

Dorothy Moriarty
Dennis & Faye Mosco
Eric and Heather Mosco
Rose Moss
John & Cindy Mullally
Bernard Nadeau
Michael Nash
Elaine Nebel
Stacey Nicholson
Ramon Nieves
James Nothem
Nova Ocean Marine, Inc.
Mark & Susan Nowinski
Patrick & Nancy O'Connor
Timothy & Joannah O'Connor
Terry & Kristine O'Mara
Tom O'Neil
Laurie & Danny Oldham
Kristine Olson
Todd Olson
Osiris Records
Mark Ostaseski
Steven Panagiotes
Randall J. Parker
Robert Parker

Michael & Merna Peden
Paul & Barbara Penfield
Perry Hall Auto Body
Chuck & Jennifer Pierce
Curt & Gerry Pindler
Larry & Susan Piper
Anthony R. Pizzuto
Fred Potts
Dale F. Powers
Dorothy Ann Proctor
Adria Prosser
Purofirst of Metro Washington
Putty Hill Body Shop
James Quattlebaum
Ed Rea
Bob Regener
Maria Reiling
John Reith
Elaine Renfro
Jim & Janet Riehl
John Rieth
Robert W. Nock Insurance
Agency, Inc.
Rotary Yacht Squadron of the
Chesapeake Bay

Jack & Janice Rubis
Rupp, Baase, Pfalzgraf &
Cunningham
Garry Rust
William Ryan
Andreas Schafer
Jim Schebil
ServiceMaster of Baltimore
ServPro of Carroll County
Servpro of the Southtowns Ltd.
Mark & Susan Setter
Bob & Clare Sevenich
Randall & Linda Shepard
Michael Sheridan
Richard & Claire Shonter, Jr.
Smallwood Small Insurance
Allen C. Smith
William J. Smith
Eric W. Snyder
Mallon Snyder
Alan & Roz Spier
Split Rock Energy LLC
Chuck & Karen Spriggs
Stainbrook Insurance Agency
Statland & Katz, Ltd.

Steve Heppler Fund
Peter Strain
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Mark Talbott
The Clark Agency
The Give Thanks Bakery Ltd.
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The Lodish Family Foundation
The Math Works, Inc.
The McArdle Agency, Inc.
The Reny Company
The Ryan Agency
J. Earl Thompson
Seton Thompson
Three Rivers Planning &
Development
Lila Tickman
Jason Tighe
Tisch Foundation, Inc.
Trader Jack's Flea Market
Cheryl Troiano
Russell F. Tromley
Walt Tucker
Peter & Richie Vanderwarker
Daneen Veres

Thomas Vicarel
James Voyles
Alan Wahl
Scott & Jackie Wallace
Jeffrey Wangler
Frank & Lois Wankerl
Nancy Watts
Joe & Cindy Weir
Amy Whipple
George & Betsy Whitehead
Robert Whitehead
Laura Whitley
Julie Wiedeman
Frank & Peggy Wies
Christopher Williams
Robert & Denise Williams
Randy Wilson
Larry & Cynthia Wind
James Wood
Valerie Wouters
Anonymous
Young Chevrolet Oldsmobile
Cadillac, Inc.
Michael S. Zavasky

In Our Laboratory:

[**ALS Therapy Development Foundation** > Annual Report 2003]

ALS TDF Scientific Advisory Board

- :: Dr. Harvey Lodish (chair)
- :: Dr. Robert Brown
- :: Dr. David Housman
- :: Dr. Ken Kosik
- :: Dr. Peter Lansbury
- :: Dr. Hidde Ploegh
- :: Dr. Hazel Sive

Clinical Research Partners

University of California at
San Francisco ALS Center

- :: Richard K. Olney, M.D.
- :: Catherine Lomen-Hoerth, M.D., Ph.D.
- :: Dallas A. Forshew, R.N., B.S.N.

