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ON THE COVER

Photo courtesy of Aaron McCourtie from Original Trails originaltrails.com

Original Trails is a travel provider for MS Climb. This image features a team of MS Climbers and was taken during their Everest Base Camp climb in October 2010. For more information visit msclimb.ca

OUR MISSION

To be a leader in finding a cure for multiple sclerosis and enabling people affected by MS to enhance their quality of life.

OUR VALUES

- ► We believe in and work toward the full integration of people with multiple sclerosis in all aspects of life.
- We keep those affected by MS at the centre of our work at every level of the organization.
- We treat everyone with respect and dignity, and protect their privacy and rights to make decisions about their lives.
- We provide meaningful benefit to the greatest possible number of people affected by MS.
- We give hope to people affected by multiple sclerosis.

- ► We acknowledge the grassroots of our organization as one of its greatest strengths.
- We recognize that volunteers, donors, and employees are critical to our success, and we celebrate their commitment.
- We demonstrate ethical behaviour and integrity.
- ► We adhere to the highest standards of quality, transparency, and accountability.
- ► We create safe and collaborative environments that foster positive relationships.



Extraordinary levels of hope, optimism and awareness characterize how the multiple sclerosis community felt in 2010. Chronic cerebrospinal venous insufficiency (CCSVI) dominated online MS discussion forums and became a familiar topic of conversation even among those not personally touched by MS. During this unprecedented year, those impacted by MS and the general public mobilized around CCSVI. These conversations raised the profile of the MS community, placing a spotlight on Canadians who live with MS and bringing attention to new avenues of MS research and treatment.

At the MS Society we continue to work diligently to reflect the needs of all Canadians living with MS. As part of these efforts, we are urgently pursuing more conclusive answers around CCSVI and MS. In 2010, we contributed both dollars and leadership to accelerating the CCSVI discovery process in a step-wise fashion, which will benefit those living with MS today and those not yet diagnosed. Within this National Impact Report, we invite you to view a special two-page feature, documenting our activities in relation to CCSVI throughout 2010.

The theme for this year's report is reaching new heights, which draws heavily on the activities being done at the community level. Consider the Windsor-Essex chapter's provision of respite services and children's programming.

These and other programs are made possible through community events like the MS Walk and generous benefactors like Doug and Sandra Bergeron, who doubled their commitment to the MS Society with a second \$1 million gift to benefit the Windsor-Essex chapter. Through the dedication and generosity of all our event participants, donors and volunteers, MS Society chapters serve as the driving force behind our ability to improve the lives of those living with MS.

Financially, we are pleased to report that our revenues grew 3.6 per cent as compared to the results of the previous year. This growth reflects the recovery of Canada's economy and the deep commitment to our mission from Canadians in every part of the country.

In closing, we would like to recognize everyone who contributed to our 2010 activities and extend a warm thanks to the MS Society volunteers, donors, event participants, sponsors, board members, and staff – you have truly made a difference. We look now to 2011 with high hopes of continued momentum, attention and excitement in our quest to end MS.

YVES SAVOIE

PRESIDENT AND CHIEF EXECUTIVE OFFICER

LINDA LUMSDEN

NATIONAL CHAIR



Today we know more about multiple sclerosis than we did 10 years ago, but many critical questions remain unanswered. The MS Society of Canada is committed to pursuing those answers through advocacy and research until a cure is found.

ACTION

In 2010, the MS Society continued to advocate for government action that would benefit people affected by MS and invested an additional \$10.25 million in MS research. In addition, significant investments were made to expand the endMS Research and Training Network.

This year the MS Society, in partnership with the National MS Society (USA), designated \$2.4 million to seven studies currently underway that investigate the possible association of chronic cerebrospinal venous insufficiency and MS (see also pages 10-11).



FACT

The endMS Research and Training Network is designed to accelerate discovery in the field of MS in Canada and aims to attract, train and retain MS researchers. Currently, the endMS Research and Training Network works with over 770 researchers from over 50 academic and health institutions across Canada. For more information visit **endMS.ca**



HOW TO ENGAGE

Learn more about MS research progress and the studies we fund msssociety.ca/research

IMPACT

Today researchers are delving deep into several promising avenues of MS-related research. CCSVI, genetics, neuroprotection and repair, environmental factors, and stem cell therapy are just some of the research areas yielding answers to important questions.

