Dear Friends,

The National Multiple Sclerosis Society strives to improve the lives of people living with MS and their families. We know that MS disrupts connections - inside the body, as well as in our lives. The connections made between those living with MS and their families, volunteers, researchers, fundraisers, and Society staff, help to reconnect us. In addition to improving lives, the Society strives to stop disease progression, restore lost function and end MS forever through the NOW (No Opportunity Wasted) research campaign. In 2012, the connections we made and the money we raised brought us one step closer to our goals and we could not have done it without your help.

In 2012 the National MS Society connected with investigators and invested $44 million in 350 research projects. Your commitment to the Society propelled research in promising directions. Research highlights include approval of oral Aubagio®, the launch of the International Progressive MS Collaborative to speed research to stop progressive forms of MS, and advances in uncovering MS triggering factors. We connected with researchers in our local community and with your support, we funded nearly $12 million in research projects at Northern California institutions. This is truly an exciting time in research.

At the Northern California Chapter, we made powerful connections with our clients, volunteers, and MS community. We are pleased to share that the chapter saw a 20% increase in the amount of programming from 2011. Our 48 Self-Help Groups served over 700 people. Chapter staff assisted 164 members with over $112,000 of direct financial assistance. Our volunteers provided over 5,000 service hours. Local MS activists visited 30 congress people and 72 state level officials. Making connections at the federal, state and local levels, we advocated for the retention of critical services and benefits for people on both Medicare and Medicaid.

The accomplishments of our Chapter could not have been achieved without the exceptional leadership and connections facilitated by our Board of Trustees and the continued commitment of our staff, volunteers and community donors. It is with gratitude and thanks that we are able to enter 2013 well-positioned to continue the pursuit of our ultimate goal – a world free of MS.

Let’s keep connecting and join the movement.

Warm Regards,

Angela Lai
Chair, Board of Trustees
ABOUT THE NATIONAL MULTIPLE SCLEROSIS SOCIETY

The National MS Society addresses the challenges of each person affected by MS. To fulfill this mission, the Society funds cutting-edge research, drives change through advocacy, facilitates professional education, collaborates with MS organizations around the world, and provides programs and services designed to help people with MS and their families move forward with their lives. In 2012 the National MS Society invested $44 million in over 350 new and ongoing projects. These new awards are part of a comprehensive research strategy aimed at stopping MS, restoring function, and ending MS forever. This commitment is the latest in the Society’s relentless research effort. Currently, the Society invests almost $12 million in critical MS research initiatives locally at J. David Gladstone Institutes, UCSF, Stanford, UC Davis and UC Berkeley.

ABOUT THE NORTHERN CALIFORNIA CHAPTER

The Northern California Chapter serves 40 counties, over 12,000 people living with MS, and the 84,000 people who care for and about them. The Northern California Chapter is a lifeline for people living with MS; by providing reliable knowledge, engaging and innovative programs and services, financial assistance and a sense of community, the Northern California Chapter keeps people moving their lives forward.

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WE ARE A DRIVING FORCE OF MS RESEARCH AND TREATMENT TO STOP DISEASE PROGRESSION, RESTORE FUNCTION, AND END MS FOREVER.
MOVENT IN RESEARCH

The National MS Society recognizes that while much progress has been made in research, much more is needed. In response, we launched an organizational research revolution, MS NOW (No Opportunity Wasted), as our comprehensive approach to propel knowledge forward. The National MS Society will devote $250 million to support our MS NOW research program. Our commitment is a broad strategy focused on three distinct outcomes:

- **STOPPING THE DISEASE**
- **RESTORING WHAT’S BEEN LOST**
- **ENDING MS FOREVER**

MS NOW will pursue all promising paths of research by supporting the most encouraging therapies from their start in lab research, all the way through clinical trials. In 2012 the National MS Society invested $44 million in over 350 new and ongoing projects. These new awards are the latest in the Society’s relentless research effort. Currently, the Society invests almost $12 million in critical MS research initiatives locally at J. David Gladstone Institutes, UCSF, Stanford, UC Davis and UC Berkeley.

