A Letter from Our Chairwoman

Dear Friends,

This year marked the 30th anniversary of the Hydrocephalus Association (HA). What began in 1983 by a small network of concerned parents has blossomed into the nation’s largest and most widely respected health advocacy organization devoted to meeting the needs of families affected by hydrocephalus. Thanks to your support, we were able to have another banner year, bringing the association one step closer to fulfilling its mission of eliminating the challenges of hydrocephalus.

As a mom of a daughter with the condition, I can’t tell you how proud I am to share the organization’s achievements in 2013. In many regards, it was a year of firsts, paving the way for a promising future for our loved ones affected by the condition. We celebrated the 30th anniversary by honoring our founders, Russell and Emily Fudge and Cynthia Solomon, during the inaugural Vision Dinner held in New York City. This intimate event was designed to raise national awareness and encourage new partners to join us on our journey to find a cure. In addition, commemorative events were held in conjunction with our WALKs, engaging 14,000 participants across the nation. We are grateful to the many volunteers for hosting these special gatherings.

We surpassed our Reason for Hope campaign goal, securing our first $3 million to advance HA’s research agenda. Because of the generous support of our many donors, HA was able to fund an additional $1 million in new research this year. This included the launch of the Adult Hydrocephalus Clinical Research Network (AHCRN), a network of six hospitals and institutes that will focus research on adult forms of hydrocephalus and address the diverse needs of our community across the age spectrum.

Committed to our roots, support and education continue to be important focal points in the work of HA—they are how my family found the organization 10 years ago. HA added webinars to our extensive support publications and products. Our first webinar entitled, “What if It Isn’t Alzheimer’s or Parkinson’s?” was conducted by Dr. Michael Williams and received great reviews. In addition, HA provided over 1,750 hours in direct patient support and continues to leverage a vibrant presence on social media to help patients and their families, and expanded our reach by restructuring the Hydrocephalus Community Networks.

We hosted our first webinar, “What if it isn’t Alzheimer’s or Parkinson’s?” presented by Dr. Michael Williams, director of The Sandra and Malcolm Berman Brain & Spine Institute, Sinai Hospital of Baltimore, during Hydrocephalus Awareness month in September. This webinar was targeted to individuals who have been diagnosed with Normal Pressure Hydrocephalus, caregivers of older adults, medical professionals, and other experts working in the field of hydrocephalus.

I hope you feel proud of what you helped to build and all that HA has accomplished. Without your support, HA would not have been able to achieve many of the successes shared in this report. We look forward to your continued involvement as we seek to fulfill our mission of eliminating the challenges of hydrocephalus.

Sincerely,

Barrett O’Connor
Chairwoman, Board of Directors
Hydrocephalus Association

Support and Education

The Hydrocephalus Association is committed to providing innovative programs to meet the diverse needs of the community. Our services ensure that patients living with hydrocephalus have a place to turn to for resources and peer-to-peer support. In 2013, we published several new educational materials, provided 1,750 hours of one-on-one support through phone, mail, email, and social media to help patients and their families, and expanded our reach by restructuring the Hydrocephalus Community Networks.

Webinar Series

We launched a new webinar series designed to deliver vital information to our community. These interactive, free sessions feature presentations from medical professionals, researchers, and other experts working in the field of hydrocephalus.

Local and Online Community Networks

There are 1 million people affected by hydrocephalus in the United States alone. Expanding our ability to reach more individuals and families impacted by the condition is critical to ensuring that we meet the demands of our community. In 2013, we restructured our support groups to serve as broader-based Community Networks. The networks provide localized support, education and empowerment by hosting educational events, support group meetings, advocacy activities and other gatherings that enable individuals and families to thrive.