On March 23, 2011, the federal government announced the creation of an innovative national monitoring system that will capture information to help identify disease patterns and track long-term treatment outcomes. People living with MS will be the beneficiaries of this monitoring system as it will shed light on important topics like CCSVI, the long-term benefits of current therapies and also serve as an invaluable tool in assisting researchers, health professionals and policy makers in their work.

PHOTO: HEALTH MINISTER LEONA AGLUKKAQ, IN OTTAWA
AT LAUNCH OF THE MS MONITORING SYSTEM

REACHING CANADIAN COMMUNITIES:

NEED

Multiple sclerosis is a chronic and unpredictable illness. It can often introduce daily physical, emotional and financial challenges into a person's life. With 55,000 to 75,000 Canadians living with MS, families across the country depend on the MS Society to provide support that enables them to enhance their quality of life.

ACTION

Through our donors' support, we are able to take action and provide programs that meet the needs of Canadians affected by MS. The impacts of these programs are felt across the country and allow people living with MS to reach new heights.

FACT

The MS Society offers a variety of programs to help people affected by multiple sclerosis effectively manage and cope with the disease. MS can be quite challenging at times, but through the support offered by the MS Society, no one has to face these challenges alone. The following programs are offered in most chapters and units:

- Information and referral
- Supportive counselling
- Support and self-help groups
- Recreation and social programs
- Financial assistance

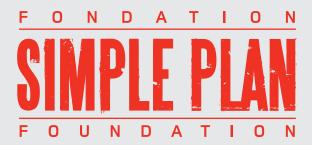
IMPACT

Whether it is through an equipment provision program or a new service created to meet the needs of a specific community, many Canadians touched by MS benefit from the generous support of corporations, foundations and individuals across Canada. Here are a few examples of our donors' dollars in action, positively impacting the lives of people touched by MS. Thanks to the overwhelming generosity of Doug Bergeron, CEO of San Jose, CA-based VeriFone Systems, Inc. and his wife Sandra, people affected by MS in Ontario's Windsor-Essex area have access to much-needed support services. The Bergerons recently made their second million-dollar gift to the MS Society, in continued honour of Doug's late father, George, who had MS and was a past chairman of the Windsor-Essex Chapter. From transportation, respite services and children's programming to the George Bergeron Healing Garden, the Bergerons have empowered the chapter to reach deeply into its community.



The Equipment Provision Program of the British Columbia and Yukon Division provides mobility aids and equipment to help those affected by MS with the physical challenges the disease presents. The MS Society is able to offer this program with support from Mr. & Mrs. P.A. Woodward's Foundation, Provincial Employees Community Services, Face the World Foundation and the Province of British Columbia.

THANKS TO OUR DONORS



A family touched by MS often faces steep medical bills on a limited income. This leaves very little room for social or recreational activities for children with MS or children of a parent with MS. Thanks solely to the generous support of the Cargill Cares Committee, the Saskatchewan Division established their Children's Recreation Subsidy Program, and now more children affected by MS in Saskatchewan are able to participate in recreation programs along with their peers.



Receiving a diagnosis of MS can be a scary moment in a person's life. Quebec Division's Hope for the Future, Coping with the Present program helps people newly diagnosed by providing them with the critical information, support and resources they need to face their new reality. "It was nice to feel supported and realize we are not alone in dealing with this condition," says a program participant. The Simple Plan Foundation acknowledged the importance of initiatives such as this and has made significant gifts totaling \$25,000 since they began supporting the program in 2009.



JC Bradford of BC received a power wheelchair and stair lifter from an MS Society support program. Now, as a direct result of her new mobility equipment, she can remain at home instead of moving into a long-term care facility. "Thanks to the MS Society paying for this piece of equipment, I have a lot more freedom to move around our home...and now I am able to remain involved in my community, where I volunteer and have lots of friends. Thank you for making my life much easier."



People living with MS deserve answers on CCSVI – an important new stream of MS research. The MS Society of Canada is committed to helping the MS community find those answers as quickly as possible to ensure that Canadians living with MS have the best information available when considering CCSVI-related procedures.

