



report from the chair and ceo

As we entered our 35th year, the Huntington Society of Canada (HSC) consolidated the steps we had taken forward in recent years as the Huntington movement in Canada and internationally climbed relentlessly toward the summit.

Thirty-five years ago at their kitchen table, Ralph and Ariel Walker began the work to found the Huntington Society of Canada providing generations of families living with the disease, a place to go for mutual support. Society members strive today to carry on in their footsteps.

In 2008-2009 the Society was able to achieve a goal set by the Board of Directors – equal support of services and research. We were able to serve the needs of both families and researchers in creating a world free from HD.

Thanks to the Devlin Fund for Families, HSC continued to strengthen services to families by:

- Completing the plan to add more hours of Individual and Family Services to underserved areas of the country
- Continuing to hold regional HD Symposiums and Family Days, and education workshops in 18 centres across the country
- Launching a series of new e-bulletins, namely “Strength and Knowledge” (caring for people living with HD),

“Discovery Pipeline” (the latest in HD research) and “Volunteers in Touch” (for Chapter executives).

HSC also continued to shore up the digital services it provides to stakeholders by:

- Taking the first steps to circulate the Horizon newsletter electronically
- Creating an online ordering system for the Amaryllis program
- Becoming more responsive to donors by adding electronic tax receipts and event registration capability online
- Creating innovative, online charitable giving opportunities like the new “One for Me, One for HD” e-campaign
- Continuing to enhance the HSC website, notably the resources available online for Chapters.

In spite of difficult financial times, HSC’s loyal, generous donors and event participants enabled the Society to exceed its budget by raising \$3.6 million. This success enabled the Society to continue to increase its investment in innovative

research. 2008/2009 marked the second year of the Society's new research program, which attracted 13 high quality proposals in basic and clinical research. For example, priorities were established to treat the most important symptoms of HD and the competition to spawn new symptomatic therapies attracted 16 proposals from clinics around the world.

Of note, the Society entered into a partnership with the Krembil Foundation, a well-known family foundation with a special interest in neurological disorders. The partnership will help to fund research at McMaster University into the mechanisms of HD and the development of an HD biomarker.

Throughout the year, your Board of Directors worked to strengthen its governance practices by installing methods to survey and touch base with clients and staff, and to evaluate the

Board, individual members and the CEO, as well as develop new policies to guide investments, conflicts of interest, and more.

During 2008/2009, the Society worked with the International Huntington Association (IHA) and Dr. Michael Hayden and Dr. Blair Leavitt, Conference Co-Chairs, to stage the World Congress on HD in Vancouver in September 2009. We look forward to welcoming the world to Canada to celebrate the substantial accomplishments of the Huntington movement.

As the Chairman of the Society and your CEO, together we thank the volunteers, Directors and staff of the Huntington Society of Canada for your untiring efforts to make life better for families living with Huntington disease, and we look forward to continuing on our journey together for another 35 years.



A handwritten signature in black ink, appearing to read 'J Stainsby'.

John Stainsby
Chair, Board of Directors



A handwritten signature in black ink, appearing to read 'DLamont'.

