“SAA’s Perfect FOUR-STAR CHARITY NAVIGATOR RATING with a 68.19 out of 70 Score for Fiscal Health, Accountability and Transparency, Places It Among The Top Ten National U.S. Charities” ~ money.msn.com
DEAR MEMBERS,

With your ongoing support the SAA continues to be the “go to” patient organization in Spondyloarthritis in North America.

It has been said that “we deliver” where other organizations only promise. And, we promise to maintain this level of activity on your behalf in the upcoming year. To accomplish this, we will not only need to seize opportunities that arise but continue to drive them. With newly published prevalence data showing that indeed Spondyloarthritis is not a rare group of diseases but rather an under recognized one, we are poised to leverage these in all relationships. These include, but are not limited to, funding opportunities, media outreach, academia and government in order to continue to gain support for this group of disgracefully underfunded diseases.

That said, I do believe that it is necessary that with our eye steadily focused on the mission, we remain a flexible operation that periodically steps back from day to day operations to check its surroundings. I believe that we must continue to review significant environmental factors—what is changing at a major level—to explore and to observe where these could lead, and how they could affect us now and in the future. And consequently outline tactical plans in order to maintain our momentum. This is and has been our process throughout these past recent years, so far standing us in good stead.

I do believe that supported by a strong volunteer base and a talented and dedicated work team, with your help, we can only continue to be well on our way. Thank you for your confidence in our efforts.

Laurie M. Savage
Chief Executive
Spondylitis Association of America

“Now is the time to not only seize opportunities but to continue to drive them.”

Laurie Savage, Executive Director
SAA’S BOARD OF DIRECTORS

Craig Gimbel, DDS, Chair
Brian MacKenzie, MBA, Vice Chair
Leslie Kautz, CFA, Treasurer
Karrie Shogren, PhD, Secretary
Eric Goldstein, CPA, MBA
David Hallegua, MD
Jeff Horn, MBA
Charlotte Howard
Ann Kittell Howat
Michael Pianin, Esq.
John D. Reveille, MD
Clarence So, MBA
Robert Ulrich, PharmD
Hilary Wilson, MBA
Laurie Savage, Executive Director (ex-officio)

HONORARY BOARD MEMBERS

Rico Brogna
Harry Bruckel, CPA
Jane Bruckel, BSN, RN
Val Halamandaris
Mimi Kennedy

SAA’S SCIENTIFIC & MEDICAL ADVISORY BOARD

Chair:
James Rosenbaum, MD .............. Portland, OR
Bruce Clark, PT ..................... Vancouver, CA
Daniel Clegg, MD .................. Salt Lake City, UT
Robert Colbert, MD, PhD .......... Bethesda, MD
Atul Deodhar, MD .................. Portland, OR
Nortin Hadler, MD ................. Chapel Hill, NC
Robert Harris, MD ................. Whittier, CA
Robert Inman, MD ................. Toronto, ON
Muhammad Asim Khan, MD .... Cleveland, OH
Walter Maksymowych, MD ..... Edmonton, CA
Allan Metzger, MD ............... Los Angeles, CA
David H. Neustadt, MD .......... Louisville, KY
Millicent Stone, MB, MRCP (UK) MSc . Bath, UK
Joel Taurog, MD .................... Dallas, TX
Ruben Burgos Vargas, MD .... Mexico City, MX
A primary objective of SAA’s mission is our commitment to research, and none of the studies we have been involved with would have been possible without your ongoing support and participation. Here is just a sampling of research projects SAA has been involved in over the last year:

**SAA SEEDS A PATIENT REGISTRY FOR ANKYLOSING SPONDYLITIS**

A patient registry is a database - a compilation of data on people with AS. In this case, the registry will be a combination of three existing patient databases that have been used in ankylosing spondylitis research. Thus, we will build a new database that can look at thousands and potentially tens of thousands of patients with AS and be able to track health trends, disease severity over time, age, gender differences (or lack thereof), race, complications and much, much more.

“This is a HUGE step in an amazing direction!!! I know these registries have greatly changed the way many diseases were thought of and treated in many other areas. Opens a lot of doors.”

~Sabrina81, SAA Forum Member

**TASC - GENETIC RESEARCH CONTINUES TO YIELD RESULTS**

This year, the Triple-A (Australo-Anglo-American) Spondyloarthritis Consortium, or TASC, identified yet another in a series of genetic variants associated with increased susceptibility to ankylosing spondylitis as well as providing new clues to how the disease may be treated in the future.