ACTION

In 2010, the MS Society of Canada and the National MS Society (USA) jointly funded seven CCSVI and MS studies valued at \$2.4 million. The studies aim to provide important information on the role of CCSVI in MS and the best method to screen for the condition. This information is essential to designing research protocols and sets the stage for a nationwide clinical trial if the evidence from these and other studies around the world show that one is warranted.

The MS Society also pressed governments to take action on the CCSVI file. We called on governments across Canada to commit funds for a pan-Canadian therapeutic trial on CCSVI if and when one is warranted, ensure no one who undergoes treatment for CCSVI outside of Canada is denied necessary post-treatment medical care and create registries of people who have obtained the CCSVI procedure outside the country.

In addition, the MS Society of Canada launched CCSVI.ca to act as a resource for those who want timely, credible and relevant information about CCSVI-related procedures.

IMPACT

Governments across Canada heard the voices of the MS community and took action on CCSVI. For an overview of government actions on CCSVI, please see page 10, CCSVI: Year in review.

Globally, more information, attention and resources are being directed towards MS issues than any other time in our 63-year history. MS is being recognized as a top-priority health issue and with this momentum, we hope to get answers on CCSVI as quickly as possible.



HOW TO ENGAGE

Take action! Help the MS Society press governments for action. mssociety.ca/advocacy



FACT

In May 2010, the MS Society requested the federal government to earmark \$10 million for CCSVI and MS research.

PHOTO: MS SOCIETY, QUEBEC DIVISION VOLUNTEERS AND STAFF AT THE NATIONAL ASSEMBLY IN QUEBEC



Canada has one of the highest incidence rates of multiple sclerosis in the world. Approximately 1,000 new cases of MS are diagnosed in our country each year, which means that three more Canadians are diagnosed with MS every day. The need to raise awareness and funds for all Canadians living with MS has never been greater.

ACTION

For the second consecutive year, A&W Food Services of Canada Inc. partnered with the MS Society and thousands of Canadians across the country to raise awareness and funds for MS. On August 26, the Cruisin' for a Cause fundraiser was relived with the sale of classic car cut-outs, donation boxes, and with \$1 from every Teen Burger® sold donated to the MS Society.



FACT

An astonishing 11,000 people sent their RSVP for Cruisin' for a Cause through Facebook; while on Twitter, the fundraiser was a top-trending topic in Canada on August 26.

IMPACT

Thousands of A&W patrons in over 700 of its locations across Canada made a contribution to the Cruisin' for a Cause fundraiser. Their generosity nearly doubled the funds raised in the event's inaugural year with over \$700,000 going toward progressing MS research and supporting programs that help to improve the quality of life for those living with MS.



HOW TO ENGAGE

Buy a Teen Burger® on August 25, 2011 at our annual A&W Cruisin' for a Cause day.

PHOTO: A&W AND MS SOCIETY STAFF AND AMBASSADORS
AT 2010 CRUISIN' FOR A CAUSE EVENT

CCSVI: YEAR IN REVIEW

November 2009

CTV and the Globe and Mail profile Dr. Paolo Zamboni's CCSVI theory of MS. MS Society of Canada issues immediate call for research proposals.

May 5, 2010

MS Society lobbies the federal government for \$10 million for funding into CCSVI and MS research.

September 13, 2010

Newfoundland & Labrador sets aside \$320,000 to fund observational studies for those who go out of country for CCSVI-related procedures.

July 20, 2010

The CCSVI Working Group is formed with the approval of the national board of directors of the MS Society to provide advice and recommendations to the board on matters concerning CCSVI.

2009

April 7 & 14, 2010

MS Society holds live informational webcasts on **CCSVI** in English and French attended by over 1,000 people.

January - March 2010

Regular national news coverage of new potential treatment theory creates broad awareness of MS and CCSVI. Canadians seek CCSVI treatment abroad.

June 11, 2010

MS Society and National MS Society (USA) jointly commit \$2.4 million to support seven operating grants that explore the relationship of CCSVI and MS.

August 26, 2010

Canadian Institutes of Health Research (CIHR) in collaboration with the MS Society hold meeting of leading North American experts in MS to identify research priorities in MS research.