Don Lamont
CEO and Executive Director

Board of Directors 2008 - 2009

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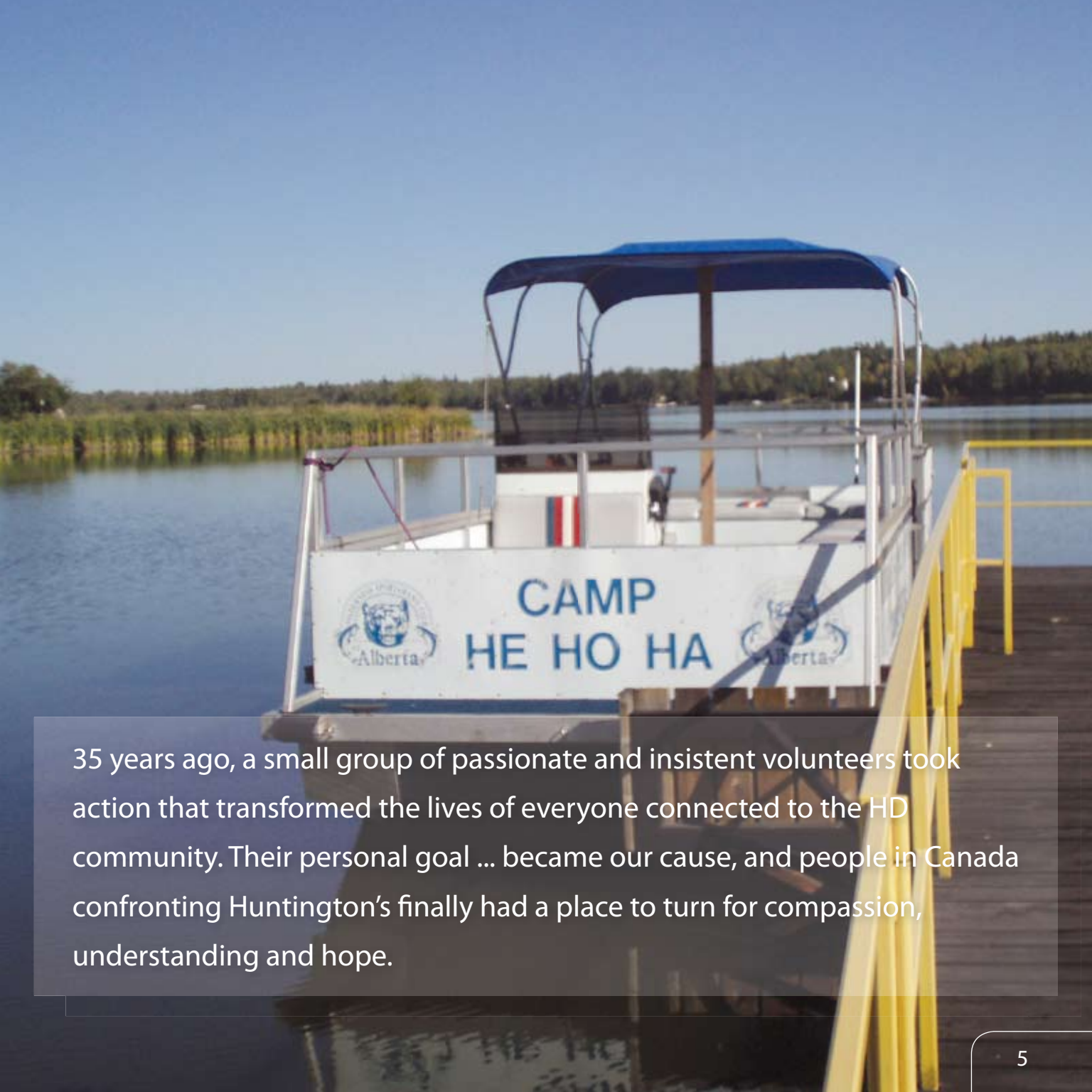
Elaine Taylor
Director

services

In a comprehensive, sustainable means, the Individual and Family Services Program (IFS) continued its relentless pursuit of supporting individuals and families of the HD community across Canada. Just as it was 35 years ago when the Society began professionals in the IFS Program traveled hand-in-hand with individuals and communities to strengthen support for members of the HD community. In the fiscal year 2008/2009, IFS served over 3,000 families in countless communities coast to coast.

The goals of the program are to:

1. Enable individuals and families to adjust emotionally and practically by supplying information and assisting them to manage the present and prepare for the future.
2. Help people identify and understand the family's need for specific services, enable them to gain access to these services on a timely basis and then to follow up to ensure individual's and family needs are met.
3. Create mutual support systems for individuals and families to lessen isolation and to learn from personal experience.
4. Ensure HD families receive a continuum of services by advocating for the development of new services designed to meet the needs of individuals and families. While ensuring existing services are tailored to include families living with HD and to serve them well.
5. Support and educate service providers to better serve individual and families living with HD.



35 years ago, a small group of passionate and insistent volunteers took action that transformed the lives of everyone connected to the HD community. Their personal goal ... became our cause, and people in Canada confronting Huntington's finally had a place to turn for compassion, understanding and hope.