“Our work shows the great value of partnering genetics research with functional investigations to determine the basic biology which leads to common diseases such as ankylosing spondylitis, the causes of which have remained an enigma for so long.”

~Professor Matt Brown, University of Queensland, AU
AS RESEARCH SURVEY - WOMEN WITH ANKYLOSING SPONDYLITIS
In April, 2012, SAA helped Robert D. Inman, MD, BA and Dharini Mahendira BSc, MD, FRCPC from the Toronto Western Hospital Spondylitis Program conduct a survey looking at women with Ankylosing Spondylitis and the potential effects of hormones, in the form of oral contraceptive pills, on AS. (Dr. Inman serves on SAA’s Scientific and Medical Advisory Board.) The response was tremendous: Over 600 participants with 557 women completing the entire survey - well over double the required participant numbers.

“I was pleased to see researchers interested in women and AS. Even with my fused spine I was able to carry my baby to term. She is 5 now. I can’t keep up with her, but I have so much fun on the sidelines!”

~Heather, SAA Facebook Friend
“I’m very impressed with how you and others at SAA treat 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. Please keep up the wonderful work!”

~Susan, SAA Member

WWW.SPONDYLITIS.ORG
With over 1,700 pages of information, spondylitis.org remains the best source of spondyloarthritis information on the web. This past fiscal year, spondylitis.org helped educate and support 665,866 unique visitors - enough to fill the largest stadium in the USA six times over.

SWIFT - SPONDYLITIS WEB INFO FOR TEENS - TEENS.SPONDYLITIS.ORG
SAA’s teen info site got a new, modern look and is the first and only site of its kind for teens with ankylosing spondylitis or related conditions.

SAA CONNECTIONS
We launched our new “Connections” program through which those who need support can find it in whatever form is most comfortable. From our forums on spondylitis.org to Facebook to in-person support groups, Connections is the new hub of all of SAA’s patient support programs.

SOCIAL NETWORKING
Facebook and Twitter have both become cornerstones of outreach for SAA where staffers regularly post news and engage fans by answering questions, posting polls and more. We hit the 3,000 fan mark on Facebook this past fiscal year and Twitter reached over 1,000 followers.

SEMINARS
SAA held free educational seminars in Portland, Atlanta and Nashville this year. Hundreds benefited from learning in person from the country’s leading rheumatologists and physical therapists.

“My past fiscal year, spondylitis.org helped educate and support 665,866 unique visitors - enough to fill the largest stadium in the USA six times over.”

Chris Miller, Director of Programs
EDUCATIONAL SUPPORT GROUPS
With over 35 active groups in 27 states, our educational support groups program continues to thrive thanks to the passion and commitment of our volunteer support group leaders. Expert guest speakers present regularly on a variety of diverse topics, from the latest spondylitis research to workplace accommodations, social security disability tips, exercise & physical therapy, and more.

eSUN & WEEKLY NEWS
Our monthly, info-packed mass email, the “electronic Spondylitis Update Newsletter” (eSUN) hit the 5,500 subscriber mark this past fiscal year, and weekly research and personal interest news items have continued to be a staple on spondylitis.org with 164,295 page views in our news section.

SPONDYLITIS PLUS & OTHER PUBLICATIONS
Our quarterly news magazine, *Spondylitis Plus* continues to be our flagship publication and is also produced, designed, written and edited in house. SAA continues to distribute thousands of brochures every year to patients and doctors alike. We have also continued our electronic distribution of publications such as our Recommended Member-to-Member Rheumatologist Directory and our Action Plan to Manage Spondylitis and more on request to those in need.

ADVOCACY
Our advocacy program boasts over 1,400 advocates, and we helped mobilize an “I Need My Rheumatologist” call-in day, and helped rally people to have Congress pass the “Arthritis Prevention Control and Cure Act” of 2010.

INFO HOTLINE & EMAIL
SAA’s toll-free assistance line and info email continue to be a sources of valuable information to those who have questions about spondyloarthritis, SAA events and support programs.

“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 diets, etc. - you write about it.”

~Elana, SAA Member & Facebook Friend

“Our educational support groups program continues to thrive thanks to the passion and commitment of our volunteer support group leaders.”

~Elin Aslanyan, Programs Manager
RAISING THE FUNDS TO LEAD THE FIGHT

We are deeply grateful for the individuals, corporations and foundations whose generous support enables the Spondylitis Association to lead the fight against AS and related diseases. As the only nonprofit organization in the US dedicated to improving the lives of the spondylitis community, SAA is committed to being efficient, effective and responsible stewards of your contributions.