HOW TO ENGAGE

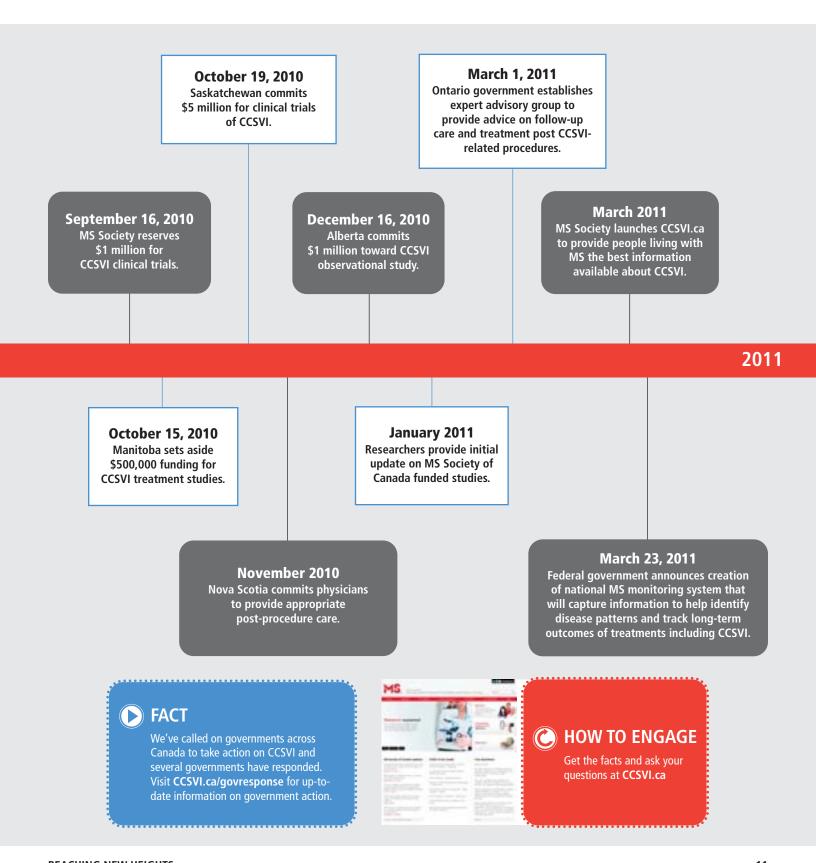
Share your story at CCSVI.ca



! NOTE

At press time governments continued to discuss possible action on CCSVI. For up-to-date developments on this important topic visit CCSVI.ca

Chronic cerebrospinal venous insufficiency or **CCSVI** is a term first used by Dr. Paolo Zamboni that suggests clogged veins can be a cause of MS. He believes that unblocking the veins can have a positive effect on the lives of those diagnosed with MS. In 2010, CCSVI dominated the attention of the MS community. Here is a retrospective look at how the story has unfolded so far.





The MS Society of Canada continues to be a trusted source of information for many Canadians impacted by MS. Education is a priority for the organization and in the past few years there has been an increase in demand for receiving quality information via new and emerging technologies.

ACTION

The National Education Series brings timely, credible, and relevant information about multiple sclerosis to communities throughout Canada. The series covers a wide range of topics that improve overall quality of life for our primary clients: people living with MS. The sessions are typically delivered in a live setting and include a panel presentation made by leading experts as well as a moderated question and answer period between the panel and the audience.

In 2010, with an unrestricted educational grant from EMD Serono, we offered a live audience education session and for the first time it was streamed live via webcast.



FACT

Nearly 87 per cent of participants of the National Education Series sessions reported that they would make at least one new choice about how they managed their MS.

IMPACT

Through locally delivered education events, people with MS, their families and support networks gain a better understanding of multiple sclerosis. These events have been offered in English and French in 74 locations from coast to coast, reaching approximately 16,000 people affected by MS.

Through our new live and on-line delivery method, we are able to reach more people affected by MS than if we had relied on the conventional approach alone. "I would like more education and information sessions provided via webcast. I live 1½ hours from the nearest MS Society office, so the webcast was very convenient for me," stated one webcast participant. Another said, "Thank you for organizing this so that I could attend from the comfort of my home."



