



Chief Executive Officer's Message

Dear Friends,

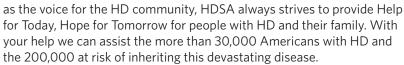
From community services and education to advocacy and research the Huntington's Disease Society of America continues to make major strides in 2013. Through grassroots initiatives on the local level to strategic planning from the national office, HDSA has built a strong foundation for future success.

On April 3rd, Senator Kirsten Gillibrand and HDSA hosted a Research Symposium that commemorated the 20th Anniversary of the discovery of the gene that causes HD with the US Capitol dome serving as the backdrop. The Symposium was the first noteworthy event in the two day advocacy experience that brought advocates from 30 states not only to hear from four of the original 'gene hunters' but also to bring their voices to Capitol Hill to advocate for the Huntington's Disease Parity Act (HR1015/S723).

This past year HDSA formed the Scientific Advisory Board (SAB) provides scientific review of research proposals to ensure that the research programs at HDSA are scientifically sounds, pertinent and provide a high impact to the HD Research community. We are confident that this is a bold first step to improve HDSA's research programs.

And last, but certainly not least, I would like to send a sincere thank you to all the dedicated and passionate volunteers that make up HDSA. From coast to coast, HDSA volunteers are on the front lines fighting to raise funds and awareness for our inspiring community. Our Team Hope Walk program did an incredible job this year and we know it will only be bigger and better next year!

As we look back on the success from this past year, please remember that



Thank you for all your support!

our Vetr

Sincerely,

Louise A. Vetter Chief Executive Officer

Year In Review

TEAM HOPE WALKS

2013 was an amazing year for the Team Hope Walk program. Tens of thousands of people around the country walked side by side in over 102 communities to support the HDSA mission.

Thank you to our Chapters, Affiliates, volunteers, participants, teams, donors, and sponsors for your commitment, dedication, and support. With your continued partnership, the Team Hope family has accomplished milestones in 2013, including:

- Over \$1.45 million dollars raised for the mission and programs of HDSA!
- Over 15,000 participants walked with us at Team Hope Walks across the country.
- 102 Team Hope Walks were held in 2013, of which 18 were new Team Hope Walks!
- A renovated Team Hope Walk website rolled out in August with updated resources, materials, functionalities, and trainings to help the HDSA community plan new Team Hope Walks and grow our current walk with more fundraising and awareness.

- Six Team Hope Walk training webinars have taken place in 2013, all of which are available on the Extranet and Team Hope website. The webinars covered topics such as Participant Fundraising, Sponsorships & Grants, Timelines for Success, Wrap Around Events, and Year After Year Success.
- Seven editions of the Team Hope newsletter "Footprints" have been shared in 2013, highlighting each month's Team Hope walks and offering fundraising tips and advice, reaching over 25,000 people across the country!

All of the Team Hope accomplishments could not have been possible without you, our committed, dedicated and selfless volunteers, participants and donors. HDSA thanks you and invites you back again for an even better 2014!

CONVENTION

On June 21-23, the 28th Annual HDSA Convention was held in sunny Jacksonville, Florida at the beautiful Hyatt Regency Jacksonville. With more than 730 members of the HD community in attendance, Keynote Speaker Marc Holder, PhD (University of British Columbia) gave a poignant speech on The Science of Happiness. Dr. Michael Hayden, MD, PhD. (University of British Columbia) served as the Research Forum Key Note Speaker and discussed strides in HD research.



New this year was the Chapter Awards Luncheon where the most outstanding volunteers from across the country were acknowledged.

The Convention featured world-class workshops about HD care, research, education and advocacy.





RESEARCH

HDSA was proud to announce that in March 2013 a new HD research program was launched to provide financial support for investigator-initiated research directed to better understand the biology of HD as it occurs in humans. HDSA expects that the research will directly impact the lives of persons with HD or help guide the future drug development for HD.

The Donald A. King Summer Research Fellowships were awarded to Jenny Lin of MIT and Jolene Luther of the University of Iowa. Jenny Lin will be working with Drs. William Yang and Steve Horvath at UCLA on a project entitled "A Systems Biology Approach to Analyze Huntington's Disease". Jolene Luther will be working with Drs. Jane Paulsen and Hans Johnson at the University of Iowa on a project called "MRI T2 Hypointensities as a Biomarker in Prodroma HD".