ALS TDF Board of Directors

- :: James Allen Heywood (president)
- :: Spiros Jamas, Sc.D. (chair)
- :: George Hughes (clerk)
- :: Keith Melanson (treasurer)
- :: Dennis A. Ausiello, M.D.
- :: Steve Fowler
- :: John Heywood, Ph.D.
- :: Timothy Hines
- :: Jennifer Huntington
- :: David Searls, Ph.D.

"I work for a large pharmaceutical company, and I certainly recognize the need for critical mass in the very challenging job of making safe and effective drugs. The people I work with are extremely dedicated to this task. However, I also recognize that, with diseases such as ALS, a small but highly motivated and imaginative group of people can make a big difference, when they are absolutely single-minded about facilitating a cure and at the same time deeply involved in education and support of patients and their families. For me, involvement with ALS TDF is a perfect complement to my 'day job.'"

David Searls, Ph.D., Vice President and Director,
Bioinformatics, Smithkline Beecham Pharmaceuticals



Moving Ahead >

ALS TDF moved into the Biotech Cluster in Cambridge, Massachusetts in 2003, continuing to lead the way for drug discovery on behalf of patients living with ALS.



Dear Friends,

As I write this letter, ALS TDF is completing its fifth year of operation. In many ways, I feel that 2003 represents the completion of our first chapter. Today, we are a state-of-the-art biotechnology research center. Today, we have made significant advances in our understanding of ALS. Today, we are better positioned than ever to deliver on our mission to slow, arrest, and cure ALS.

My brother Stephen has had ALS for six years now. Although his body in large part has been destroyed by ALS and he now uses a ventilator, his mental strength and outlook continue to astound those who know him. Stephen is connected to the world through some amazing technology, and he is an active member of both our family and society. He spends time doing what he loves—being a good father to his three year old son Alex and continuing his craft as a custom builder.

Three years ago, we began a project to restore the carriage house behind my home that was virtually falling apart. Wood was rotting, the roof had huge holes and remnants from the previous owner were everywhere. Stephen knew that it was the perfect project for him—he could transform it into a work of art. Today, the carriage house has a solid structure, new three-inch-thick cedar doors, and windows overlooking

the woods behind it. There is still much to do inside, but the first stage of construction is complete.

This is similar to ALS TDF. Our first few years laid the extensive groundwork for the program that exists today. We started with a vision and began to build an infrastructure to cure ALS. Today, we have made great strides and have many reasons to be proud. We remain focused on developing therapies for today's patients and because of our progress, we now work from a much stronger foundation—with clear direction and new tools. We now have a controlled, systematic process for drug studies in ALS mice—the leading program in the world. Our translational research program collaborates with leaders in the biotech and pharmaceutical industries and our unique business model allows us to do so without conflict or delays. We have made significant scientific advances into understanding both the underlying mechanisms of the disease as well as potential treatments—exemplified by the start of our clinical trial at the University of California at San Francisco.

There is still a great deal for us to accomplish, but I know that today ALS TDF is more efficient, driven and focused than ever before. Each day we better equip ourselves to fight this disease, learning from both our successes and failures. Most importantly, we do this with the lives of ALS patients at the forefront of our minds.



James and Stephen Heywood

Your support makes ALS TDF a leader in ALS research and gives hope to my family and to thousands of others with ALS everyday. With your passion and leadership, we are writing the next chapter together—I have never been more excited by the prospect of what lies ahead.

Thank you,

James Allen Heywood
d'Arbeloff Founding Director



Gwendolyn Wong, Ph.D., Director of In Vivo Pharmacology

OUR RESEARCH

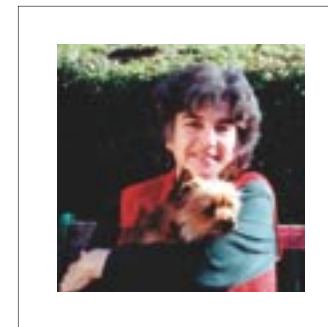
ALS TDF's Research & Development team initiates clinical trial for ALS at University of California at San Francisco (UCSF).