Thanks to continued support from donors, volunteers and professionals across the country, HSC will continue to connect and support the HD community.

IFS worked to achieve these goals in 2008/2009 through two initiatives: Direct Services and Service Development.

Direct Service: The IFS team members worked with individuals and families and assessed the need, provided education about HD, identified community resources, and facilitated referrals. The aim of direct services was and continues to develop a specific strategic plan with individuals and families; to provide guidance and support through problem solving. The provision of direct services can include direct one-on-one, short-term counseling, and support groups.

Service Development: Members of the IFS team worked in communities across Canada and actively sought opportunities to link key players to improve medical, social and other community services. This was achieved by identifying gaps in service, providing education, and facilitating linkages.

Supports to Families

HSC is continuing to develop and advocate on behalf of families for multi-disciplinary clinics across Canada, along with developing more Huntington Study Group sites. We've been successful in helping to increase the number of HD study group sites in Canada with two new locations opening in Halifax, Nova Scotia and Winnipeg, Manitoba in 2008.

Another exciting new development revolves around our involvement with the Ontario Cancer Biomarker Network (OCBN) and the FuRST-pHD program in Ontario. Intended for presymptomatic carriers of the HD gene, the project aims to develop a method of measuring very early HD symptoms that will ultimately help to bring new drugs to market.

Genetic Discrimination

HSC has led the charge on the Canadian Coalition for Genetic Fairness (CCGF) and has been successful in getting the



coalition's mandate positioned in the Conservative platform. On behalf of the coalition, HSC representatives have also had opportunities to meet with various government officials in several ministries, including Justice, Industry, the Office of the Privacy Commissioner and the Prime Minister's office.

Coalition development is well underway. The first meeting of 14 organizations supporting the coalition was successfully held in fall 2008, and many plans and events are in the works for the remainder in 2009, including an official launch on the Hill on October 6th, 2009.

Neurological Health Charities Canada

HSC is a founding member of Neurological Health Charities Canada. The organization received \$15 million from the Canadian federal government for an epidemiological study on neurological diseases in Canada. This research will help to identify the occurrence of neurological diseases, such as HD in Canada, as well as the burden of living with these diseases. This will help to create a case for further advocacy.

We're also working on the development of the Ontario Neurological Strategy to support people living with

neurological diseases. The next step is a meeting with NHCC and several stakeholding ministries. We have partnered with the Ontario Ministry of Health to take important precursive steps to develop this strategy.

Supporting two MSW students

By providing support to two Masters of Social Work (MSW) students, we were able to bring the "Living Life to its Fullest" project to completion. The project resulted in the development of a brochure offering an exchange of information, including tips and suggestions, from families affected by HD.

35 years ago, a small group of passionate and insistent volunteers took action that transformed the lives of everyone connected to the HD community. Their personal goal – to connect and unite HD families in the shared struggle with a cruel and misunderstood disease – became our cause, and people in Canada confronting Huntington's finally had a place to turn for compassion, understanding and hope. Thanks to continued support from donors, volunteers, and professionals across the country HSC will continue to connect and support the HD community.

research

In fiscal 2008-2009, we marked the second successful year of the Huntington Society of Canada's new research funding strategy. The Society maintains a portfolio of strategic projects designed to yield unique and meaningful results in both the near term and the long run.

This strategy is designed to complement the work done by other members of the international coalition of scientists and organizations fighting Huntington disease (HD). To attract top flight proposals, the Society increased the size of the awards for its NAVIGATOR program for basic research and the Symptomatic Relief competition, which is administered by the Huntington Study Group.

NAVIGATOR Program

In total, 11 new applications were received for this research program, which has done much over the years to increase our basic understanding of the mechanisms of HD.

In 2006, Dr. Michael Hayden's lab found that mice resistant to caspase cleavage-6 do not develop Huntington disease. This year the Society funded Dr. Michael Hayden at the University of British Columbia to examine the role of proteolysis and palmitoylation and their interactions with caspase cleavage for the purpose of developing new treatments.

Dr. Eileen Denovan-Wright of Dalhousie University received a grant for her work on regulating transcription, an important process in HD.