HOW TO ENGAGE

Missed this year's education session? You can view the live webcast at mssociety.ca/webcasts



For some people, a diagnosis of multiple sclerosis leads to an inevitable lifestyle change. The financial impact on people living with MS and their families can be significant. Approximately 80 per cent of Canadians with MS are unable to work full time.

ACTION

With the help of our volunteers, the MS Society ensures that information developed as part of the national publications program is up-to-date and relevant for people affected by MS. Jeff McCartney, wealth advisor at Scotia McLeod, is a long-time volunteer of the MS Society. He offers his expertise in finance to contribute to resources about financial planning for people with MS.

As noted by Jeff, "An important aspect about navigating the journey with MS is managing your money and planning wisely for the future."



FACT

The national publications program was created in 2006. Through this program, we have developed 30 publications in English and French to address specific unmet needs of those affected by MS.



HOW TO ENGAGE

For more information on financial planning and living with MS, three videos developed by Globe Investor are available for preview at **mssociety.ca** (visit Living with MS).

This year, through a newly formed partnership between the MS Society and the Investor Education Fund, *Adapting: Financial Planning for a Life with Multiple Sclerosis* was created. It addresses key financial issues for Canadians affected by MS.

IMPACT

"This publication offers important information and guidance for people trying to prepare, plan and stay well-informed for the future," says Jeff. "Being prepared for the unpredictability of living with a chronic illness is essential both personally and when one is thinking of others in their family such as young children and other dependents."

PHOTO: JEFF MCCARTNEY, WEALTH ADVISOR, SCOTIA MCLEOD



With over 400 employees spread across seven provinces, efficiency in our work practices requires diversity and inclusiveness among staff nationwide.

ACTION

Implement a national human resources strategy that aims to increase diversity among the workforce, improve online collaboration and enhance communication across the country. As part of this strategy, a nationwide employee survey is conducted every second year to measure improvements across the country.



FACT

The MS Society of Canada has seven regional divisions and over 120 chapters across the country – all working to find a cure for MS and enable people affected by MS to enhance their quality of life.



HOW TO ENGAGE

Visit **mssociety.ca** and use our interactive map to find an MS Society of Canada office near you.

IMPACT

In 2010, the level of diversity and collaboration across the organization increased from 68 per cent (2008 Nationwide Employee Survey) to 73 per cent (2010 Nationwide Employee Survey). This five per cent increase strongly reflects improvements made to HR processes including recruitment and performance management as well as the creation and launch of an online collaborative platform. We are proud to report on these nationwide survey results that reflect our internal successes, and allow us to better serve people whose lives have been touched by MS.



PRESIDENT'S AWARD

Suzanne Deschamps of Alberta was presented with the National President's Award in recognition of her valuable contributions to the MS Society and to the community-at-large.

A highlight of Suzanne's many accomplishments was her work on getting four MS drug therapies on the Alberta drug formulary, which was approved in June 1998. Since her diagnosis of MS in November 1992, Suzanne has been the president of the Lakeland Chapter and has raised over \$68,000 for the RONA MS Bike Tour.

OPAL AWARD FOR CAREGIVERS

Richard Jurewicz of Ontario was presented with the National Opal Award for Caregivers in recognition of his commitment to taking care of his wife, Teresa, who was diagnosed with relapsing-remitting MS in 1986.

Aside from being a caregiver to Teresa, Richard also participates in the RONA MS Bike Tour in Ottawa and has rasied \$57,000 to date.

PHOTO: NATIONAL OPAL AWARD FOR CAREGIVERS WINNER RICHARD JUREWICZ AND WIFE TERESA

AWARD OF MERIT, NON-MEMBER

Dr. Katherine Knox of Saskatchewan received the National Award of Merit, Non-Member for her outstanding contribution to the mission of the MS Society.

Dr. Knox is involved with the endMS Research and Training Network and the Canadian Collaborative Project on Genetic Susceptibility. She has been a keynote speaker in numerous education sessions related to MS such as the Saskatchewan Division's MS family conference and CCSVI information sessions in Regina, Saskatoon and Lethbridge, Alberta. Dr. Knox is also collaborating with Dr. Anthony Traboulsee on research on CCSVI and MS.