HA’s Community Network consists of 39 networks, which have held 80 events hosting nearly 1,500 individuals. In addition, we have 25 active virtual networks with online Facebook Groups consisting of more than 1,200 members. These closed groups allow those living with hydrocephalus, and their loved ones, to connect and share in a private forum in their own region as well as throughout the world. We are grateful to our many volunteers who are involved in raising local awareness and facilitating community gatherings.
Awards and Scholarships

Resident’s Prize

The Hydrocephalus Association Resident’s Prize was awarded to Jennifer Strahle, MD, for her paper, “Early Treatment of Neonatal Intraventricular Hemorrhage-Associated Hydrocephalus with Systemic Deferoxamine,” which was presented at the 2013 Pediatric Section meeting of the American Association of Neurological Surgeons (AANS)/Congress of Neurological Surgeons (CNS) in Toronto, Canada.

This prize is awarded each year to the most promising hydrocephalus-related research paper presented by a neurosurgical resident at the Pediatric Section Meeting of the AANS/CNS. The prize is designed to encourage young doctors to focus their research efforts on advancing the treatment and care of individuals with hydrocephalus.

Scholarships

Since the Hydrocephalus Association’s scholarship program was established in 1994, we have awarded 103 scholarships to deserving future leaders of our community. This year, we were able to offer nine educational scholarships to young adults living with hydrocephalus who have exhibited promising leadership skills and are involved in their communities. Despite the tremendous challenges and obstacles they face, these students continue to excel in the classroom, volunteer and give back to their communities and inspire their peers and all of us. The 2013 recipients are Anna Elise Jones, Juliette Ohan, Ana L. Ubierna, Melissa Bruebach, Jordan Rogers, Timothy Beighley, Emma Jane Maurer, Abigail Fazelat and Lindsay Kerr. These awards were made possible through the support of the following endowed scholarship funds: Gerard Swartz Fudge Memorial Scholarship Fund, Morris L. and Rebecca Ziskind Memorial Scholarship Fund, Anthony Abbene Scholarship Fund, Justin Scot Alston Memorial Scholarship Fund, Mario J. Tocco Hydrocephalus Foundation Scholarship Fund and Giavanna Marie Melomo Memorial Scholarship Fund.

For the second year, we also awarded five additional scholarships through our Teens Take Charge (TTC) Program to Amanda Korcal, Emily Reed, Ruby Poole, Danielle Denise Kukuckwich and Samantha Richardson. TTC’s scholarship awards were made possible through the continued support of the Medtronic Foundation and all the individuals who held events and contributed to the program.

Looking Back...1986

Our first educational booklet, About Hydrocephalus — A Book for Parents, is published. This is our first publication in a line of numerous publications that have become mainstays for individuals and families.

Teens Take Charge

The Teens Take Charge (TTC) program continues to facilitate an active online community of more than 1,500 teens and young adults affected by hydrocephalus, and their siblings.

This forum provides an opportunity for young adults to openly share their journey and provide peer-to-peer support, encouragement and advice. TTC members are involved in various fundraising and awareness activities, from presenting at school assemblies and holding classroom discussions on hydrocephalus, to taking part in local parades and representing the program at our WALK events. In addition, TTC’s advisory council and members publish articles and share their personal stories of encouragement that are posted on our website and various social media platforms to inspire youths living with this condition.
Research Update

This year marked the end of phase one of our Reason for Hope research funding campaign, which started in 2009 to fund the Hydrocephalus Association’s (HA) Research Initiative.

We are proud to report that as a community we surpassed our $3 million funding goal, raising a total of $3,065,046 for hydrocephalus research. Over the past five years, HA has funded a total of $2.7 million in both basic and clinical hydrocephalus research. These research projects are focused on understanding the causes of hydrocephalus and developing better treatments in order to, ultimately, find a cure.

In 2013, we partnered with the Rudi Schulte Research Institute (RSRI) to fund a new research venture. RSRI chose Mark Wagshul, PhD, for his proposal, “MR Elastography: A noninvasive tool for the management of shunted pediatric hydrocephalus.” Dr. Wagshul will utilize this noninvasive MRI-based technique to see if brain compliance can help determine the best treatment option for pediatric hydrocephalic patients.