We are proud that 82 cents of every dollar goes directly toward funding critical programs and services, and we pledge to continue to prove our value to the patrons and supporters who place their confidence and trust in us.

“Thanks everyone at SAA for all you do for me and everyone else suffering from AS. Your help has been invaluable!”

~Michele, SAA Member & Facebook Friend
TOP 10 CHARITY - OF ANY SIZE - IN THE U.S.
An analysis performed by personal finance news website MainStreet (www.mainstreet.com), also reported on MSN Money, named SAA as the ninth-highest rated national charity in America.

SAA received a perfect Four-Star rating from Charity Navigator, America’s leading independent charity evaluator, for the third consecutive year in 2011 and a 68.19 out of 70 score for its fiscal performance and operational responsibility, ranking it higher than thousands of other charities in the nation.

In its compilation of highest-rated charities, MainStreet looked at the leading charities in the U.S. based on their 2011 Charity Navigator scores, filtering out organizations that operate only locally and those that work mainly to collect and distribute funds to other charitable organizations.

“As the only nonprofit serving the needs of the 2.7 million people in the U.S. affected by spondylitis, SAA is honored to be recognized for our ongoing commitment to accountability, fiscal responsibility and program excellence,” said Laurie M. Savage, SAA Executive Director and CEO. “We are exceptionally proud of the fact that an organization with only nine employees can achieve the level of impact that we have.”

For more information about Charity Navigator scores, visit:

www.charitynavigator.org

“We are exceptionally proud of the fact that an organization with only nine employees can achieve the level of impact that we have.”

Laurie Savage, Executive Director
“As SAA’s Treasurer, I am proud that SAA is built on a foundation of strong financials and good governance. SAA uses its resources wisely to assure both stability and innovation in the programs our members count on. Effective oversight is a key component of this, as SAA’s finances are reviewed carefully by Staff, a Board Member Finance Committee, and outside auditors on an ongoing basis. SAA’s supporters can be assured their contributions are stretched to the maximum in supporting our mission.”

~ Leslie B. Kautz, CFA, Treasurer
## STATEMENT OF FINANCIAL POSITION

### YEAR ENDED JUNE 30, 2011

<table>
<thead>
<tr>
<th>Assets</th>
<th>2011</th>
<th>2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cash and cash equivalents</td>
<td>582,067</td>
<td>509,454</td>
</tr>
<tr>
<td>Investments, at market value</td>
<td>1,205,128</td>
<td>1,188,131</td>
</tr>
<tr>
<td>Government contract receivable</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Grants receivable</td>
<td>15,000</td>
<td>220,450</td>
</tr>
<tr>
<td>Accounts receivable</td>
<td>10,500</td>
<td></td>
</tr>
<tr>
<td>Prepaid expenses</td>
<td>8,289</td>
<td>15,100</td>
</tr>
<tr>
<td>Property, at cost</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Furniture and equipment</td>
<td>112,607</td>
<td>105,397</td>
</tr>
<tr>
<td>Less: Accumulated depreciation</td>
<td>(93,530)</td>
<td>(83,800)</td>
</tr>
<tr>
<td>Total:</td>
<td>19,077</td>
<td>21,597</td>
</tr>
<tr>
<td>Deposits</td>
<td>5,441</td>
<td>5,441</td>
</tr>
<tr>
<td><strong>Total Assets</strong></td>
<td><strong>1,845,502</strong></td>
<td><strong>1,960,173</strong></td>
</tr>
</tbody>
</table>

### Liabilities and Net Assets

<table>
<thead>
<tr>
<th>Liabilities and Net Assets</th>
<th>2011</th>
<th>2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>Accounts payable</td>
<td>28,085</td>
<td>8,243</td>
</tr>
<tr>
<td>Accrued vacation</td>
<td>21,610</td>
<td>24,425</td>
</tr>
<tr>
<td><strong>Total Liabilities</strong></td>
<td><strong>49,695</strong></td>
<td><strong>32,668</strong></td>
</tr>
</tbody>
</table>

Net assets:

- Unrestricted net assets | 1,342,809 | 1,382,548 |
- Board designated fund | 101,174 | 79,195 |
- Temporarily restricted net assets | 351,824 | 465,762 |

| **Total Net Assets** | **1,795,807** | **1,927,505** |
| **Total Liabilities and Net Assets** | **1,845,502** | **1,960,173** |
# STATEMENT OF ACTIVITIES