This year, researchers from Prana Biotechnology and UCSF have published positive data on the small molecule, PBT2, in two different animal models of Huntington's disease. PBT2 is a novel compound that is known to cross the blood-brain barrier, appears safe to humans and has demonstrated neuroprotective effects in various brain disease models.

The results showed that PBT2 may positively impact cognition in Alzheimer's disease, Prana set out to discover if PBT2 will have similar effects in HD patients. PBT2 was tested in a Phase II clinical study called Reach2HD for patients with early to mid-stage HD in Australia and the United States.

ADVOCACY

In March 2013 Adam Kinzinger (R-IL) and Bob Pascrell (D-NJ) introduced the Huntington's Disease Parity Act in the U.S. House of Representatives as HR 1015. Thus far, the bill has garnered co-sponsorship from nearly 30 House legislators. In April 2013, Senator Kristen Gillibrand (D-NY) reintroduced the HD Parity Act in the Senate as S723.

HDSA Advocacy has been very active on social media. In a two week period, more than 1,000 people shared an HDSA petition to thank the two House Champions who introduced the HD Parity Act of 2013.

HDSA also produces excellent advocacy resources such as, the Advocacy Toolkit, Disability Application Toolkit and Genetic Information Nondiscrimination Act (GINA) Toolkit.



Rep. Adam Kinzinger (R-IL)



Rep. Bob Pascrell (D-NJ)



Sen. Kristen Gillibrand (D-NY)

Statement of Financial Position

AS OF DECEMBER 31, 2013 AND 2012

	2013	2012
SSETS		
Cash and cash equivalents	3,039,561	2,546,676
Investments, at fair value	563,280	580,191
Contributiions receivable, net	324,380	562,951
Pledges receivable, net	374,628	0
Prepaid expenses and deposits	120,502	86,329
Cash surrender value of life insurance	129,271	124,821
Property and equipment, net	49,728	77,867
Total Assets	4,601,350	3,978,835
ABILITIES		
Accounts payable and accrued expenses	180,566	421,172
Accrued compensation	175,379	187,951
Research grants payable	1,547,370	798,848
Deferred rent	53,270	82,881
Obligation under capital lease	15,942	21,267
Deferred revenue	10,945	7,323
Total Liabilities	1,983,472	1,519,442
ET ASSETS		
Unrestricted		
Undesignated	(1,066,803)	(761,203)
Board-designated	510,152	510,196
	(556,651)	(251,007)
Temporarily restricted	2,924,529	2,460,400
Permanently restricted	250,000	250,000
Total Net Assets	2,617,878	2,459,393

Consolidated Statement of Activities

FOR THE YEARS ENDED DECEMBER 31, 2013 AND 2012

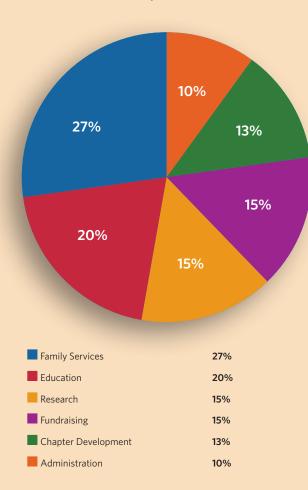
SUPPORT & REVENUE	2013	2012
Public donations	2,653,792	1,967,876
Foundation grants and corporate contributions	1,472,230	1,363,462
Federal campaign	286,049	298,203
Investment income	12,022	5,311
Donated services	187,265	340,735
Other	24,485	21,983
Total Support and Revenue	4,635,843	3,997,570
EXPENSES		
Program services:		
Research	1,207,444	570,986
Community services	2,145,314	1,262,045
Education	1,532,395	1,544,895
Chapter development	1,056,693	1,071,126
	5,941,846	4,449,052
Supporting services:		
Management and general	817,947	832,279
Fundraising	1,120,500	1,524,435
	1,938,447	2,356,714
Total Expenses	7,880,293	6,805,766
Change in net assets before non-operating income	(3,244,450)	(2,808,196)
Non-operating income:		
Special events:		
Gross receipts	4,108,409	4,044,123
Less: direct costs	(705,474)	(618,928)
Net income from special events	3,402,935	3,425,195
Change in Net Assets	158,485	616,999
Net assets - beginning of year	2,459,393	1,842,394
Net Assets - December 31	2,617,878	2,459,393

2013

2012

HDSA DOLLARS AT WORK

Flscal Year 2013 Functional Expenses



75 / 25: Mission / Administration Ratio

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To improve the lives of people with Huntington's disease and their families.

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