Dr. Franchesca Cicchetti at Laval University received funding to continue her work to better understand how the environment of the brain affects the success of neural transplants.

During fiscal 2009, research continued on NAVIGATOR projects undertaken earlier by Dr. Michael Hayden, in regard to the role of phosphorylation on serine 421 in Huntington disease. Research by Dr. Simonetta Sipione at the University of Alberta also continued to further our understanding of the role of gangliosides in HD.

Research by Dr. Ronald Wetzel at the University of Pittsburgh also continued in regard to the functions of the Huntingtin N-terminal Domain, a small segment of amino acids that contributes to the aggregation of huntingtin fragments and perhaps the disease process.

NEW PATHWAY Program

This program is designed to further work on the next generation of targets for comprehensive HD therapies. Dr. Vanessa Wheeler at Massachusetts General Hospital continued work on her project to identify the factors that contribute to instability in CAG repeats in mice in order to identify potential therapies.

There is mounting evidence that mitochondria, the power plant of human cells, play an important role in Huntington disease. Blair Leavitt at the University of British Columbia's Centre for Molecular Medicine and Therapeutics received funding to examine alterations in mitochondria in the brain cells of mice with HD, and to understand the relationship to their age.



The Huntington Society of Canada is continuing the legacy began by Ralph Walker of investing in the most promising HD research to help find a treatment and cure for HD tomorrow, while helping families today, and putting Canada as a leader in the worldwide HD research movement.

SYMPTOMATIC RELIEF Program

The goal of this project is to test experimental interventions that show promise of providing demonstrable symptomatic benefit in the near term for the cognitive (intellectual), motor, behavioural or weight maintenance problems facing patients with manifest HD. With the help of an international panel of scientists, the Society assigned the highest priority to research proposals that addressed the following symptoms: (listed in rank order): irritability and aggression, depression, obsessive compulsive symptoms/perseveration, apathy, sleep disorder, bradykinesia/rigidity and dysphagia. Sixteen proposals were received from applicants in North America and Europe.

In partnership with the Huntington Study Group a new project at Emory University in Atlanta received support. This project will look into the safety and effectiveness of Aripiprazole, a drug already available, in treating irritability and aggression in Huntington disease.

Throughout 2009, work continued on the following projects, which were funded through the Laura's Hope Fund: Dr. Kloos and Dr. Kegelmeyer at Ohio State University on the efficacy of home based biofeedback exercises to reduce falls in HD and Dr. Van Dunjin and Dr. Priller, Leiden University Medical Centre (Netherlands) and Charite-Universitätsmedizin Berlin (Germany) on the use of Bupropion (an antidepressant) for apathy in HD.

Krembil Foundation

In fiscal 2009, the Society formed a partnership with the Krembil Foundation to support promising work by Dr. Ray Truant at McMaster University in Hamilton, Ontario. This project involves a series of related steps pertaining to oxidative stress in HD, analyzing how huntingtin localizes to coflin rods and investigating the role of Transglutaminases (TG2) while determining the chemical modulators of coflin rod formation.

The project also involves the identification of compounds, including kinase inhibitors that may prevent coflin from interacting with huntingtin, along with the development

of a biomarker to measure the progression of HD for use in clinical trials when seeking regulatory approval. This project also has the potential to identify important links between Alzheimer's and Huntington's disease.

New HD Scientists

Working with the Canadian Institute for Health Research (CIHR), which organizes the competition, the Society attracted Canadian scientists to HD research by offering pre and postdoctoral awards to promising young investigators doing HD research.

A new doctoral award was granted to Lise Nicole Munsie at McMaster University, for work related to the project outlined above, to advance our understanding of the role the huntingtin protein plays in the nuclear stress response toward developing a biomarker for HD.

The Society continued to fund two post doctoral projects underway at the University of British Columbia. The first project by Dr. Dagmar Ehrnhoefer, concerns chemical compounds inhibiting caspase cleavage; while the second project, conducted by Dr. Austin Millnerwood, deals with polyQ length and caspase cleavage in mice.

Finally, Dr. Michael Hayden received a prestigious award for research excellence from the Society as the top scoring scientist working on Huntington disease in open CIHR competitions for research regarding intermediate alleles (IA). Individuals with an IA will never develop HD but there remains a risk that their children or grandchildren could develop the disease.