The ALS Therapy Development Foundation and the Larry L. Hillblom Foundation, Inc. announced the initiation of the Ritonavir and Hydroxyurea clinical study in 2003, to be held at the University of California at San Francisco (UCSF). The study will enroll at The ALS Center at UCSF under the direction of ALS specialist Dr. Richard K. Olney. Ritonavir and Hydroxyurea will be tested separately; however, future studies may look toward combining the drugs.

Ritonavir and Hydroxyurea emerged from ALS TDF's drug screening pipeline and are both FDA approved. ALS TDF strives to test approved drugs whenever possible, shortening the time between discovery and the clinic.

Ritonavir is an antiretroviral protease inhibitor used for the treatment of HIV infection. Hydroxyurea is approved for chronic myelogenous leukemia and sickle cell disease. Hydroxyurea is also used off label for malignant gliomas since it is very CNS penetrant. Each compound achieved at least two positive studies before being selected as clinical candidates. The effect of each compound exceeded the effect of Rilutek under identical test conditions.

This study is jointly funded by ALS TDF and the Larry L. Hillblom Foundation, Inc.



Vanna Forrester

Courageous Pioneer & Fighter

Vanna Forrester, the mother of ALS TDF's Vice President of Drug Development, Sean Scott, passed away in October 2003 after a difficult three-year battle with ALS. Vanna's first symptoms appeared in early 2000, when she had difficulty breathing after climbing stairs. While her prognosis was grim, she chose to fight aggressively.

With determined assistance from neurologists Rick Olney and Josh Novic, and information from ALS TDF's research program, Vanna bravely tried many experimental treatments. Her experimentations directly caused the discovery of Hydroxyurea for ALS. ALS TDF tested Hydroxyurea after her first anecdotal reports of benefit and found that the drug showed a positive effect in multiple animal studies.

Vanna leaves as her legacy a clinical trial for Hydroxyurea, which is in its early stages at UCSF. She wanted nothing more than to see others benefit from her efforts and to avoid the horrors she faced. She was a hero to all who knew her or knew of her efforts. Sean joined ALS TDF in 2000 in an effort to help her, setting an aggressive pace for the research program and bringing an appropriate sense of immediacy to the decisions we make.



Nancy Kelly, Director of Business Development, and Sean Scott, Vice President of Drug Development

OUR LAB

ALS TDF's laboratory continues its aggressive pace while fine-tuning its drug studies for the most accurate results.

In 2003, ALS TDF completed the first and most difficult phase of its drug screening efforts, emerging with a viable strategy for attacking the disease. To date, the R&D team has run close to 250 animal studies—150 of those studies were full size drug studies and the remaining 100 were pilot studies for toxicity, drug exposure, and biomarkers.

Over the course of these studies we have identified variations in lifespan of the SOD mouse due to litter, gender, batch, and protein load. We have introduced appropriate controls for these variables and have introduced endpoints sufficient to detect drug effect. We have developed new surgical techniques and skills to deliver drugs to the SOD mouse—resulting in the identification of various challenges in delivering drugs to the

spinal cord, which may be preventing drugs from working as well as they could. The completion of this initial phase of research leaves us poised to target the disease in new and more powerful ways.

In 2004, there are fundamental goals which will build on the foundation we've laid. First, we must continue and expand the search for an ALS biomarker or surrogate marker to help detect drug effect; the search for these markers will be integrally tied to our search for better disease targets as we refine the hits we have in hand. Additionally, we look forward to solving the drug delivery and exposure issues in this mouse model that have gone unrecognized in this field of research until now.



Dr. Harvey Lodish

Chair of ALS TDF's Scientific Advisory Board

ALS TDF proudly announced the addition of a Scientific Advisory Board in 2003, with Dr. Harvey Lodish presiding as Chair. Lodish, a renowned leader in the field of membrane biology and his results have important implications for the treatment of cancer, diabetes, heart disease, and obesity.

