seminars.

Platinum Sponsor

AbbVie

Gold Sponsors

Amgen
Novartis
UCB

Silver Sponsor

Pfizer

Bronze Sponsor

Lilly

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Robert & Mary Jo Lund
Nancy Regan
John Reveille, MD
James Rosenbaum, MD
Joe & Cindy Wickwire

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Jake Aronov
Harry & Jane Bruckel
Arnold Burgemeister
Joseph D’Ambrosia
Judy Fiskin
Britt-Louise Fletcher
Margo & Stanley Itskowitch
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Jonathan Lathrop
Jane & Scott Morrill
Paula Olsemowski, PhD
Harold Phillippi
Michael Pianin
Scarlet Pearl Casino Resort
Milt B. Schulle
Herb & Barbara Shear
Barry & Christine Tobias
Alison Toth, MD

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Keith & Celia Arnaud
Michael Barton
David Baxter, PhD, DO
Steven & Deborah Benner
Bradley & Marla Bockhorst
Ellen Carroll
Hu & Vivian Chao
Chris Deininger
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Herwin Gill
Craig Gimbel, DDS
Bill Giser
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Ellen Gonzalez
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Buffa Hargett
Robert & Barbara Hasty
Steve & Ann Herendeen
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Mike Hileman
David & Katharine Hutchinson
Shari & Craig Jankowsky
Daniel Kautz
Sharon Kornhaus
Naomi Langer-Seltzer & Larry Seltzer
Jerry Lim
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Susan M. Mandell
Mcgivney Foundation Inc
Nancy & Dan Morgan
Ed & Therese Neely
Mariah & Martin Nyvall
Catherine Otto, MD
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Sol Pinchuk
Katherine Poppe
Timothy & Suzanne Quinn
Luther Ragin, Jr
Laurie & Thomas Saylak
Esther Seto
Karrie Shogren, PhD
Theodore Theophilos
Danielle Thomas
Genevieve & Fenton Tom
Hanan Watson
Tom & Barbie West
Orna & Keenan Wolens
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. Quest Legacy gifts ensure that SAA can continue advancing research, supporting, educating, and advocating on behalf of everyone within our spondylitis community. Information about the Quest Legacy Society can be found at spondylitis.org/quest. For more information on planned giving, please contact Richard Howard at legacy@spondylitis.org or (818) 855-2103.

SAA is honored to recognize these Quest Legacy Society members:

- Charles Anderson* (2014)
- Morey & Lisa Anderson
- 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
- Eli Klepacz* (2017)
- Colin Kraft
- Jennifer Layden
- Batya Bilmes Lemler
- Linda Lovello
- Emiliano Mazzarella
- Brian MacKenzie
- Charles Mol
- Barbara Murchison* (2008)
- Christopher & Stephanie Oleksy
- Paula J. Olsiewski
- Kevin & Karen Padrick
- Diann Peterson
- Harold Phillippi* (2016)
- Barbara & Michael Pianin
- Sol Pinchuk* (2017)
- 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
- Laurie M. Savage
- Rachel & Justin Shifrin
- Katherine Sprouse
- Shirley Symington* (2007)
- Ira Thompson* (2002)
- Bob & Janet Ulrich
- Gregg Umek
- Shirley van Leeuwen
- Hilary Wilson
- Darren B. Wolpert

* Remembered in perpetuity and (year of donation)
The SAA does not receive funding from the government. Through generous donations from our members and individuals, like you, we are able to live out our mission and continue our efforts.

We have made every effort possible to ensure that this list is accurate and complete. In the event of an error or omission, we extend our apologies and encourage you to contact Diann Peterson, Associate Executive Director at 800-777-8189, ext. 226 or by email at diann.peterson@spondylitis.org so that proper recognition may be ensured in the future.

**$500 – $999**
Charles Adams, Jr  
Kristy & James Arnold  
Barbara Arrighi  
Deborah Beggin  
Shirley Boundy  
Rafael Bras  
Cynthia Butler  
Marshall & Hedy Butler  
David Campbell  
Alan Carmack  
Joanne Chamberlin  
Curtis Cheatham  
Marc Cohen  
Jim Colombo  
Niel Constantine  
Susan Coppole  
Nancy Davis  
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Tracey Dellaripa  
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William Enfield  
Martha Escobar  
Nicola Fuentes Toubia  
Susan Gallagher  
Robert & Rhonda Gendron  
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Alan Glaser  
Guy Glimpse  
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Greg Gulley  
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Michael Hensley  
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Jeff & Nathana Horn  
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Ted Lambrinos  
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Eileen & John Pacilio  
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Diann Peterson  
Sheila Pohly  
Eleanor Price  
Kalees Rayner  
Steve Roethe  
Pamela Rogacki  
Ron & Christine Rogers  
Kathleen Rose  
Fran Rosier  
Michael Rothermund  
Lucy Ellen Sallick  
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John Trotter  
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Robert Warren, MD PhD  
Donald Weber  
Jonathan White  
Darren B. Wolpert  
Mary Ann Wray

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Joseph Allard  
Malikah Alsaud  
Timothy P. Anderson, PhD  
Trudy Angel  
Diana, Richard, & Brian Annett  
David Arnett  
Joanna Atkinson  
Betty & George Balfa  
Andrea Barnett  
James Barnett  
Mark Barry  
Jennelle & Jesse Barton  
Angela Bartzen  
Sarah Bates  
Jeff Baumann  
Erica Behr
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Kathleen Brown
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Joanne Liang
Linda Lovello
Sandra Ludwig
Henry Lynch
Denize Machit
Sason Mahdi
Diane Malaquias
Matthew McCutchen
Helen McLane
Cheri & Scott Mensching
Janet Meyer
Alan Mibab
Rebecca Middlebrook
Michael & Kristin Milner
Barry & Amy Misbin
Perry Montgomery
Martin Montoro
Amanda Moore
Jeffrey & Hilarie Moore
Thomas W. Morgan
Gary Mozenter
Margaret Myers
Amy Myers
Ford & Catherine Nicholson
Renée Novak
Scott & Laurie Obert-Thorn
B.A. Blue Olson
Kristi Otto
Carolyn Pace
Mary Padilla
Laila Pence
Luis Perelman
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Wristory
Marilyn Wyatt
Gayle Wyman
Lola Wynter
David T. Yu, MD
Volunteer fundraisers play a valuable role in helping to support SAA’s programs. Whether it’s a donation received through a portion of sales or some other creative fundraising project, we are always thankful for the donations that benefit our spondylitis community and enable us to live out our mission to lead the fight to find a cure.

Successful fundraising begins with just an idea and a desire to support those impacted by spondylitis. Regardless of the amount, we want to thank you for entrusting us to be good stewards of your support and inspired by your efforts to make a difference.

For more information about Spondylitis Association volunteer fundraising, please contact Sean Ewert at (818) 855-2106.