The Huntington Society of Canada is continuing the legacy began by Ralph Walker of investing in the most promising HD research to help find treatments for HD tomorrow, while helping families today, and placing Canada as a leader in the worldwide HD research movement.

Special thanks to Dr. Scott Zeitlin of the University of Virginia for preparing supplementary review reports for the Research Council.

Research Funded During 2008/2009

NAVIGATOR Awards for Basic Research

- Dr. Francesca Cicchetti, Laval University, *Investigation of the Effects of Pathological Environment on Graft Survival in Huntington's Disease*
- Dr. Eileen Denovan-Wright, Dalhousie University, *Regulation of transcription in Huntington's Disease*
- Dr. Michael Hayden, University of British Columbia, *The Role of HIP14-mediated Palmitoylation and Proteolysis in the Pathogenesis of Huntington's Disease*
The role of Huntingtin Phosphorylation on serine 421 (pS421) in the Pathogenesis of HD
- Dr. Simonetta Sipione, University of Alberta, *Exploring the Role of Gangliosides*

In partnership with CIHR

Huntington Society of Canada Prize for Research Excellence (prize for top scoring HD researcher in the CIHR open competition)

Dr. Michael Hayden, University of British Columbia, *Intermediate Alleles*

CIHR Pre-Doctoral Award

Lisa Nicole Munsie, McMaster University, *The HD protein has a role in the nuclear stress response: Mechanism for HD Biomarker*

Galareh Mazarei, *Novel striatal-specific gene expression in a mouse*

CIHR Post-Doctoral Award

Dr. Austin Millernwood, University of British Columbia, *PolyQ length and caspase cleavage on corticostriatal synaptic function in mice*

Dr. Dagmar Ehrnhoefer, University of British Columbia, *Chemical compounds inhibiting caspase cleavage*

In partnership with the Huntington Study Group

Dr. Dylan Wint, Emory University, *"Safety and Effectiveness of Aripiprazole in treating irritability and aggression in Huntington disease"*

Krembil Foundation

Dr. Ray Truant, McMaster University, *Cofilin Rods*

New Pathways

Dr. Blair Leavitt, *Mitochondrial Alterations in the Mouse HD Model*

Research Council 2008-2009

Dr. Ray Truant (Chair)

McMaster University

Dr. Michael Hayden

University of British Columbia

Dr. Patrik Brundin

Wallenberg Neuroscience Centre

Dr. Harold Roberston

Dalhousie University

Dr. Marcy MacDonald

Massachusetts General Hospital
Harvard Medical School

Dr. Eileen Denovan-Wright

Dalhousie University

Dr. Lynn Raymond

University of British Columbia

Dr. Stephen Ferguson

University of Western Ontario

Dr. Blair Leavitt

The Centre for Molecular
Medicine and Therapeutics

report from the treasurer

The results of the 2009 fiscal year are a testament to the commitment and perseverance of the Huntington Society's staff, volunteers, families and donors. Despite the massive down turn of the economy the Society's revenue grew this year by 4%.

Once again, our Chapters and volunteers did an outstanding job organizing a number of successful fundraising events. These events generated more than \$960,000, which represents 26% of the total revenues.

We deeply appreciate the generosity and support of an anonymous donor, whose "matching gift incentive" along with all of our loyal supporters enabled us to accomplish our highest level of success ever.

In line with strategies developed in prior years, the Society continued to increase its expenditures on services, this year by 11%, to families living with Huntington disease and on research, by 55%. Our investments in research have grown significantly in the last few years, to the point where direct expenditures on research and service are now approximately equal. Thoughtful planning has allowed both programs to achieve a sustainable level. Important research projects now have multi-year funding commitments from The Ralph Walker Research Fund and allocations from the General Fund.

Along with almost every investor, the Society experienced market value losses during the year in its investment portfolio. Thanks to wise counsel of the Investments Committee, the losses were relatively small at 2% of book value, while the Toronto Stock Exchange dropped 40% over the same period.