HA awarded a second grant to Timothy Vogel, MD, of Cincinnati Children’s Hospital. Dr. Vogel’s proposal, “The role of neural progenitor cells in neonatal hydrocephalus,” focuses on cilia, hair-like structures on the surface of neuron progenitor cells (NPCs) in the brain that contribute to CSF movement. He will be exploring how defects in cilia motility contribute to the development of neonatal hydrocephalus.

Looking Back...2009

Mentored Young Investigator (MYI) program is started, with 5 awards made to new researchers interested in hydrocephalus. The MYI program intends to stimulate innovative research such that scientists go on to apply to various programs at the NIH for further grant funding.

HOPE in Progress

SONIA PODVIN, PhD
HOPE: Develop Pharmacological Agent to Treat Hydrocephalus

Sonia Podvin, PhD, is a post-doctoral fellow at the University of California, San Diego. Dr. Podvin was a recipient of one of HA’s Mentored Young Investigator (MYI) Awards. Her MYI grant funded her research as a molecular pharmacologist aiming to develop safe, specific drugs to treat hydrocephalus. Her study investigates an anti-inflammatory hormone called augurin that circulates in CSF. Augurin may be able to control brain hydrodynamics and ultimately may be pharmacologically manipulated to treat hydrocephalus non-invasively.

JOHN KESTLE, MD, FRCSC, FACS
HOPE: Reduce Shunt Infections

John Kestle, MD, FRCSC, FACS, is a Professor and Vice Chair of Clinical Research in the Department of Neurosurgery at the University of Utah, Salt Lake City, Utah. His clinical practice specializes exclusively in pediatric neurosurgery, with a specific interest in pediatric epilepsy surgery. Dr. Kestle has developed a clinical protocol for treating hydrocephalus that has helped reduce post-operative infection rates for shunt surgery by more than 35%. He co-founded and chairs the Hydrocephalus Clinical Research Network, which is supported with funds from the Hydrocephalus Association.

MARK HAMILTON, MD, FRCSC
HOPE: Improve Treatment and Diagnostic Methods for NPH

Mark Hamilton, MD is a neurosurgeon and director of the Adult Hydrocephalus Program at the University of Calgary. Dr. Hamilton is currently the chair of the Adult Hydrocephalus Clinical Research Network (AHCRN). AHCRN is focused on addressing the needs of both transitional patients and adults that acquire hydrocephalus and is being funded by HA. AHCRN will conduct multisite research while developing an electronic image database and cerebrospinal fluid (CSF) biobank. These research efforts are focused on finding better treatments for hydrocephalus and identifying new diagnostic methods for normal pressure hydrocephalus (NPH).
Hydrocephalus Association Annual Report 2013

Hydrocephalus Clinical Research Network (HCRN)

The HCRN expanded its network from seven to nine centers with the additions of Vanderbilt University and the University of British Columbia. Initially, the two centers focused on adding patients to the HCRN registry and participating in the HCRN Quality Improvement Study for post-surgical infections. They are now completely incorporated into the HCRN. Dr. Wellons is heading the center at Vanderbilt and continues to be the lead Principal Investigator on the study of post-hemorrhagic hydrocephalus, which is nearing the completion of its accrual of patients. Dr. Doug Cochrane is heading the center at the University of British Columbia. In addition to expanding to nine centers, the HCRN has been busy publishing three papers in 2013.

Adult Hydrocephalus Clinical Research Network (AHCRN)

The success of the HCRN provided the catalyst for establishing an Adult Hydrocephalus Clinical Research Network (AHCRN).

In 2012, HA began working with distinguished adult hydrocephalus researchers including, neurosurgeons, neurologists, and neuropsychologists to establish an adult research network. The goals of the network are to improve the lives of adults with hydrocephalus, raise awareness of the condition and tackle the unique issues transitional patients, adult congenital patients and other adult-onset patients face. Six sites were selected and progress was made throughout 2013 developing essential data elements and protocols for the network. The six participating researchers and sites in the network include:

- Mark Hamilton, MD, University of Calgary
- Heather Katzen, PhD, University of Miami
- Petra Klinge, MD, Brown University
- Mark Luciano, MD, PhD, Cleveland Clinic
- Norman Relkin, MD, PhD, Weill Cornell Medical College
- Michael Williams, MD, Sandra and Malcolm Berman Brain & Spine Institute, Sinai Hospital of Baltimore

Dr. John Kestle, co-founder of the HCRN, also joined this group effort to study adult hydrocephalus, bringing his rich history in hydrocephalus research to benefit the collaboration.