## YEAR ENDED JUNE 30, 2011

### Changes In Unrestricted Net Assets

<table>
<thead>
<tr>
<th>Revenue and Support</th>
<th>2011</th>
<th>2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>Corporate and foundation grants</td>
<td>331,845</td>
<td>291,559</td>
</tr>
<tr>
<td>Contributions</td>
<td>475,916</td>
<td>469,081</td>
</tr>
<tr>
<td>Interest income</td>
<td>13,545</td>
<td>26,615</td>
</tr>
<tr>
<td>Program revenue</td>
<td>21,786</td>
<td>25,462</td>
</tr>
<tr>
<td>Realized &amp; Unrealized Loss / Gain on investment</td>
<td>13,461</td>
<td>1,604</td>
</tr>
<tr>
<td><strong>Total unrestricted revenue</strong></td>
<td>856,553</td>
<td>814,321</td>
</tr>
<tr>
<td><strong>Net assets released from restrictions</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>satisfaction of program restrictions</td>
<td>353,237</td>
<td>491,542</td>
</tr>
<tr>
<td><strong>Total unrestricted revenue and other support</strong></td>
<td>1,209,790</td>
<td>1,305,863</td>
</tr>
</tbody>
</table>

### Expenses

<table>
<thead>
<tr>
<th>Expenses</th>
<th>2011</th>
<th>2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>Salaries</td>
<td>549,107</td>
<td>524,511</td>
</tr>
<tr>
<td>Employee benefits</td>
<td>40,094</td>
<td>43,815</td>
</tr>
<tr>
<td>Payroll taxes</td>
<td>45,250</td>
<td>42,008</td>
</tr>
<tr>
<td>Office and administration</td>
<td>65,969</td>
<td>86,848</td>
</tr>
<tr>
<td>Medical research consulting</td>
<td>29,520</td>
<td>112,500</td>
</tr>
<tr>
<td>Physician Education</td>
<td>50,844</td>
<td>51,643</td>
</tr>
<tr>
<td>Patient Education</td>
<td>62,373</td>
<td>50,884</td>
</tr>
<tr>
<td>Legal and accounting</td>
<td>57,149</td>
<td>52,225</td>
</tr>
<tr>
<td>Rent</td>
<td>61,502</td>
<td>40,506</td>
</tr>
<tr>
<td>Medical conferences presented</td>
<td>107,821</td>
<td>233,337</td>
</tr>
<tr>
<td>Travel</td>
<td>28,742</td>
<td>38,020</td>
</tr>
<tr>
<td>Postage</td>
<td>45,761</td>
<td>50,751</td>
</tr>
<tr>
<td>Printing</td>
<td>50,860</td>
<td>46,984</td>
</tr>
<tr>
<td>Outreach</td>
<td>22,829</td>
<td>21,925</td>
</tr>
<tr>
<td>Depreciation</td>
<td>9,730</td>
<td>9,061</td>
</tr>
<tr>
<td><strong>Total expenses</strong></td>
<td>1,227,551</td>
<td>1,405,018</td>
</tr>
</tbody>
</table>

### Decrease in unrestricted net assets

| Decrease in unrestricted net assets    | (17,761) | (99,155) |

### Changes in temporarily restricted net assets

<table>
<thead>
<tr>
<th>Changes in temporarily restricted net assets</th>
<th>2011</th>
<th>2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>Corporate and foundation grants</td>
<td>239,300</td>
<td>403,142</td>
</tr>
<tr>
<td>Net assets released from restrictions</td>
<td>(353,237)</td>
<td>(491,542)</td>
</tr>
<tr>
<td><strong>Increase in temporarily restricted net assets</strong></td>
<td>(113,937)</td>
<td>(88,400)</td>
</tr>
</tbody>
</table>

### Total increase in net assets

| Total increase in net assets               | (131,698)| (187,555)|

### Net assets, beginning of year

| Net assets, beginning of year              | 1,927,505| 2,115,060|

### Net assets, end of year

| Net assets, end of year                    | 1,795,807| 1,927,505|
CORPORATE PARTNERS

The Corporate Partnership Program provides a way for the Spondylitis Association’s pharmaceutical partners to positively impact the lives of those affected by spondylitis by contributing to the organization’s general operating budget. SAA also receives additional corporate support for special programs.