Overall, the financial statement reflects the Society's continued prudent stewardship practices, careful planning and strategic implementation. The Society continues to maintain responsible administrative costs, with the maximum allocation of donated funds to the Society's programs.

Heartfelt thanks to our Huntington Society of Canada volunteers, staff and services team across the country for another exceptional year of service to families living with Huntington disease.



Karen Koester
Treasurer

Statement of Financial Position

At June 30, 2009

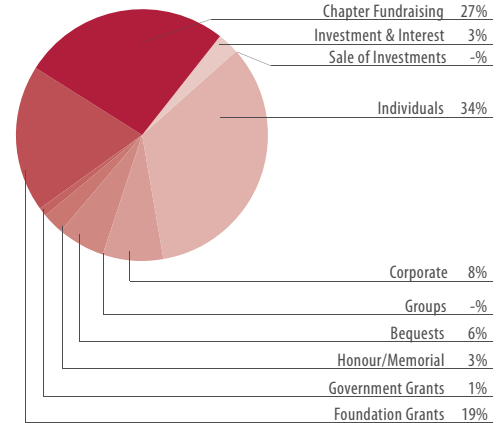
	2009	2008
Assets		
Current Assets:		
Cash	\$ 234,379	\$ 492,395
Investments (at fair value)(note 6)	3,291,712	3,408,733
Accounts receivable	131,459	102,361
Pledge receivable	-	76,813
Accrued interest receivable	18	903
Prepaid expenses	79,802	63,475
	\$ 3,737,370	\$ 4,144,680
Equipment	46,958	60,226
	\$ 3,784,328	\$ 4,204,906

Liabilities and Fund Balances

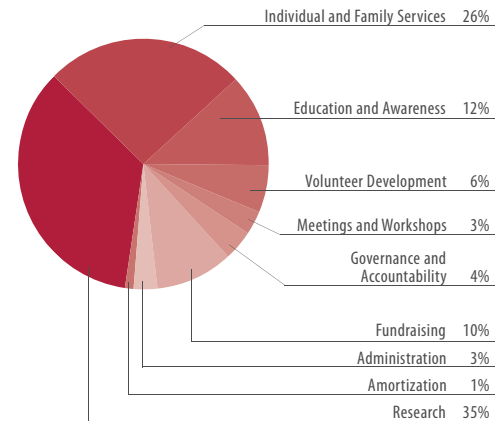
Liabilities:		
Accounts payable and accrued liabilities	\$ 371,574	\$ 130,542
Deferred contributions	371,685	974,277
	\$ 743,259	\$ 1,104,819

Net assets:		
General Fund (note 1)	\$ 705,885	\$ 750,883
Invested in Capital Assets	46,958	60,226
Restricted Endowment Fund (note 2)	1,429,866	1,450,215
Ralph Walker Research Fund (note 3)	650,116	651,175
Laura's Hope Fund (note 4)	208,244	187,588
	\$ 3,041,069	\$ 3,100,087
	\$ 3,784,328	\$ 4,204,906

