Milestones on a Three-Decade Journey

1981
National Alopecia Areata Foundation is formed.

1985
First of 26 NAAF Conferences is held in Las Vegas.

1989
NAAF awards a record $134,000 for research.

1991
NAAF testifies before a Congressional Subcommittee, the first of 10 oral testimonies.

1992
First of 1,218 NAAF Conference Scholarships is awarded.

1995
The 10th Annual Conference, the largest to date, welcomes over 700 attendees in Boston.

1997
NAAF leads 600 volunteers to advocate on Capitol Hill on behalf of alopecia areata research.

2007
First federal bill (HR5936) providing insurance coverage for alopecia areata patients is introduced.

2008

2010
Newly initiated timeline for NAAF’s Treatment Development Program (TDP) accelerates our quest for a viable treatment.

Annual Report for the Fiscal Year Ending December 31, 2011
Dear Friends,

Take a trip back in time with us. The year is 1981. Ronald Reagan is sworn in as the nation's 40th president. Indiana Jones first cracks his whip in *Raiders of the Lost Ark.* “Bette Davis Eyes” rules the airwaves, at least until the arrival that summer of a small cable channel called MTV. The Dodgers beat the Yankees in six games to clinch the World Series. And a rare autoimmune disease that has afflicted men and women, children and adults since the beginning of recorded time at last meets its equal in the founding of an organization called the National Alopecia Areata Foundation (NAAF).

It may sound strange now but in 1981 we barely knew how to spell alopecia areata, let alone treat it. What we did know was that this was an unusually tenacious disease, and as such it would require an unusually tenacious organization, led and staffed by unusually tenacious people, to combat it. Researchers, board members, support counselors, staff—everyone affiliated with NAAF—would have to be uncommonly committed to supporting the people who suffer from alopecia areata, raising the awareness of their families and friends, and eradicating the disease itself.

While alopecia areata has plagued sufferers for several millennia, in just the past three decades NAAF has made bold strides in genetic and dermatological research barely imaginable in 1981. And NAAF will continue to do so until the causes of this disease are discovered, the people who are susceptible are protected, and alopecia areata is finally defeated. In 2011, our Treatment Development Program scored an almost dizzying amount of accomplishments—such as funding an alopecia areata autoantibody study at the University of Colorado, leveraging resources for a new alopecia areata biomarker study at Columbia University, and officially contracting with MD Anderson Cancer Center in Houston, Texas, to take over the responsibility for funding the National Alopecia Areata Registry.

Until alopecia areata is a thing of the past, our efforts remain unabated. Our peerless support network continues to make real differences in the lives of both individuals and their families. We are finding new ways to raise awareness, most recently through successful events with Major League Baseball teams. New media allows us to spread news, support, and hope through Facebook and Twitter in addition to more traditional venues like our website, informational materials, DVDs, electronic and print newsletters, the Ascot Fund and the Alopecia Areata Marketplace.

In 2011, we saw the cumulative effect, the real progress, stemming from thirty years of persistent effort. Today NAAF renews its pledge to provide all of its energy, resources and guidance to our community; we hope you know we will always advocate, always research, always fight beside you and with you on behalf of people with alopecia areata. NAAF and its members stand more united than ever in a common purpose staked by three decades of research, support and awareness.

One can only imagine what the future years will bring . . .

Brian Ter Haar  
Chair, Board of Directors, 2012

Vicki Kalabokes  
President and Chief Executive Officer

*The nation’s leading nonprofit watchdog group, the BBB Wise Giving Alliance, has found that NAAF operates with the utmost integrity, transparency, and accountability. NAAF is proud to be one of the very few nonprofits (out of the almost 1.5 million nonprofits in the United States) that meet these strict standards. In addition, NAAF meets all 44 of the standards required for membership in the National Health Council.*
“Thank you for the opportunity to learn and be part of this year’s conference. It is a wonderful thing to have the scholarship fund. It made the conference happen for me! Thanks for all efforts.”

- NAAF’s 26th Annual International Conference, held in Los Angeles, was a huge success thanks to a program full of new and fun events, a lineup of great guest speakers, a fantastic hotel, and the hard work of our incredibly devoted volunteers and staff. It was an unforgettable experience for all 832 attendees.

“Thank you so very much for the scholarship you gave me to attend the LA Conference! It was great to see everyone again and make new friends. It was my first time to California and I loved it! I am considering becoming a Support Group Leader and at the conference I met a family from nearby that is in need of support. So that was a step toward success!”

- 97 Conference Scholarships were awarded to people in need, enabling them to attend the conference. A total of 1,218 scholarships have been awarded since the inception of our conference scholarship program.