PLATINUM PARTNER 2011

Centocor Ortho Biotech Inc.

GOLD PARTNER 2011

Abbott

A Promise for Life

FOUNDATIONS

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

Blank Charitable Foundation
Cecile & Fred Bartman Foundation
Community Foundation Alliance
Davidson & Gentry Families Fund
Ellen & Marshall Cole Philanthropic Fund
Jean and E. Floyd Kvamme Foundation
Kautz Family Foundation

Koven Foundation
Murdy Foundation
Silicon Valley Community Foundation
Sydney T. Levenson, MD Foundation
Thistle & Rose Foundation
Young Foundation

GIFTS IN KIND

The Spondylitis Association would like to thank the following supporters who have donated in-kind goods and service between July 1, 2010 and June 30, 2011.

Dennis Bolton Enterprises, Inc.
Tom Contrino
Alan B. Hirsh
Quexion, LLC

Jay Ross at GR8PCDR, Inc.
Sundance Press
Jennifer Visscher
Over sixty-four percent of the Spondylitis Association’s revenue in fiscal year 2011 came from individuals and family foundations. More than twenty-six percent of that revenue was from Leadership Circle members.

The Leadership Circle is a distinguished group of dedicated individuals who made annual gifts of $1,000 or more between July 1, 2010 and June 30, 2011. Through their participation, Leadership Circle member contributions support all SAA programs. Their generosity demonstrates the strength of their commitment to the mission of SAA and underscores the important role that Leadership Circle members play in helping to sustain the organization today and in the future.

**$50,000 AND ABOVE**
Jean & E. Floyd Kvamme Foundation

**$10,000 - $20,000**
Cecile & Fred Bartman Foundation
Kautz Family Foundation
Herb & Barbara Shear
Diane Williams

**$5,000 - $9,999**
Stephen & Grace Becker
Elizabeth & Daniel Davis
Leslie Kautz
Deidra & Niki Krutop
Murdy Foundation
Laurie M. Savage
Barbara Van Alstine

**$1,000 - $4,999**
Charles Adams, Jr
Keith & Celia Arnaud
Vasser Bailey
Brian Berman
Blank Charitable Foundation
Michael Borghi
Harry & Jane Bruckel
Hu Chao
Tom Contrino
Judy Coseglia
George Cuccia
Davidson & Gentry Families Fund
John P. Davin
Chris Deininger
Nero Deliwala
Margaret & Bob Ferrarone
Judy Fiskin
Britt-Louise Fletcher
Andrew & Lauren Forbes
Alan J. Fraser
Susan Gallagher
Craig Gimbel, DDS
Bill Giser & Margaret Locke
Susan & Alexander Goldberg
Eric Goldstein
Charles Grisemer
David Hallegua, MD
Laura Hamilton
Elmer & Arlene Hansen
Dr. Robert & Barbara Hasty
Steve & Ann Herendeen
Todd & Kathleen Herzog
Kenneth Honer
Jeff Horn
Charlotte Howard
Ann Howat
Stephen & Michele Hunter
Wallace Hwang
Margo & Stanley Itskowitch
Susan & Stephen James
Daniel Kautz
Ted Lambrinos
Jonathan Lathrop
Brian Lucas
Brian MacKenzie
Brian R. Mann
Brian Mason
David Eric Moore
Nancy & Dan Morgan
Jane Morrill
Karen Nickerson

Maria & Martin Nyvall
Paula Olsiewski, PhD & John H. Healey, MD
Catherine Otto, MD
Michael Patterson & Toni Cole
Donald Petersen
Michael Pianin
Mukesh Prasad
Paul Provost, RPh
Timothy Quinn
Luther Ragin, Jr & Deborah Fish Ragin, PhD
Nancy Regan
John Reveille, MD
Carol & Frank Robl
Cathy Rutter
Laurie Saylak
Laura G. Schafer
Ronald Schanze & John Pritchett
Jeffrey Schmitt
Karrie Shogren, PhD
Silicon Valley Community Foundation
Traude Steidlmayer
Thistle & Rose Foundation
Barry & Christine Tobias
Alison Toth, MD
Fadlo & Justine Touma
Robert W. Ulrich, PharmD
Robert Wick
Tom Wilkinson
Janet & David Williams
Hilary Wilson
Steve & Mimi Young
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 the organization’s most valued supporters, members of the *Quest Legacy Society* have decided to continue the fight against spondylitis beyond their own lifetimes, bestowing upon all those affected by the disease, their families, and future generations a reason for hope. Information about the *Quest Legacy Society* can be found on SAA’s website at:

[www.spondylitis.org/quest](http://www.spondylitis.org/quest)

*SAA is honored to recognize these Quest Legacy Society members:*

Stephen & Grace Becker  
Cecelia Bunch  
Gary Cadle  
R. Means Davis, Jr  
Richard & Deanna Day  
Harvey R. Derscheid  
Alan J. Fraser  
David Fulle  
Dr. Robert & Barbara Hasty  
Phil & Eileen Hipe  
Jennifer Layden  
Brian MacKenzie  
Christopher Oleksy  
Nancy Regan  
Laurie M. Savage  
Katherine Sprouse  
Gregg Umek  
Darren B. Wolpert

*In fiscal year 2011, planned gifts were received from The Estate of Raymond and Mary Jean Cassidy (given through the Community Foundation Alliance).*
Gifts from individuals, whether membership dues or straight donations, are the cornerstone of SAA’s financial support. This year, individuals’ gifts to SAA accounted for 61% of the organization’s overall revenue.

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

Also note that many donations given through workplace campaigns such as the Combined Federal Campaign and United Way are received anonymously. If you have given through one of these campaigns, please let us know so that we may acknowledge your generosity and ensure that you continue to receive your member benefits.

### $500 - 999
- Alan W. Anderson
- Robert & Christine Baldoni
- John Barnes, MD
- Gerald & Estelle Becker
- Bradley & Marla Bockhorst, DMD
- John Brimsek
- William Buehler
- Donald & Judy Bunin
- Joanne Chamberlin
- Curtis Cheatham
- Judith & Ed Christian
- Sharon T. Clement
- Robert A. Colbert, MD PhD
- Community Foundation Alliance
- R. Means Davis, Jr
- Nancy Davis
- Deb Detiveaux
- Raymond & Mimi Diller
- John & Cathie Duniway
- Jane & Dane Farnworth
- David Fulle
- Jason Ganetsky
- Douglas Gant
- Robert & Rhonda Gendron
- Alan Glaser
- Buffa Hargett
- Mark & Ruth Holland
- Shari & Craig Jankowsky
- Bradley P. Johnson
- Marjorie B. Joseph
- Muhammad Asim Khan, MD
- Rachel Knopoff
- Koven Foundation
- Eric Laube
- Donald G. Lee
- Sandor Lehoczky
- David M. Lintern
- Jacqueline & Richard Litchfield
- Heather Lopane
- Anna & Chris Lynch
- Florence Magassy
- Susan M. Mandell
- Robert Mann
- Mick Mars
- Dorothy Martin
- Eddie & Victoria Mazanec
- Inna Mednikov
- Joshua Nadel
- Newport Business Media
- Ford & Catherine Nicholson
- Hazel Nix
- Christopher Noble & Chris Farrow-Noble
- Betty Opbroek
- Sam Paul
- Candy Payne
- Diann Peterson
- Len Potoshman
- Michael Rothermund
- Aaron Rupert
- William Rutherford
- Milt B. Schulle
- Lee & Gail Silver
- Roger Tachuk
- Jayne & Kelly Tien
- David Towery
- John Trotter
- Stephen Utkus
- David & Christine Vernier
- Mary Vieth
- Dirk Visser
- Robert W. Warren, MD PhD
- Andy Weisenfeld
- Tom & Barbie West
- Joe Wickwire
- Maria Wilcox
- Young Foundation

### $250 - $499
- Caroline & Rob Abrom
- Jake Aronov
- Ellen Ashby
- Betty & George Baffa
- James Barnett
- Jennelle Barton
- Michael Barton
- Michelle Beggs
- Bryan Bradbury
- Rafael Bras
Garry M. Brown
Duncan Brown
Jeanne Bruno
Thomas Bunin
James Burke
Paula Bustamante
Larry Caldwell
Lisa Calladine
Lynn Camino
Web Carr
Ellen Carroll
John Cella, III, MD
Stephen Chappell
Gaylon & Debbie Childers
Jennifer Clark
James & Louise Clark
Michael A. Cohen
Hoble Cohen
Don Cohen
Ellen & Marshall Cole
Philanthropic Fund
Jim Colombo
Rick Coscia
Aldene & Kenneth Croswell
Dennis Dempsey
John Digianni
Mark Dominey
Amy & Garr Dorey
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