The development of the adult network is essential for HA’s mission to improve the lives of all people with hydrocephalus. The adult network will increase in importance as pediatric patients transition to adulthood and awareness of adult-onset hydrocephalus increases.

HA Publishing: Hydrocephalus Funding Landscape

HA is committed to advancing hydrocephalus-focused research but little is known about past hydrocephalus funding.

To understand the past funding environment, to create benchmarks to measure against future progress and to quantitatively bolster the case for increased hydrocephalus funding, HA analyzed the amount of funding allotted for hydrocephalus research by the National Institutes of Health (NIH).

The resulting paper, “Hydrocephalus research funding from the National Institutes of Health: a 10-year perspective,” was published online in the Journal of Neurosurgery Pediatrics in December 2013. The study analyzed public funding for hydrocephalus research during the 10-year period between 2002 and 2011. It concluded that hydrocephalus needs more researchers conducting high-impact, innovative research and alternative funding is very important given the challenging public funding environment.
The Hydrocephalus Association’s (HA) WALKs and Special Events had another record-breaking year, raising close to $1.3 million towards our mission.

With 32 local events across the country and more than 15,000 participants, our volunteer-coordinated WALKs grew by ten percent, raising over $1.1 million. Special events, primarily “third party events,” raised more than $168,000 – a 13 percent increase over last year. This was only possible due to the hard work and hundreds of hours of time expended by our wonderful volunteer network – WALK Chairs, Special Events leaders and participants, sponsors and donors. THANK YOU!

The inaugural Los Angeles WALK raised more than $130,000 – the largest first year event in HA history. Chicago’s WALK ranked as the second largest WALK in the country, posting a 33 percent increase from the prior year by raising more than $90,000. The Seattle WALK continued its success, finishing in third place with $85,000.

The Hydrocephalus Association’s special events also demonstrated great results. Team Hydro, participating in Sharkfest for the sixth year, raised $53,000 for hydrocephalus research. Other events included a holiday concert organized by (and starring) Lyn Leach from Lucas, Kentucky, raising $4,000. The 17th annual “Thanks for Running 5K” in Fitchburg, Massachusetts, (held Thanksgiving Day morning) raised more than $6,700. “MX for Children,” an annual motocross event, had another great year by raising more than $43,000. One of our newer events, “Trick or Treat for Hydrocephalus,” showed incredible growth by raising more than $30,000 from a small group of schools. The Hydrocephalus Association was the charity of choice for Atlanta’s Manheim Company, which held a golf tournament that raised over $30,000.

* Third party events are events which are not sponsored by the Hydrocephalus Association but are conducted by local volunteers to benefit the Hydrocephalus Association and its mission.
HYDROCEPHALUS ASSOCIATION AT 30

A History Rooted in Community

In 1983, a small group of parents of children with hydrocephalus living in the San Francisco Bay area came together seeking community and support. They were all patients of Dr. Michael Edwards at the University of California, San Francisco Medical Center. Dr. Edwards and Lori Hilliard, his neurosurgical nurse, graciously hosted a few small, informal gatherings after Emily and Russell Fudge expressed their feelings of isolation and frustration when their son, Gerard, had his second shunt revision. This was a time very different from today. In the early 80s there was no widespread public acknowledgement that the majority of shunts would need to be revised multiple times during a lifetime. There were no programmable or adjustable valves. There were no MRIs. There were no such people as Board certified pediatric neurosurgeons and no Patient’s Bill of Rights. This was a time before the internet, leaving parents, patients, and loved ones to seek out the scant information available in libraries and from their doctors. And there was nowhere to turn for support and guidance.