"I've been very impressed by the way ALS TDF has uncovered promising drugs by testing a mouse model of ALS. My hope is that over the next year, the Scientific Advisory Board, working together with ALS TDF scientists, can devise new assays that are more rapid and less expensive and that will allow us to screen many more compounds for their potential as ALS TDF therapeutics."

Lodish, a founding member of the Whitehead Institute for Biomedical Research, and Professor of Biology and Bioengineering, MIT, joined the MIT faculty in 1968 and has been a professor of biology since 1976. He earned his Ph.D. at Rockefeller University in 1966 and was elected a Fellow of the American Association for the Advancement of Science in 1986, a member of the National Academy of Sciences in 1987, and a fellow of the American Academy of Arts and Sciences in 1999. Lodish serves on advisory boards for the Cleveland Clinic Research Institute and the Division of Biology at CalTech. Currently, he is the President of the American Society for Cell Biology.

Harvey Lodish Photo: Sam Ogden



> Jack Orchard and wife Eve Tetzlaff



> Betty Luke



> Chris Pendergast

OUR FUNDING

ALS TDF leads the way in drug discovery today due to a dedicated group of individuals and families that have banded together to help us reach our goal to slow, arrest, and cure ALS – thank you. Together, we have raised more than \$14 million since 1999.

Jack Orchard ALS Foundation

Jack Orchard got involved with ALS TDF in the summer of 2002 after his friend told him about our unique approach. After visiting ALS TDF later that year, Jack was most intrigued by the drug screening model that couples the scientific approach with a real sense of urgency.

“At ALS TDF, where the focus remains on screening existing compounds known to be effective in treating conditions that share a biochemical pathway with ALS, hope accompanies every assay,” says Jack.

Today, the Jack Orchard ALS Foundation has raised nearly \$1 million to create awareness and fight ALS, directing almost \$150,000 of those funds to ALS TDF in support of our unique research program.

Betty Luke

Betty Luke went to school with Peggy Heywood, Stephen and Jamie’s mother, in the 1960’s at the Simmons College School of Social Work. Over the years, she, Peggy, and a few of their classmates would still get together for luncheon and kept in touch. When Betty heard that Stephen was diagnosed with ALS, she wanted to help and support the family and his friends when they founded ALS TDF.

“I’m fond of Peggy, and I had met John and her sons. I felt compassion and admiration for Stephen and for this very fine cause to find a cure or at least slow this disease. The story of this important undertaking is compelling and inspirational,” said Betty.

Betty has contributed generously to ALS TDF’s research and lives in North Andover, Massachusetts with her dog Gracie, a Welsh Corgie, and is a psychotherapist with a private practice in Andover.

Ride for Life

Ride for Life’s annual event draws scores of patients that ride their electric wheelchairs from Manhattan to the tip of Long Island to raise money to fund research for a cure along with public awareness about ALS.

“Basic biological research will someday unravel the mystery of ALS,” says Christopher Pendergast, President and Founder of Ride for Life. “Most patients living with ALS will die while waiting for this type of research to bear fruit. On the other hand, the type of research that is the main focus at ALS TDF is identifying currently available agents that will slow or stop the progression. Much like how aspirin was accidentally discovered to be a powerful treatment for cardiovascular disease.”

Ride for Life donated nearly \$50,000 to ALS TDF’s laboratory in 2003, and more than \$150,000 since ALS TDF’s inception.

>> Ronya Kozmetsky

Ronya Kozmetsky committed a \$1 Million gift in 2003 to ALS TDF, representing her and her family’s commitment and desire to have the greatest possible impact on the advancement of a cure for ALS. The Kozmetsky family was introduced to ALS TDF in the fall of 2002. The patriarch of the family, Dr. George Kozmetsky, had ALS, and they decided to get involved in the ALS research arena. Dr. Kozmetsky, who passed away in April 2003, was the co-founder of Teledyne and the former Dean of the McCombs School of Business and Graduate School of Business at the University of Texas, Austin.