JOHN ALEXANDER AWARDS

2009 RECIPIENTS

Print Award: Hope blossoms in those living with MS during carnation campaign, Gary Kean, *The Western Star*, May 7, 2009. **Broadcast Award:** One Family's Battle, Annie DeMelt, CTV Montreal, April 24, 2009.

2010 RECIPIENTS

Print Award: Depression often unwelcome symptom of multiple sclerosis, *Abigail Cukier*, *Stoney Creek News*, *March* 25, 2010

Broadcast Award: Multiple Choice, Cindy Sherwin, Marc D'Amours and Pascal Sevigny, CTV Montreal, November 24, 2010.

LEADERSHIP DONORS

The MS Society of Canada is proud to celebrate the following individuals, corporations, and foundations that provided generous gifts to the MS Society's Leadership Giving program to support research and client services.

EMD Serono RBC Financial Group, through the **RBC** Foundation

\$100,000 - \$249,999

Doug & Sandra Bergeron Pfizer Canada Inc.

\$50,000 - \$99,999

Anonymous

Alberta Culture and Community Spirit Alberta Seniors and Community Supports Biogen Idec Canada Inc. The Calgary Foundation In honour of Bill & Isabel Kelley

The Law Foundation of BC

Novartis Pharmaceuticals Canada Inc. Alanne O'Gallagher

\$25,000 - \$49,999

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Karen Lauer

The Catherine & Maxwell Meighen Foundation Provincial Employees Community

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Teva Canada Innovation Vancouver Foundation

\$10,000 - \$24,999

Anonymous (3)

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| Gluteus to the Maximus Greatful Tread | Leanne Anderson Martha Corbin | ON MB | TeamBP | Shauna Bygrave | AB | PricewaterhouseCoopers LLP | Jonathan Simmons | ON |
| Greattui iread | | MB | The Brotherhood | Team Paul Lafond | ON | Rusty Links SLOW SPOKES | Shawn Mole | AB |
| | Grace Lackman | | VIA Rail WindRiders | Denis Lemieux | QC | | Jack MacDuff | NB |
| In an an Dandah ann ann | Angela Morelli | MB AB | windkiders | Robert McDonald Colin Wensley | SK SK | Stantec Stingers | Jamie Burke Dave Macmurchie | AB BC |
| Jasper Rockhoppers John Ernewein Limited | Greg Van Tighem John Ernewein | ON | V | | SK BC | Team Addison Team Docktor | | AB |
| | John Ernewein | ON | YouGoGang | Myrna Hastings | | | Pamela Haskell | |
| – Mary's Crew Kilimandjaro-Gaz Metro | Marie Bourbonnais | QC | Y-Ride YMCA | Ike Henry | AB | Team Jannie | Jan Van Velzen Shawn Kupfert | ON QC |
| L'Équipe qui a du chien | Josée Morneau | QC QC | | | | Team Kupfert The Merry Striders | Snawn Kuptert Sandra Corbett | QC ON |
| Lequipe qui a du chien Les Dérailleurs | Mario Sorel | | | | | The MSFits | Soren Schou | BC |
| les Guy-Dons | Guy Auger | QC OC | | | | The Wishing wenches | Carmen Amundsen | BC BC |
| ies duy-polis | Guy Auger | ŲĊ | | | | With The Grain | Ralph Jackson | MB |
| | | | | | | with the Grain | varbii jackzon | IVIB |
| DEACHING NEW HEI | | | | | | | | 24 |

EVELYN OPAL SOCIETY

We'd like to extend our deepest gratitude to the following supporters, who in 2010 graciously chose to create a legacy towards a future free from MS, and in so doing, joined the growing ranks of our Evelyn Opal Society members.

Anonymous Daphne Tucker Faulkner Karen Mychaluk Larus Thorarinson

Gayelene Bonenfant Douglas Homer-Dixon Anne Power

We are humbled by the generosity of our donors and the extraordinary commitment they show towards the effort to end MS. We are honoured to recognize the contributions of the following Evelyn Opal Society members in the past year.

\$250,000 +

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Estate of Audrey May Trainor

Estate of Anne E. Warner

Estate of Peter James Wakeford

We apologize for any errors or omissions.