In 1984, Emily and Russell decided to host a more formal meeting, asking Dr. Edwards to speak about hydrocephalus and inviting a large number of his patients. At that meeting, they proposed the idea to create an organization. With a $1,000 grant from UCSF and Dr. Edwards, the Hydrocephalus Foundation of Northern California was formed, later to be renamed the Hydrocephalus Association. The association was initially organized out of the basement of the Fudge home by a dedicated core group of individuals that included Emily and Russell Fudge, Jennifer Henerlau, Pip Marks, and Cynthia Solomon.

This year we commemorate the 30 year anniversary of the Hydrocephalus Association. We honor the resolve of this dedicated group of parents who recognized the need for an independent, patient-focused organization to support individuals affected by hydrocephalus and to provide their families and friends, who were desperate for information, with life-changing educational materials and group interactions. Thirty years ago this determined group could never have imagined that the Hydrocephalus Association would become the nation’s largest and most widely respected provider of information, support services, advocacy initiatives and private research funding for hydrocephalus.

It is this rich history rooted in our community that drives our core values and mission today.

FIRST ANNUAL VISION DINNER

A Time for Awareness. The Hope for a Cure.

The Hydrocephalus Association’s first annual Vision Dinner, “A Time for Awareness: The Hope for a Cure,” was held on October 11, 2013, at the Apella Alexandria Center in New York City. Generously hosted by Vicki and Craig Brown, major benefactors of the Hydrocephalus Association (HA), the evening united patients, neurosurgeons, scientists, advocates and business leaders to raise the public profile of hydrocephalus and to share the need for greater research funding to cure the condition.

The evening debuted HA’s Reason for Hope video that profiled the unique stories of members in our community. Attendees also had the opportunity to hear from two parents leading the way to further HA’s research initiative and learn about the hope in progress, currently funded by the association. Renowned pediatric neurosurgeon and 2012 MacArthur Foundation Fellow, Dr. Benjamin Warf, was the keynote speaker, sharing his revolutionary new surgical procedure to treat hydrocephalus in the infant population.

The evening also honored three individuals for their tireless dedication to eliminate the challenges of hydrocephalus through their affiliation with the Hydrocephalus Association. Paul Gross, Co-Founder, Hydrocephalus Clinical Research Network, received the Inspiration Award; Marion Walker, MD, Professor, Neurosurgery, Primary Children’s Medical Center, received the Service Award; and Emily and Russell Fudge, Co-Founders, Hydrocephalus Association, received the Leadership Award. United States Senator Charles E. Schumer and New York State Senator John J. Flanagan provided congratulatory letters praising the dedication of the awardees and the progress of the Hydrocephalus Association.
Advocacy in Action

Your Voice in Washington

HA’s national advocacy efforts started in May 2006, when we first organized the National Hydrocephalus Awareness and Advocacy Day on Capitol Hill. We were a small but mighty voice sharing our collective story throughout the halls of Congress. In 2012, recognizing the importance and need for more federal dollars to advance our research agenda, the board relocated the association to the Washington, D.C. area, minutes away from the National Institutes of Health and Capitol Hill. As part of our advocacy efforts, we have trained thousands of members and arranged visits with elected officials to inform them about the impact this condition has on families across the nation.

Advocacy Steering Committee

To help develop and drive the association’s advocacy strategy, a community-led Advocacy Steering Committee was established in 2013. The committee will develop tactics each year to ultimately achieve our primary advocacy goal to ensure the federal government invests adequate funds to support our community. Specifically, the committee will ensure that Congress: understands the breadth and depth of the impact of hydrocephalus on our people; prioritizes funding hydrocephalus research for improved treatments and cures; supports those affected by hydrocephalus with appropriate programs. These goals require targeted awareness campaigns to assure that decision makers understand that hydrocephalus is a condition that affects a broad array of people, is not cured with the insertion of a shunt and is a growing problem with significant burden on our people and the economy.