“I hope that Jamie can beat the odds for his brother,” says Ronya Kozmetsky. “Dr. Kozmetsky really appreciated and applauded the efforts of the ALS Therapy Development Foundation on his behalf and on behalf of all those living with ALS.”



Mrs. Ronya Kozmetsky and the late Dr. George Kozmetsky

“MY HUSBAND DIED ONE YEAR AGO FROM ALS — IT’S A HORRIBLE, HORRIBLE DISEASE. HE AND MY SON GREG GOT INVOLVED FIRST WITH ALS TDF, AND NOW IT’S IMPORTANT TO ME FOR GEORGE’S SAKE TO HELP ELIMINATE THIS DISEASE SO THAT NO ONE ELSE HAS TO SUFFER.”

- RONYA KOZMETSKY -



> The Nodaros Family



> The Gerbick Family



> The Douglass Family Christmas Tree Sale



> Tony Tucker with his daughter

FAMILIES FUNDING OUR RESEARCH

Andy Nodaros & Family

Andy Nodaros originally found ALS TDF through his friend Dave Eltschlager. Both patients, they came together to ALS 101 in 2002 and met the ALS TDF team. From then on Andy was determined to help support our research. A charismatic and charming personality, full of wit, Andy convinced reporters, local businesses, and friends and family to support ALS research in a variety of ways. Sadly, Andy passed away in 2003 but continues to be a catalyst for change, even now.

Today, Dave continues this work in his memory, along with a group of people in the Pittsburgh area now connected through ALS. The families and friends of Walt Knopp, Jon Russell, Scott Furner, Andy, and Dave continue their work toward a cure. Collectively, this group of ALS TDF supporters raised nearly \$60,000 toward the research in our laboratory in 2003.

Paul Gerbick & Family

Paul Gerbick was diagnosed in January 2002, and shortly afterward attended ALS 101. He came away enthusiastic about ALS TDF's mission, and in June

he and his family put together their first letter-writing campaign to their friends and community. He later decided to auction off his beloved classic red Thunderbird to benefit ALS TDF's research as well.

Wife Betsy, daughters Emily, Caroline, and Abigail all supported Paul's efforts to raise money for a cause he believed wholeheartedly in. His brother Lawrence and sister Veronica contributed by designing raffle tickets and educating others on ALS and why they should contribute.

At Paul's workplace, TAC Worldwide Companies, his coworkers and friends all sold and bought tickets. Paul passed away in the summer of 2003, but the effort continued on his behalf. By the end of the year, the Gerbick family raised more than \$60,000 in support of ALS TDF's research program.

The Douglass Family

When Ken Douglass was diagnosed with ALS, he and his parents searched the Internet for answers. They soon recognized drug companies would not develop and bring a drug to market in time for him, and appreciated ALS TDF's focus on treatments that could benefit patients today.

Dick and Vivian Douglass soon decided to use the tree farm on their retirement

property as a way to fund research. For the last three years, the family has held a Christmas Tree Sale on their property, raising more than \$10,000 in 2003. Today, Ken has a vent and feeding tube but is doing quite well, active through his computer. His family works together to care for him, particularly his wife Mary.

The Tucker & Haley Families

Tony Tucker was diagnosed with ALS nine years ago, and continues to live at home, cared for by his loving wife, friends and family. Though he encounters the daily struggles with ALS, he manages to find beauty and enjoys being surrounded by a large community of family and friends, recently watching his daughter, Bridget, enter college.

When Tony was diagnosed, his wife's family and his own gathered around him in support. Today, Tony's brother, Walt Tucker, heads up Golf 4 ALS, an annual golf tournament to support the laboratory at ALS TDF. The Haley family, Tony's in-laws, also started an annual event nine years ago. Together, the two events raised more than \$40,000 in 2003—honoring the life of Tony and funding critical research at ALS TDF.