Statement of Revenue and Expenditures

The following presents the statement of revenue and expenditures for the Multiple Sclerosis Society of Canada for the fiscal year ended December 31, 2010. There are three columns of financial data this year because of the decision approved by the national board of directors to change the year-end of the Multiple Sclerosis Society of Canada from August 31 to December 31 to better match our business cycle and to match the taxation year for the majority of our donors. As a result of this change, a short fiscal year running from September to December, 2009 was necessary as Canada Revenue Agency requires a reporting period that is no longer than 12 months. The four months of operating results are not a normal fiscal year because most fundraising events happen from the spring to the fall but this period was necessary to bridge from our year ending August 31, 2009 to our new fiscal year ending December 31, 2010. The deficit for this short fiscal year was anticipated and there was sufficient cash and net assets within the organization to absorb this shortfall. The most appropriate comparison of our financial performance is between the two 12 month fiscal years.

COMBINED STATEMENT OF REVENUE AND EXPENDITURES

| (in thousands of dollars) | Twelve months ended Dec. 31, 2010 \$ | Four months ended Dec. 31, 2009 \$ | Twelve months ended Aug. 31, 2009 \$ |
|---|--------------------------------------|--|--|
| Revenue | | | |
| Leadership giving activity | | | |
| Bequests | 3,032 | 1,527 | 3,684 |
| endMS Research & Training Network | 2,332 | 1,224 | 1,165 |
| Corporate giving and major donors Grants from governments | 1,585 1,344 | 636 514 | 1,472 |
| Grants from pharmaceutical companies | 407 | 119 | 1,846 516 |
| Other grants | 933 | 320 | 662 |
| | 9,633 | 4,340 | 9,345 |
| Community based fundraising events | 27,180 | 2,044 | 26,056 |
| Individual giving and direct marketing | 17,088 | 7,860 | 16,975 |
| Dinners, tournaments, and third party events | 4,349 | 1,865 | 4,516 |
| Gaming Sale of goods | 1,569 349 | 415 403 | 1,259 416 |
| United Way and HealthPartners | 1,727 | 467 | 1,779 |
| Public awareness activities | 1,304 | 16 | 1,511 |
| Investment income | 1,023 | 423 | 241 |
| Miscellaneous | 306 | 53 | 204 |
| Memberships | 79 | 34 | 85 |
| Total Revenue | 64,607 | 17,920 | 62,387 |
| Fundraising expenditures | | | |
| Leadership giving | 1,925 | 785 | 2,069 |
| Community based fundraising events | 10,934 | 2,778 | 11,270 |
| Individual giving and direct marketing Dinners, tournaments, and third party events | 10,616 1,706 | 5,066 680 | 10,849 2,063 |
| Gaming | 340 | 106 | 433 |
| Cost of goods sold | 236 | 235 | 281 |
| Indirect fundraising | 1,564 | 520 | 1,689 |
| | 27,321 | 10,170 | 28,654 |
| Program and administration expenditures | | | |
| Client services | 9,640 | 3,719 | 10,524 |
| Research | 7,118 | 576 | 7,324 |
| Research – endMS Research & Training Network | 2,442 | 1,387 | 1,165 |
| Research – CCSVI | 699 | 4.507 | - 5.467 |
| Public education and awareness Chapter and volunteer support and development | 6,995 3,762 | 1,587 939 | 5,167 3,013 |
| Government and community relations | 1,690 | 501 | 1,570 |
| MS Clinics | 922 | 286 | 889 |
| Administration | 4,130 | 1,521 | 4,467 |
| | 37,398 | 10,516 | 34,119 |
| Total Expenditures | 64,719 | 20,686 | 62,773 |
| Deficiency of revenue over expenditures for the period | (112) | (2,766) | (386) |



1-800-268-7582 info@mssociety.ca mssociety.ca

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175 Bloor Street East Suite 700, North Tower Toronto, ON M4W 3R8 416-922-6065

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1501-4330 Kingsway Burnaby, BC V5H 4G7 604-689-3144

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9405 - 50 Street, Suite 150 Edmonton, AB T6B 2T4 780-463-1190

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