Coalition Building

HA works with a number of national health advocacy groups to promote the importance of federally sponsored research. We joined ResearchAmerica, the nation’s largest not-for-profit public education and advocacy alliance working to make research to improve health a higher national priority. Through this coalition, we urged Congress and the administration to increase funding for the National Institutes of Health (NIH), Centers for Disease Control and Prevention (CDC), Agency for Healthcare Research and Quality (AHRQ), Food and Drug Administration (FDA) and National Science Foundation (NSF) at levels that keep pace with scientific opportunity. We also advocated for federal funding for global health research and a legislative and regulatory climate that stimulates growth in industry research and development. In addition, HA was active in advocacy meetings and sign-on letters put together as part of the National Health Council (NHC), the American Brain Coalition (ABC), the Rare Disease Legislative Advocates (RDLA), and the National Organization for Rare Disorders (NORD). Topics have included the implementation of the Affordable Care Act, the effects of sequestration on research funding and the Orphan Drug Act.

Congressional Directed Medical Research

HA advocated for the inclusion of hydrocephalus on the list of conditions eligible for funding under the Department of Defense’s Congressionally Directed Medical Research Programs (CDMRP). Great strides were made to educate the House and Senate spending and authorizing committees that allocate money for federal health care research. HA reached out to health and defense appropriators and authorizers to help increase funding for hydrocephalus research at NIH and to create a new funding stream for hydrocephalus research through the Department of Defense’s Peer Reviewed Medical Research Program (PRMRP). We were successful in gaining support on the Senate appropriations side for the hydrocephalus-specific language that Representative Andrews included in the National Defense Authorization Act. Unfortunately, due to partisan battles within Congress over other issues, the language included in a bill passed by the House of Representatives was not included in the final DOD Appropriations bill passed by a conference committee of both the House and Senate.

National Institutes of Health Partnership

The National Institutes of Health (NIH)-National Institute of Neurological Disorders and Stroke (NINDS) hosted their annual Nonprofit Forum. The forum, “Progress through Partnership,” emphasized the role nonprofits play in linking scientists at NINDS with the patient community for better research outcomes. Paul Gross, HA board member and member of the NINDS Advisory Council, served on the planning committee that helped build the agenda for the event. Representatives from over 80 nonprofit organizations had the opportunity to listen to and interact with key NINDS officials, including Dr. Story Landis, director of NINDS, and Dr. Kathy Hudson, deputy director for science, outreach and policy at NIH.
In the News

AARP Informs Members About Normal Pressure Hydrocephalus
Dr. Michael Turner, neurosurgeon, talks about Normal Pressure Hydrocephalus on AARP Radio.

Looking Back…2003
The Visionaries documentary series season 10 spotlights HA on PBS stations across the country.

HCRI Included in US News & World Report 2013
Children’s Hospital Rankings
US News & World Report acknowledges the importance of HCRI’s role in the hospitals ranked in the 2013 Best Children’s Hospitals, which includes stronger treatment protocols and research around the causes and treatment of hydrocephalus.

Famous Rock Star Treated for NPH
ABC News features a story about rock legend, Dick Wagner (lead guitarist for Alice Cooper, Aerosmith and KISS) and his journey with Normal Pressure Hydrocephalus. After receiving a shunt, he was back on the road making music.

The Saturday Evening Post Spotlights Normal Pressure Hydrocephalus
The Saturday Evening Post spotlights Normal Pressure Hydrocephalus (NPH), an often misdiagnosed condition that affects approximately 375,000 individuals in the U.S.

NPH Featured on Boston National Public Radio Show
National Public Radio Boston affiliate WBUR’s Here and Now host, Robin Young, interviews Dr. Mark Johnson, neurosurgeon and head of the Adult Hydrocephalus Program at Brigham and Women’s Hospital, about Normal Pressure Hydrocephalus (NPH).

Leadership Award for Two Founding Members of the Hydrocephalus Association
Long Island Newsday reports on Emily and Russell Fudge, recipients of the Vision Award for Leadership, for being part of the initial group of founding members of the Hydrocephalus Association. They received the award at the Hydrocephalus Association’s First Annual Vision Dinner held in New York City on October 10, 2013.