>> Driving 4 Life

On June 16th, 2003, professional golfer Tom Watson and his caddy Bruce Edwards along with his wife Marsha announced the Driving 4 Life campaign to support ALS TDF's research on the Today show. Bruce was diagnosed with ALS in 2001, and had been Tom's caddy for 27 years. Just days before, Tom Watson had stunned the golf community with a 5-under par 65 in the opening round of the U.S. Open.

Another ALS patient and former PGA professional, Jeff Julian, along his wife Kim, also led the way for Driving 4 Life, and held an event to support ALS TDF. By the end of the year, the campaign had raised nearly \$1 Million to fund ALS TDF's laboratory, and fans around the country banded together in support of Bruce Edwards and Jeff Julian through individual donations, sponsorships, golf tournaments, and other creative ways of raising money for ALS TDF. Bill Denney, Darren McBratney, and Mark Burrow formed Driving 4 Bruce—a team that drove a golf cart from their hometown in Kansas City to Pebble Beach, California in honor of Bruce, recruiting sponsors and donations in their honor. ALS TDF and Driving 4 Life thank all those who have invested in our research and in today's ALS patients.



> Tom Watson and Bruce Edwards



ALS Therapy Development Foundation

"Doctors tell us (a cure) may be out there, but money is needed for critical research. That's where Driving 4 Life will help."

TOM WATSON

"Over the past two years we have searched for and gone after every possible lead. That search led us to ALS TDF. We have not met a more aggressive, determined organization than the team at ALS TDF."

JEFF JULIAN



> Jeff Julian

FINANCIAL REPORT 2003

	2003	2002	2001
Assets			
Cash	375,845.00	1,265,312.00	890,304.00
Accounts Receivable	1,100,000.00	-	-
Investments	-	574,260.00	974,999.00
Fixed Assets, Net	130,209.00	151,340.00	117,763.00
Other Assets, Net	63,802.00	40,287.00	18,912.00
Total Assets	1,669,856.00	2,031,199.00	2,001,978.00
Liabilities & Net Assets			
Liabilities	627,983.00	435,021.00	225,126.00
Net Assets	1,041,873.00	1,596,178.00	1,776,852.00
Total Liabilities & Net Assets	1,669,856.00	2,031,199.00	2,001,978.00
Support & Revenue			
Contributions	2,524,562.00	2,240,704.00	2,863,153.00
Grant Receipts	120,000.00	641,185.00	1,000,000.00
Fundraising Events	1,257,055.00	1,524,614.00	58,116.00
In-kind Donations	80,143.00	19,072.00	21,828.00
Interest, Dividends & Gains Realized and Unrealized	9,011.00	(90,158.00)	74,972.00
Total Income	3,990,771.00	4,335,417.00	4,018,069.00
Expenses			
Research & Development	3,649,068.00	3,600,967.00	2,204,794.00
Patient Services	253,258.00	218,503.00	71,639.00
Fundraising	391,463.00	451,439.00	333,795.00
Management & General	251,287.00	245,183.00	372,667.00
Total Expenses	4,545,076.00	4,516,092.00	2,982,895.00
Increase in Net Assets	(554,305.00)	(180,674.00)	1,035,172.00
Net Assets: Beginning of Year	1,596,178.00	1,776,852.00	741,680.00
Net Assets: End of Year	1,041,873.00	1,596,178.00	1,776,852.00

Donors

Leadership Circles

As a member of one of the Leadership Circles, you demonstrate your leadership through annual gifts of \$1,000 or greater. You provide support for ALS TDF's laboratory operations, patient services, and clinical research into potential treatments for ALS. Thank you to all members of our Leadership Circles who continue to enable ALS TDF to make significant strides toward slowing, arresting, and curing ALS.

Founders' Circle

\$100,000-plus annually or \$1 million or more cumulative giving

ALS Hope- The Chris Hobler/James

Maritz Foundation

Anonymous

Alex & Brit d'Arbeloff

Dee Dee Fornengo Fund

Driving 4 Life Effort

Fran Delaney Fund

William Eckhardt

George & Daphne Hatsopoulos

Ronya Kozmetsky

Mary Lou Krauseneck Fund

Stephen Milne Society

continued on next page...