MS research continued to advance on many fronts in 2012. This year saw:

- The approval of a second oral therapy for relapsing forms of MS and other emerging treatments progressing through the development pipeline
- The launch of the International Progressive MS Collaborative, the largest effort to date to speed research to stop progressive forms of MS
- The discovery of what could be a target of the immune attack in people with MS, which may lead to new understanding of the disease and new treatment strategies
- The completion of the first human trial of an experimental therapy targeting myelin repair
- Progress in restoring functions using innovative rehabilitation techniques, including memory enhancement using a technique involving stories and imagery to solidify learning, and improving balance and mobility with specific exercises
- Advances in uncovering MS triggering factors, bringing us closer to finding ways to prevent the disease; and many other advances pushing us closer to a world free of MS

A woman in Mill Valley is able to make meaningful connections in her personal and professional life because of her disease-modifying treatments.

“Because of my treatments, today I am a healthy person who just happens to have MS. As the years go by in a blink, my MS treatments remain critical to my well-being and longevity, allowing me to continue my active professional and personal life with my family. Maybe, just maybe, the world again will be my oyster.”

- Christina, Mill Valley, CA
LOCALLY FUNDED MS RESEARCH

J. David Gladstone Institutes

- Natacha Le Moan, Ph.D.
  Funding: $143,223 | Summary: Exploring early events that lead to nerve tissue damage in MS.

Stanford University Medical Center

- Ben Barres, M.D., Ph.D.
  Funding: $496,237 | Summary: Determining how the protective barrier between the bloodstream and the brain and spinal cord breaks down in MS.
- Ben Barres, M.D., Ph.D.
  Funding: $487,716 | Summary: Investigating how hormone promotes growth of nerve-ensheathing myelin for clues to increasing myelin formation in MS.
- Alexandra Goodyear, M.D.
  Funding: $130,000 | Summary: Developing the skills involved in the design, implementation, and analysis of clinical trials in MS.
- Michael Kurnellas, Ph.D.
  Funding: $135,948 | Summary: Exploring a naturally occurring substance produced by cells for its therapeutic potential in MS.
- Lawrence Steinman, M.D.
  Funding: $448,898 | Summary: Engineering a substance produced in the brain as a possible approach to a new treatment for MS.
- William Talbot, Ph.D.
  Funding: $446,243 | Summary: Identifying genes involved in the production of myelin to find new ways to repair damaged myelin in MS.

University of California, Davis

- Wenbin Deng, Ph.D.
  Funding: $295,256 | Summary: Investigating the therapeutic potential of using cells derived from adult skin to repair nerve-insulating myelin damaged during the course of MS.
- David Pleasure, M.D.
  Funding: $433,947 | Summary: Exploring the role of specific immune reactions in nerve fiber damage and testing ways to block them to protect the nervous system in MS.

University of California, San Francisco

- Sergio Baranzini, Ph.D.
  Funding: $1,139,921 | Summary: Cataloging patterns of genes found in the blood of people with MS for clues to factors controlling disease susceptibility, severity, progression and response to therapy.
- Sergio Baranzini, Ph.D.
  Funding: $699,836 | Summary: Evaluating the factors that cause MS to find new ways to treat it.
- Jeffrey Bluestone, Ph.D.
  Funding: $498,211 | Summary: Looking for ways to use the immune system’s regulatory mechanisms to stop immune attacks in MS.
- H. Christian von Büdingen, M.D.
  Funding: $485,808 | Summary: Examining the relationship between B cells in the blood and those in the brain to improve MS diagnosis and therapy.
- Jonah Chan, Ph.D.
  Funding: $379,167 | Summary: Looking for ways to encourage the growth of new myelin-forming cells to repair damage in MS.
- Jennifer Graves, M.D., Ph.D.
  Funding: $130,000 | Summary: Developing the skills involved in the design, implementation, and analysis of clinical trials in MS.

University of California, Berkeley

- Farren Briggs, Ph.D.
  Funding: $135,448 | Summary: Exploring how exposure to specific factors may interact with a person’s genes to trigger MS.
• **Ari Green, M.D.**  
  **Funding:** $771,773  |  **Summary:** Developing a technique to measure the health and injury of nerve cells as a potential tool for quickly evaluating the potential of therapies to protect the nervous system.

• **Angela Hahn, Ph.D.**  
  **Funding:** $150,800  |  **Summary:** Searching for a way to rebuild myelin at sites of damage by stimulating oligodendrocytes.