Cynthia Solomon Receives Spirit of Excellence Award
Cynthia Solomon, a founding member of HA for which she won the Distinguished Achievement Award, is awarded the Spirit of Excellence Award by the California Office of Health Information Integrity (CaOHII) for founding FollowMe, a web-based personal health record company.

South Florida Radio Program Delves Into Hydrocephalus
South Florida Medical Insider radio show on 1230 WBCT interviews pediatric neurosurgeon Dr. Neil Patel and Hydrocephalus Association South Florida Walk Chair Eileen Rodger for an in-depth story about hydrocephalus.

The Right Questions to Ask When Caring for a Parent with Hydrocephalus
Jillian Keenan of The New York Times puts together a list of questions all adult children need to ask of their parents before a critical medical emergency takes place. In her case, her father had Normal Pressure Hydrocephalus.

Olivia Maccoux Dubbed ‘Iron Athlete’ By High School Teammates
The Minneapolis Star Tribune reports on our very own Olivia Maccoux who has shone as a star athlete, despite having endured over 100 brain surgeries to manage her hydrocephalus.

2013 Audited Financial Statement

Statement of Financial Position
At December 31, 2013

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<td>TOTAL NET ASSETS &amp; LIABILITIES</td>
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Statement of Activities
For the year ended December 31, 2013

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<td>TOTAL SUPPORT &amp; REVENUE</td>
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2013 Donor List

Thank you for your support of the hydrocephalus community. Your contributions allow us to move one step closer to fulfilling our mission of eliminating the challenges of hydrocephalus.

$100,000 or more
Theodore W. Battman Family Foundation
Craig and Vicki Brown
Estate of Frank Clinton
Nancy and Hugh Davey
Paul Gross and Lori Polski

$50,000 - $99,999
Clare Rose Foundation
Jan Churchill and Randy Keill
Modotronic Foundation
Modotronic Neurosurgery

$25,000 - $49,999
Alpen Family Foundation, Inc.
John and Liz Devlin
Cliff and Amy Goldman
Richard H. Goldman Memorial Fund
Rut Stichle Research Institute

$10,000 - $24,999
Adobe Foundation
Assael
Ahn Family Foundation
Anonymous
Howard Sobol and Sarah Billingsgurt
Bostocks Family Foundation
Assem and Amina Chandra
Herman E. and Estelle Chandra
Aseem and Amita Chandra
Howard Solomon and Aseem and Amita Chandra
Ahn Family Foundation
Adobe Foundation

$1,000 – 2,499
Enterprise Products Partners L.P.
Express Rale, LLC
Eye Surgical and Medical Associates, Inc
Lauran Fumaro
Kathy and Roger Farmer
Frederick Ferman
Donald and Joanne Ferrini
Pam and Grant Finsloyn
Florida Panhandle Foundation
Russell and Emily Fudge
Genesis Automation, Inc.
Steven and Ruth Glazer
Global Gals
Godillard School (TME Preschool)
Adam Goldberg
Goodfellow Foundation
Anny Ooov and Shilpi Chandra
David Gravo
Joyce and Daniel Gregson
Edith Bross
Guardian Energy, LLC
Scott and Sherry Haberman
Hawkins Commercial Appliance Service
Paul Holman
Carolyn Hill
Hilt Construction
Richard Hopple
Robert Hughes
Mary Hatzopoulos
Image Outfitters
Integra LifeScience

$500 – $999
Abreus Foundation
Amigo and Christopher Battman
Brick’s Gap Intermediate School
John R. and Dorothy C. Caples
Andrew Carderich
Michelle and Glenn Engleman
Noris and Mary Finkayson
Susan florina
PIMC Ice Sports
Jolene Grau
Vera and Paul Guern
Lyn Lasham and Stephanie Oltman
Madison Diastom Partners
Majestic Realty Co.
Koan and Raia Mcleod
Memorial Hermann
Meschelle Neuroscience Institute
Motorcycle Superstore
Pat Murphy
Opalask Foundation
Kevin O’Shea
Diana Peterson
Portland Northwest Partners
Tracy Taback
Annie Vonghiai
Scott Voygals
Berne and Nancy Wandler
Garland and Suzanne Wood