Family & Friends Community Partners

The Family and Friends Community is a key part of ALS TDF's fight to develop treatments and a cure for ALS.

This fast-growing group of family and friends has increased our motivation and strength to find a cure for ALS and together raised approximately \$2 million in 2003. Thank you to each and every person that made the following efforts possible, and exhibited ALS TDF's values of Courage and Love.

Bryan & Mark Allain Fund
ALS Charitable Fund, Inc.
Andrew Nodaros Fund
Anthony Tucker
Branman/Weintraub
David Green
David Reinstein Fund
Dean Adraktas Fund
Deane Jewell Team
DeeDee Fornengo Fund
Dick Douglass Family Effort
Driving 4 Life Effort
Elliot Macht
Fran Delaney Fund
Friends of Harry Gianneschi
Friends of My Heart Regina Bisson Fund
Heywood Friends
Jack Orchard ALS Foundation
Jason Becker Fund
Jean Angell Fund
Jeff Ureel Fund
Jennifer Nichols
Jim & Susan Airey ALS Fund
Jim Raspanti Fund

Joe Shambo
Larry Manes
Laura Gould
Luc Blais Fund
Marie Garratt Memorial Fund
Mary Lou Krauseneck Fund
Matt Dowd Fund
Paul Gerbick Fund
Peggy Lanza Fund
Pittsburgh Families Efforts
Ruth Kulow Fund
Sharon Stahly Fund
Silverback Technologies
Stanfield Family ALS Fund
Stephen Heywood
Stephen Milne Society
Steve Fowler Fund
Steve Heppler Fund
Steve Lewis Fund
Ternet Friends Fighting ALS Fund
The Carter Family ALS Research Fund
The Dunphy Sister's Fund
The Heather McDonald Fund
Vanna Forrester Memorial Fund

Walter Knopp Fund
Will Hubben ALS Research
Wolfmom Family Fund
Alan Inglis Fund
ALFH Fund
Ben Heywood Effort
Doug Duren Effort
John Russell Fund
Judy Oakley Effort
Kathy Rivers Fund
Kenny Alber Fund
Lauretta Stott Fund
Maes Family Fund
Paul Barrett
Phil Rapin Fund
Steve Carlin
Trude Klugman
Marty Stayer ALS Research Fund
Tom Larsen Challenge
Jan Richardson Fund
Jack Hoffman
Darin Lily Effort

continued from previous page...

Governors' Circle

\$25,000-plus annually

ALS Family Charitable Foundation, Inc.
Anthony Tucker Effort
Steven Brouman
The Carter Family ALS Research Fund
Connecticut Friends for ALS Research
David Reinstein Fund
Elliot Macht Effort
HMS Host
The Jack Orchard ALS Foundation
Jean Angell Fund
Matt Dowd Fund
McDonald's Corp
Paul Gerbick Fund
PGA TOUR, Inc.
Pittsburgh Families Efforts
Ride For Life
Christy Sloan/ The ALS Race for Research

Jennifer Huntington
Jim & Susan Airey ALS Fund
Jim Raspanti Fund
Gail Kendall
Tom Larsen
William and Betsy Leitch
Betty Luke
Stephen J. McKendrick
Palguta's Barracudas Golf Outing
Mark Reed III
David Searls
John & Lola Soback
Stanfield Family ALS Fund
Stephen Heywood Fund
Steve Lewis Fund
Ternet Friends Fighting ALS Fund
The Leonard & Claire Tow Charitable Trust Inc.
The Thorson Foundation
Woodbine Bend/Jenny Stoddard
Brian Wruble