• **Stephen Hauser, M.D.**  
  **Funding:** $835,014  |  **Summary:** Banking genetic material from individuals and families with MS as a shared resource for studies searching for genes that confer susceptibility to MS.

• **Shen-Yi Howng, Ph.D.**  
  **Funding:** $150,800  |  **Summary:** Studying brain cells that contribute to myelin formation and destruction in MS for clues to reducing nervous system damage.

• **Sabeen Lulu, MBBS**  
  **Funding:** $130,000  |  **Summary:** Training in how to design and conduct clinical trials to find better treatments for people with MS.

• **Ellen Mowry, M.D.**  
  **Funding:** $1,312,902  |  **Summary:** A clinical trial investigating whether vitamin D supplements can alter disease activity in people with MS who are taking a standard therapy.

• **Jae Kyu Ryu, Ph.D.**  
  **Funding:** $156,515  |  **Summary:** Studying a molecule that may trigger immune attacks that cause nerve tissue damage in MS.

• **Ulf Schulze-Topphoff, Ph.D.**  
  **Funding:** $175,804  |  **Summary:** Studying the extent that immune system cells influence immune attacks that damage the nervous system in MS for clues to improving therapy.

• **Michel Varrin-Doyer, Ph.D.**  
  **Funding:** $175,804  |  **Summary:** Investigating how immune system cells are activated in a disease with some features similar to MS.

• **Emmanuelle Waubant, M.D., Ph.D.**  
  **Funding:** $1,064,639  |  **Summary:** Clinical trial to determine if a drug approved for ALS can protect brain and spinal cord tissues in MS.

• **Tracy Yuen, Ph.D.**  
  **Funding:** $163,103  |  **Summary:** Studying how glucocorticoids may influence the repair of myelin.

• **Scott Zamvil, M.D., Ph.D.**  
  **Funding:** $614,600  |  **Summary:** Distinguishing immune responses in MS and neuromyelitis optica.

• **Scott Zamvil, M.D., Ph.D.**  
  **Funding:** $755,378  |  **Summary:** Investigating mechanisms involved in the development of immune cells that can turn off the immune attack in MS, and how to stimulate their activity.
WE DEVELOP AND ALIGN HUMAN, BUSINESS AND FINANCIAL RESOURCES TO ACHIEVE BREAKTHROUGH RESULTS.
A woman in Sacramento connects with the Northern California Chapter to receive financial assistance and improves the quality of her life.

“I want to thank the National MS Society for helping to relieve this immense financial obstacle in my life. The Society’s generosity and good will are a tremendous inspiration to me, and has afforded me much-needed peace of mind. I am humbled by the National MS Society’s assistance. I will never forget this gift that you have given me and hope to pay forward the very spirit of the National MS Society to others.”

- Kimberly, Sacramento, CA
WE ARE ACTIVISTS.
MAKING OUR VOICES HEARD

Change happens through MS activism. The National MS Society and MS activists nationwide relentlessly advocate every day for federal policies and government programs to benefit the lives of people with MS and their families. The MS California Action Network and the chapter’s Government Relations Committee advocate for research money and healthcare rights, striving to promote public policy in the best interest of people with MS.

Nationally, MS activists raised their collective voice to encourage legislative progress on a wide range of issues that benefit the lives of people with MS and their families, spreading awareness, cultivating positive change, and achieving real-world results. In Northern California, MS activists advocated for change at the state and local level.

IN 2012, WE ACCOMPLISHED THE FOLLOWING:

- Activists helped to secure an additional $3.8 million in MS research funding through the Congressionally Directed Medical Research Programs, bringing our total through this funding source to more than $20 million since MS was first listed as eligible for funding in fiscal year 2008.

- Activists advocated for passage of state legislation curbing out-of-pocket costs for MS therapies and other prescription drugs. They collected personal stories of community members, as well as data illustrating the effects of high out-of-pocket costs, sharing them during in-person meetings with all 42 members of the state legislature.

- The Chapter activated our 5,000 member Action Alert network to write letters in support of five National MS Society supported federal bills before Congress and four pieces of legislation before the state legislature.

- Activists conducted over 30 face-to-face visits with members of Congress, advocating for the retention of critical services and benefits for people on both Medicare and Medicaid.