- Conference attendees came not just from 37 states, but from Australia, Canada, Hong Kong, Japan, New Zealand, Panama and Sweden.

“I’m writing to express my most sincere gratitude for making it possible for my daughter and I to stay at the conference hotel for this year’s NAAF conference. As a single mom, it’s been hard over the years to make out-of-town trips happen for us. This was by far the most awesome traveling experience we’ve had together. When my daughter was 7 she was diagnosed with alopecia areata. Until then I’d never even heard of it. The last 3 years have seemed to make me an expert on the subject, but no amount of research can prepare you for the feeling of comfort, community, joy, and acceptance that she and I both experienced over the conference weekend.”

- The NAAF website provided up-to-the-minute information on research, support and resources, programs and events, and much, much more. As of December 31, 2011, it had received over 13 million hits. Go to www.naaf.org to see for yourself!

- The NAAF Facebook page and Twitter feed offered the latest news regarding the alopecia areata community and its doings. Best of all, you got to interact with us by sharing your comments and questions. We loved hearing from you! As of December 31, 2011, the NAAF Facebook page had over 400 fans and over 4,000 likes, and the NAAF Twitter feed had over 800 followers.
• Our NAAF Electronic News, *Beneath the Surface*, continued to provide a quick-reading roundup of information important to some 15,000 readers in the alopecia areata community.

• A total of 87 men, women and children received financial grants from the Ascot Fund in order to purchase hair prostheses. These grants make an incredible difference in the lives of those affected. Since this very special program was established in 2004, a total of 501 people have been helped.

• Some 54 different merchants were accessible through the Alopecia Areata Marketplace, each providing hard-to-find and much-needed products to those with alopecia areata. The variety and quality of the products were exceptional.

• NAAF had 130 knowledgeable volunteers in its International Support Network, serving 111 cities worldwide, organizing local support groups and providing telephone support. NAAF highlighted support volunteers in each quarterly newsletter.

• Alopecia areata kids are awesome . . . and they know it! That’s why NAAF offered a wide selection of resources just for them, including a nine-minute DVD/video that kids could share with family, friends, and schoolmates; a school guide for use by parents and teachers; a bibliography of books for kids with alopecia areata; children’s support groups; our Children’s Conference Camp attended by 250 kids ages 5 to 17; our Pen Pals program; secure and private message boards for children and teens; a special KidNet section of the NAAF Newsletter; and brochures for parents and kids.

• 1,300 participants, both adults and kids, actively took part in our Pen Pal program. (And you can too!)

• NAAF helped more people because we had more people helping us, in the form of thousands of volunteers worldwide.

• Our quarterly award-winning *NAAF Newsletter* kept you updated.

• Our four primary NAAF brochures (*Alopecia Areata: What You Should Know; A Parent’s Guide; Helping You Cope; and From a Child’s Point-of-View*) were all revised to be more up-to-date and easier to read.

“In helping others, we shall help ourselves.”
—FLORA EDWARDS

1995
The 10th Annual Conference, the largest to date, welcomes over 700 attendees in Boston.

2001
Veteran character actor Barry Corbin closes the Annual NAAF Conference in Oakland with a recitation of a soliloquy from Shakespeare’s *As You Like It*.

2004
First of 501 Ascot Fund grants is awarded.

2001
The Alopecia Areata Marketplace is established.

2007
22nd Annual NAAF International Conference draws a record 1,000 participants to Washington, DC.
Miss Delaware 2010 Kayla Martell used her unique position as a Miss America contestant to work on behalf of alopecia areata awareness, as well as attending NAAF’s 26th International Conference in Los Angeles.

“As Miss Delaware and now in my regular life my priority each day is to help one more person know what it means to have alopecia areata; then I know I’ve done a good deed that day and I’ve changed one life.” — KAYLA MARTELL

Kayla was also present at the first annual “Have a Ball and Bowl” fundraiser, presented by the Wilmington Delaware Support Group. The kids enjoyed learning from Kayla that alopecia areata has not prevented her from achieving her goals. Everyone loved a day of knocking down pins, and close to $9,000 was raised for alopecia areata research.

NAAF partnered with 14 Major League Baseball teams to present 22 Awareness Events, beginning with Rebecca Hibbs singing the National Anthem at a Nashville Sounds game in May and climaxing with a slew of events during Alopecia Areata Awareness Month in September. Events included pregame ceremonies, awareness booths, pitching clinics, honorary batboys and girls, video PSAs, honorary first pitches, scoreboard messages . . . even an in-game interview. In total 14,000 NAAF bracelets were distributed and $15,000 was raised.