Margaret Cooper
corin Family
Coydog Foundation
Coydog Insights
Blake Crouser
James and Pamela Crouthamel
Crown Imports
Troy Daniel
David Daniel
Dark Millennium Tattoo Designs
Carson Donlan
Carlton Deaton
Richard and Marilyn Dodson
DigiCarli, Inc.
Pall Donnelly
Sue Duda
Tom Duvall
Ann Edin
Elevation Raps of the Rockies, Inc.
Richard Ellenberger, MD
Jonathan Elsbach
Jill Elsdon
Shannon Ethoe
Janice Ethridge
Stefania Fair
Krystal Fay
Samantha Finlayson
Mary and Paul Finnegan
Judy and Kenneth Fisch
Michael and Kristin Flar
Mary Fleming
Food Services, Inc.
James Garmly
Gavin Garza
QDFS Engineers
Anthony Gaudia
Holly Gilmier, MD
Global Impact
Maez Carol Giddlock
Alyssa Greene
Doug Grisom
Dana Grisom
Lisa Guerin
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Remi Gupta
Gregory Gurnsey
H. Pearce Real Estate
Stephen Haines, MD
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Michael Handler, MD
Dborah Harrison
April Hart
Davin Harwell
Jessica Hais
Hue Arthur Wellsfield
and Helbog
Jonathan and Sarah Hodge

Rumy and Nitin Gupta
Gregory Gurnsey
H. Pearce Real Estate
Stephen Haines, MD
Thomas and Linda Hala
Michael Handler, MD
Dborah Harrison
April Hart
Davin Harwell
Jessica Hais
Hue Arthur Wellsfield
and Helbog
Jonathan and Sarah Hodge

Looking Back…2010
The Fudge Solomon Legacy Society is established to encourage and recognize all those leaving a legacy gift to HA.
In Memory

Jacob Woodward-Adams
Betty Beardsley
Yvette B. Bialko
Nicholas Brinkley
Chin Chang
Leslie Colburn
Peter Coppola
Gerald C. Dorn
Burt Eaton
Jack Galanis
Kate Finlayson
Gerard Fudge
Marvin Israel Goldman
Jean Alice Gorisek
David H. Swanson
Frank Heery
Corley David Heidenreich
Sean Heisal
Harold C. Korstan
Mary Louise Lan Duffy
Ellie Mackiewicz
Meliam McCollom Prouty
Joan Missaki
Midhat Natale
Calib Nelson
Brian Newcomer
Sam Perkins
Meliam Prouty
William and Judy Rogers
Ashley Ruth
Lee Sager
Alex Saldivar
Stephen Sciantrakis
Maurie Shinsky Gross
Lauren Sebowey
Norman Andrew
Stephenson
Rugena E. Stevens, II
Mrs. Elvera Marie Sticker
Jethry Stunkard
Jeanne Thompson
Michael Toski
Gertrude Waik
John Walters
Claesson H. Washburn
Halley Williams

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Hydrocephalus Association

The Legacy of Frank L. Clinton

Frank Clinton was a pillar in the community of Paris, Illinois. He served as the town mayor for 16 years, and there was not a person or family in the area that he didn’t know. Mr. Clinton was introduced to HA shortly after being diagnosed with Normal Pressure Hydrocephalus (NPH). He received treatment and blossomed after the surgery with no complication. In appreciation for HA’s services during this difficult time and eagerness to see the organization’s mission fulfilled, Mr. Clinton placed HA in his will. He continued to engage with the association for many years. On April 4, 2013, Mr. Clinton died from other health challenges. HA received notification that he selected the organization to be the sole beneficiary of his estate. His confidence in HA’s direction and loyalty to see our work advance is evident in his final gift that will ensure his legacy continues.