- For our priority legislation to curb out of pocket medical costs, the Chapter garnered media coverage in The New York Times, The Huffington Post, Washington Post, NPR Radio and KCRA TV.

- Over 40 new MS Activists enrolled in an extensive training program designed to enhance their ability to work directly with policy makers on the federal, state and local levels of government. They have each committed to recruit at least two more people into our Chapter’s Advocacy efforts within the next year.

A man in San Francisco meets with key legislators in the MS Congressional Caucus to discuss the importance of continuing to fund MS programs in the context of the fiscal cliff, and connects them to the mission to create a world free of MS.

“Having been involved in politics my entire professional career, I know first-hand the value of contacting, getting to know, and supporting (if it is justified) your members of Congress and your state legislature.”

- Al, San Francisco, CA
WE ARE LEADERS IN THE WORLDWIDE MS MOVEMENT, MOBILIZING MILLIONS OF PEOPLE TO DO SOMETHING ABOUT MS NOW.
The National MS Society values every contribution of the more than 500,000 Society volunteers throughout the country who provide support for people living with MS and their families. The total number of volunteer hours dedicated to the Northern California Chapter in 2012 was 13,043, which is equivalent to 7.2 full-time employees.

In 2012, the Northern California Chapter engaged with:
- 614 individuals who contributed 1,842 hours at 12 Walk MS: Northern California events.
- 250 individuals who contributed 1,900 hours at Bike MS: Waves to Wine Ride.
- 18 individuals who volunteered 1,444 hours in Northern California Chapter offices, the equivalent to almost one full-time employee working for one year.
- 73 volunteers who led 48 self-help groups in Northern California.
- 25 Government Relations Committee volunteers who advocated for policy and legislation change state-wide.
- 10 peer counselors who provided emotional support.

Event volunteers not only shared with us their time and hard work, but also raised over $20,000!

**EVENTS TO CREATE A WORLD FREE OF MS**

**Do It Yourself Fundraising**
Do It Yourself Fundraising is an opportunity for people with a deep commitment to the MS cause to raise awareness and critical funds for the MS Movement in new and creative ways. In 2012, events like crab feeds, runs, lemonade stands, and motorcycle rides raised over $48,000.

**DuskBuster**
DuskBuster is a 5K Run/2-Mile Walk starting at the South Tunnel of Polo Fields in San Francisco’s Golden Gate Park. The evening event is great for running teams and enthusiasts, casual joggers, walkers, families and more. In 2012, over 200 people raised over $33,000.

“A woman in San Francisco volunteers for the Northern California Chapter and makes connections that count.

“I still have some good tools left in my toolbox and I will continue to use them as long as I can and volunteering at the MS Society has allowed me to use those good tools.”

- Joyce, Walnut Creek, CA
Walk MS connects people living with MS and those who care about them. It is an experience unlike any other - a day to come together, to celebrate the progress we’ve made and to show the power of our connections. In 2012, over 7,500 participants raised $1.51 million across 12 walk sites in Northern California communities. Walk MS was held in the East Bay (Oakland), Eureka, Folsom, Modesto, Monterey Bay, Sacramento, San Francisco, San Jose, Santa Rosa, Solano County, Walnut Creek, and Yuba City.

**TOP INDIVIDUALS**

1. Janine Vanier - $60,820  
2. Perry Ann Jeveli - $34,424  
3. Steve Apperson - $16,570  
4. Mindy Zuckerman - $12,800  
5. Navid Mansourian - $11,585  
6. Jamie Zaninovich - $10,745  
7. Linda Pfeiffer - $8,180  
8. Shauna Silva - $6,985  
9. Monica Kirkorian - $6,870  
10. Brian Garber - $6,700

**TOP TEAMS**

1. Team MSters - $64,267  
2. Pear’s Pack - $44,650  
3. Linda’s Marvelous Supporters - $33,885  
4. Team Apperson - $17,105  
5. Team Victorious - $16,115  
6. Fighting Five - $13,095  
7. Laurie’s MS Stompers - $12,980  
8. Val’s Pals - $12,775  
9. KeepOnSmyelin :) - $12,215  
10. Team Zaninovich - $11,390

A woman in Modesto participates in Walk MS so she can raise money for others diagnosed with MS and connects to say thank you.