Revenue



Expenditures



Statement of Operations and Fund Balances

For year ended June 30, 2009

	General Fund	Capital Fund	Endowment Fund	Ralph Walker Research Fund	Laura's Hope Fund	Total 2009	Total 2008
Revenue							
Donations:							
Individuals	\$ 1,003,366	\$ -	\$ 10,000	\$ 201,031	\$ 6,060	\$ 1,220,457	\$ 1,122,542
Corporate	190,958	-	-	67,650	12,247	270,855	316,588
Groups	10,703	-	-	-	-	10,703	6,398
Bequests	223,439	-	-	-	-	223,439	143,530
Honour/Memorial	75,945	-	-	21,348	-	97,293	145,883
Government grants	42,569	-	-	-	-	42,569	28,183
Foundation grants	454,056	-	-	219,500	-	673,556	350,129
Chapter fundraising projects	961,695	-	-	-	-	961,695	1,153,658
Investment and interest income	117,581	-	-	13,906	2,349	133,836	185,658
Gain (loss) on sale of investments	(24,690)	-	-	(9,805)	-	(34,495)	(2,662)
Gain (loss) on sale of equipment	153	-	-	-	-	153	450
	\$ 3,055,775	\$ -	\$ 10,000	\$ 513,630	\$ 20,656	\$ 3,600,061	\$ 3,450,357
Expenditures:							
Research (note 5)	747,000	-	-	513,123	-	1,260,123	814,086
Individual and family services	934,544	-	-	-	-	934,544	849,242
Education and public awareness	433,093	-	-	-	-	433,093	400,716
Volunteer development	200,043	-	-	-	-	200,043	176,963
Meetings and workshops	114,022	-	-	-	-	114,022	90,927
Governance and accountability	160,176	-	-	-	-	160,176	156,619
Fundraising	351,601	-	-	-	-	351,601	347,819
Administration	112,520	-	-	-	-	112,520	72,066
Amortization	-	36,479	-	-	-	36,479	28,142
	\$ 3,052,999	\$ 36,479	\$ -	\$ 513,123	\$ -	\$ 3,602,601	\$ 2,936,580
Excess of revenue over expenditures (expenditures over revenue)	2,776	(36,479)	10,000	507	20,656	(2,540)	513,777
Fund balances, beginning of year	750,883	60,226	1,450,215	651,175	187,588	3,100,087	2,706,327
Unrealized loss on investments classified as available for sale (note 6)	(24,563)	-	(30,349)	(1,566)	-	(56,478)	(120,017)
Interfund transfers	(23,211)	23,211	-	-	-	-	-
Fund balances, end of year	\$ 705,885	\$ 46,958	\$ 1,429,866	\$ 650,116	\$ 208,244	\$ 3,041,069	\$ 3,100,087



We deeply appreciate the support of the anonymous donor whose matching gift incentive ... enabled us to accomplish our highest level of success ever.

Notes to the Statements

1. The General Fund is an unrestricted fund used at the direction of the Board of Directors to fulfill the mission of the Society in all areas including services, research, education, and administration. All unrestricted donations are allocated to this fund.
2. The Endowment Fund is restricted and accounts for donations received which the donor has designated for specific endowment purposes. In exceptional circumstances and subject to approval of 2/3 of the Board of Directors of the Huntington Society of Canada, funds may be transferred from the capital of the Endowment Fund to the General Fund to finance special projects or unbudgeted annual deficits. Annual interest earned on the Endowment Fund will be used to fund normal operational expenditures.
3. The Ralph Walker Research Fund is a restricted fund established in 1983 to commemorate the 10th anniversary of the Huntington Society of Canada and to pay tribute to the Society's founder, Ralph Walker. Donations that have been designated by donors for general research are reported in the Ralph Walker Research Fund, along with the related expenses.
4. The Laura's Hope Fund is a restricted research fund established in 2003 in honour of Laura Evans, by her parents. The purpose of the fund is to support pre-clinical and/or clinical research towards developing a treatment or a cure for Huntington disease. Revenues designated by donors to the Laura's Hope Fund have been restricted for pre-clinical and/or clinical research.
5. All research payments are expensed as payments are made.
6. Effective July 2007 the Society adopted the new Canadian Institute of Chartered Accountants Handbook Section 3855 "Financial Instruments – Recognition & Measurement". The Society has classified its financial instruments as follows: Cash: Held for Trading; measured at fair value with changes in fair value recorded in the statement of operations. Accounts Receivable & Accrued Interest Receivable: Loans & Receivables, measured at amortized value using the effective interest rate. Investments: Available for Sale; measured at fair value with changes in fair value recorded in changes in fund balances. Accounts Payable & Accrued Liabilities and Deferred Contributions: Other Liabilities, measured at amortized value using the effective interest rate. A complete copy of the audited financial statements is available at www.huntingtonsofcanada.ca

donor recognition

Every effort has been made to ensure this is an accurate list of donors who made charitable contributions to the Huntington Society of Canada between July 1, 2008 and June 30, 2009. If you notice an error or omission, please accept our apology and let us know so we can update our records.