Associates' Circle

\$5,000-plus annually

Allain-Lebreton Co. LLC
Apex Foundation
Bruce & Lissa Barron
Bryan & Mark Allain Fund
Harold & Phyllis Brouman
Caddie Master Enterprises, Inc.
Elizabeth Courtney
Gary & Donna Dake
DGA Builders, Inc.
Richard & Vivian Douglass
Bruce & Veronica Erhart
Foxwoods Resort Casino
Friends of Harry Gianneschi
L.R. & Caroline Gardiner

Gary Player Foundation, Inc.
General Atlantic Service Corp. MG
Amy Goldberg
John & Peggy Heywood
Rajive Johri
Bob & Mary Leisure
Larry Linden
Herbert L. & Suzanne S. Lynskey
MacDonald Illig Jones & Britton LLP
MasterCard International
Patricia McCabe
Newburg & Company, LLP
Jennifer Nichols Effort
Olson ALS Foundation, Inc.
"Ponaks"- Elizabeth Courtney
Radford Foundation
SAFELITE Glass Corp.
Joerg Schwalfenberg
Tom Sharp
Sharp Residential Builders & Developers
The William G. Gilmore Foundation
Robert & Dany Theuerkauf
Anonymous Foundation
Gerard Ventura
Walter Knopp Fund
Waste Management, Inc.
Basil G. Witt

Michael & Jean Artin
Thomas W. Austin
Auto Body Images, Inc.
Joe & Joyce Baclawski
Anthony Balsamo
Yvonne Balsamo
Jim & Kaye Barker
Anonymous Foundation
Bob Basten
Michael A. Beachan
Bryan & Nancy Beaver
Ray Bell
James Berkowitz
Marc Blais
Body-Borneman Associates Inc.
Boizelle Insurance
Richard & Cathy Boskey
Bovenzi, Inc.
George & Leslie Brett
Charles Brodie
Janie Brown
Budow and Noble, P.C.
Nat & Rosemary Buono
Greg & Sally Buss
C. F. O'Connell & Sons, Inc.
Capezio Contractors, Inc.
CCC Information Services, Inc.
Christoph Charle
Orville Chenoweth
Deron L. Cherry
Bill Childers
Jerome & Pam Circo
Clarksville Auto Center, Inc.
CNY Insurance Agency
Stephen & Bernadette Cooper
Mary Corrigan
Luciano Cortese
Barry Coughlin

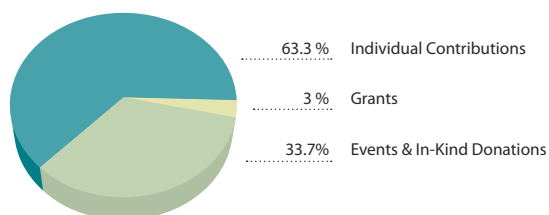
Benefactors' Circle

\$1,000-plus annually

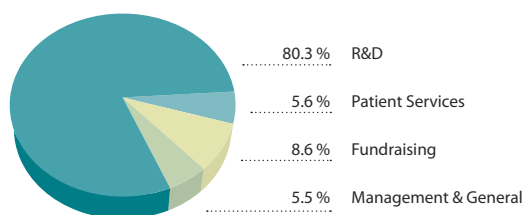
A.L. Howes Agency, Inc.
Michael & Bobbie Agnello
American Insurance Outlet, Inc.
Andrew Nodaros Fund
Gebran & Suzanne Anton
John & Diana Appleton

Robert Cox
Craig Mader Insurance Agency Inc.
James S. Cunningham
Kathleen Davis
Timothy D. Dees
Diamond Auto Glass
Harry DiSimone
Dean Adraktas Fund
Deane Jewell Team
DL Masonry
Jim & Joan Dopp
Stephen & Pam Douglass
Robert Duckett
Bill & Pam Eltschlager
Enterprise Rent-A-Car Company
Faye and Lucille Stewart
Foundation
Fayette Foods
James Ferency
Fife Family Foundation Inc.
First American Title Insurance Co.
Jeannie Moller Fontana
Jay & Susan Forrester
Forry, Ullman, Ullman & Forry, P.C.
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