“I walk simply to walk... To find time to stop and thank all the people who are walking to make a difference in my future. I fundraise for the future of my fellow mankind, to discover a cure so someday we all can walk.”

- Sheila, Modesto, CA
Bike MS: Waves to Wine, September 22-23, 2012

Don’t just ride, Bike MS. It’s not just the miles that matter - it’s the unforgettable journey. In 2012, over 2,000 participants rode up to 175 miles over two days from San Francisco to Sonoma County, raising $2.36 million.

TOP INDIVIDUALS

1. David Patterson - $38,320
2. Chace Schornstein - $33,114
3. Nicole Schiereck - $30,874
4. Kate Aks - $27,532
5. Tom Galizia - $19,346
6. James Lau - $17,675
7. Jon Richards - $14,200
8. Rob Lytle - $14,185
9. Jon Lane - $13,630
10. Jim Forni - $12,768

TOP TEAMS

1. Salesforce.com & Friends - $186,904
2. Team Menstrual Cycles - $116,239
3. The Olympic Club’s Team Lemonade - $68,797
4. Deloitte Difference - $64,993
5. Team Chevron - $63,915
6. Cruisin’ with Susan - $59,724
7. NetApp & Friends - $54,704
8. Berkeley’s Anti-MS Crew - $47,760
9. Mitzvah Milers - $43,533
10. Barreling Up the Coast - $42,017

A woman from Pleasant Hill participates in Bike MS and connects with others committed to a world free of MS.

“Everyone associated with Bike MS is so nice and friendly and loyal to the event and the riders. It moved me to tears!”

- Diane Dodd, Pleasant Hill, CA
WE DEVELOP, DELIVER AND LEVERAGE RESOURCES TO ENHANCE CARE FOR PEOPLE WITH MS AND QUALITY OF LIFE FOR THOSE AFFECTED BY THE DISEASE.
EVERY CONNECTION COUNTS

The Northern California Chapter offers a variety of community-based programs to facilitate education, recreation, physical and emotional wellness, connection with others living with MS, and family communication.

Someone newly diagnosed first connects with the Society at one of the Chapter's Newly Diagnosed Orientation Programs.

The Chapter touched over 3,000 members during 120 educational, emotional health and outreach programs. This is a 20% increase in the amount of programming from 2011.

The Partners in MS Care program recognizes and supports quality MS care. The Chapter is partnering with health care professionals who demonstrate knowledge and experience in MS care, have a special interest in treating people living with multiple sclerosis, and work closely with the Society.

MS Specialists work closely with Chapter staff to connect patients to valuable services like financial assistance.

The Chapter assisted 164 people with over $93,000 of direct financial assistance for things like durable medical equipment co-pays, home modification projects, transportation and respite care. We were also able to increase our assistance by an additional $27,000 with funds leveraged from other community agencies.

A man in Merced takes advantage of a chapter service and gains freedom and mobility.

“Thank you, Terry’s Loan Closet, for making my life so much better. You have given me the opportunity to stay mobile and keep walking. Since I got the WalkAide I haven’t fallen once.”

- Danny, Merced, CA
Participants in community programs, such as self-help groups, connect and share resources with each other.

In Northern California 48 Self-Help Groups served over 700 people in 2012. National MS Society self-help groups have different purposes and goals. Groups may focus on support, advocacy or education. Each group gives its members an opportunity to connect with others living with the disease and learn about important community resources.

A member of a local self-help group learns about the MS Navigators at the National MS Society and calls to receive local resources.

The Information and Referral Center served over 1,700 individuals through 6,000 interactions. The highly skilled MS Navigators at the IRC offer the latest information about MS and provide referrals to local MS Care Centers, physicians and service providers.

A person living with MS who is worried about their employment is speaking to an MS Navigator at the IRC and learns about the Chapter’s Employment and Benefits program.

52 people had individual consultations with an employment and benefits specialist to learn about things like applying for disability benefits and reasonable accommodations at work.
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The following corporations, foundations, and individuals donated or raised $1,000 or more between October 1, 2012 and September 30, 2012. Our sincere apologies if we have inadvertently omitted anyone from this list.

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WHAT WILL YOUR CONNECTION BE?