Devlin Fund for Families

\$500,000 & Above

Jill & Dan Devlin

Cornerstone Donors

\$250,000 - \$499,999

Anonymous

Co-Op Atlantic

Krembil Foundation

Leadership Donors

\$100,000 - \$249,999

Anonymous

The Andy and Beth Burgess

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Mackenzie Financial Corporation

Maxim Power Corporation

Marilyn McDonald

Sun Life Financial Inc.

The Temerty Family Foundation

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Richard & Margaret Rivard

The Sharp Foundation

TELUS

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Ariel Walker

Supporters

\$10,000 - \$24,999

Anonymous

The Bearspaw Ladies Legacy

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Fred & Joan Nabb

Bob & Norma Stevens

TELUS

Venice House Restaurant

Andrew & Tory Wright

Partners

\$5,000 - \$9,999

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Bishop Allen Academy

Norman Bloom

Bluewater Investment Management

John & Anne Brace

Alan Carson

Bob & Brenda Comtois

Dirtt Environmental Solutions

Ralph Dorr

EnCana Cares Foundation

Giffels Design-Build Inc.

Stephen Gould

Alvin & Darlene Law

Dan Owen

Gisele Reklitis

RBC Foundation

Heath & Kate Sterling

Bruce & Elaine Taylor

Mark & Catherine Taylor

Joan Tweedle

Nic & Mieke Wales

Roy & Ruth Weber

Jim Wiswell & Ellen Foster

Susan Wright & John Sankey

Alan Yeung

Friends

\$1,000 - \$4,999

Anonymous (9)

ACE INA Insurance

AIG Commercial Insurance

Company of Canada

Albi Homes

All Charities Campaign

Alan Antoniuk



Our sincerest thanks to the generous and faithful support of our donors, who have brought us that much closer to our summit of success.

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Patricia Babet
Karen Barkley
Barry Gibbs Sales Ltd.
Bassi Construction & Masonry Ltd.
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Scott & Katherine Beattie
Stephen & Jane Beatty
Norma Bergeron
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Anton Emerick

Emery Investments
The Erin Mills Development Corporation
Esterhazy & District Donor's Choice
Evton Capital Partners Inc.
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First Gulf Development Corporation
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Thank you for remembering those who were close to us. Gifts to the Huntington Society of Canada were made for the following people in the last fiscal year.
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part of the climbing team



Halifax HD Clinic

The Halifax HD Clinic opened its doors in December 2007 to provide multidisciplinary care to HD families in Nova Scotia and the Maritimes. The clinic comprises of Barb Horner, HD Social Worker (the Director of the Nova Scotia and PEI HD Resource Centre), Dr. Shannon Johnson, a neuropsychologist, Dr. Kerrie Schoffer and Dr. Roger McKelvey, neurologists, Dr. Allan Cook, a neuropsychiatrist and clinic nurse Donna Bouchard. This dedicated team supports and enhances the quality of life for individuals with HD and their families through education, clinical care, and medical treatments.



Joe & Susan Hunter

Joe and Susan Hunter have been involved with the Huntington Society of Canada since the days of Ralph Walker. They have been involved with the Society in many capacities over the years- monthly donors, annual supporters, Board member, Chapter members, and event committees. Last year Joe and Susan decided it was time to take the next step in their support of HSC and create a lasting legacy and tribute by starting a Family Fund. Their Family Fund is helping to provide HSC with a guarantee of support for crucial research, education and care services- now and for the future.



Maxim Power

Maxim Power is an Independent Power Producer located in Calgary, Alberta, and this past year MAXIM decided to help support an innovative pilot program to reach children and families living with Huntington disease. The Kids for HD program helps children adjust and adapt to living with HD, staying strong throughout the process. This program fit perfectly with MAXIM's community investment philosophy. The company seeks to find charities that are perhaps less advertised and not as well funded but specifically provide benefits to children with challenges – and their families as well – that can't access the resources that they otherwise need. As President & CEO John Bobenic said "It's about the children and it's about the impact that their struggles have on the entire family unit. That's a perfect fit for us and a very near and dear cause for us."

151 Frederick St., Suite 400
Kitchener, ON
N2H 2M2

Telephone: (519) 749-7063
Toll Free: 1-800-998-7398
Fax: (519) 749-8965

info@huntingtonsociety.ca
www.huntingtonsociety.ca

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