Happythankyoumoreplease, a comedy-drama featuring a female character with alopecia areata, came to theaters in March. Written and directed by Josh Radnor (Ted Mosby on How I Met Your Mother), the film featured a character named “Annie” based on Josh's real-life friend, Rachel Fleit. The Fleits have been longtime supporters of NAAF, and Rachel was our guest speaker at the 2010 Annual NAAF Conference in Indianapolis. Happythankyoumoreplease is now available on DVD, and we highly recommend it!

“William James once said, ‘Act as if what you do makes a difference. It does.’ That is exactly the feeling of the Tortoise & Hair™ Pittsburgh staff. In 2007, my feelings about this disease were ones of anger, loneliness and sadness. Today? I just feel blessed.” — AMANDA SHANNON

The Tortoise & Hair™ run/walk continued as NAAF’s premier nationwide awareness and fundraising event. Almost $7,000 was raised at the 5K run in Wadesboro, North Carolina, put together by Hoy and Gwen Lanning on October 7. Two days later at a Pittsburgh run/walk organized by Amanda Shannon 120 people registered to participate and 15 organizations sponsored the event.

On October 27, 2011, Pat’s Fund (www.patsfund.org) held its annual Living with Autoimmunity luncheon, focusing on the tendency of autoimmune diseases to run in families. The luncheon was also an opportunity to announce the launch of the Autoimmune Advocacy Alliance (A3). A3 seeks to coordinate the advocacy organizations related to more than eighty autoimmune diseases in one consortium that can more effectively raise awareness of these conditions. With the launch of 1985 Congress declares July 7 to 13 Alopecia Areata Awareness Week.

1990 NAAF gets full hour segment on the Geraldo Show, as well as segments on Northwest Washington Afternoon, P.M. Magazine, People Are Talking, and The Today Show.

1994 Benefit screening of the film North raises $60,000 for NAAF and the Children’s Village in New York.

1998 NAAF receives the Excellence in Education Award from the American Academy of Dermatology (AAD) for the information it provides to patients, families, physicians, politicians and the general public.
A3, we are all hopeful that the term “autoimmune” will become a household word and will give the 50 million people who are afflicted with these diseases, and their family members who are at risk, a louder, more focused voice to ask for help, understanding, and answers.

- Nearly 160 people attended Let Your Hair Down for Sarah in Chicago. The event, presented in May by Mike and Cheri Walczak for their daughter, raised $6,474.
- More than 400 gathered at Wilmington, Delaware’s Sanford School Sports Center in September to take part in the Incredible Back-to-School Wig Out, a fundraising event hosted by Sanford School supporting NAAF. J. Christian and his team volunteered their time and talent to cut and style hair, providing free haircuts for those donating their hair. Proceeds included 16 donated ponytails totaling 168 inches of hair, and $3,200.

“Thank you, NAAF, for all you do, and may we find a cure one day. We know that the money we raised will be used wisely and will help many others.” — JEN, MARK AND MAX TRINKO AND FAMILY

- Milwaukee witnessed the Second Annual Tournament of Champions and Family Cookout for NAAF in September. Presented by Josh and Judy Lang, and Jen and Mark Trinko, the event featured “Kiah’s Carnival” (named for the Lang’s daughter Kiah, who suffers from alopecia areata) with lots of activities for kids, and a silent auction drew more than 150 sponsors. Approximately 350 attended, raising $10,341.
- The second annual Christopher’s Run for Alopecia Areata 5K Run and Mile Walk, was held in Laurel Springs, New Jersey, in September. The run was organized by both the Passarella and Manuala families to honor Christopher Passarella, a 10-year-old who has suffered from alopecia areata since the second grade. Over 75 attended the event, including the Mayor of Gloucester Township, raising $6,400 for NAAF.
- NAAF continued its fight for insurance coverage of alopecia areata on the Federal Level while volunteers like Heidi Bratt in Massachusetts continued to advocate at the state level.
- NAAF volunteers hosted fundraising and awareness events around the world, generating almost $218,760 for NAAF. All of these events are listed by state (or province) in this annual report. We thank all who hosted—or participated in—these fantastic events!
NAAF gratefully acknowledges all of the AWESOME individuals and families that raised money and spread awareness on our behalf during 2011.

Total Raised: $218,760!

Alabama
Jennifer McCrary Family & Friends

California
Kylie Bamberger Family & Friends
Gannon Hubbell Family & Friends
Natalia & Marco Castillo Family & Friends
     Human Race
Los Angeles Tortoise & Hair
San Diego Support Group
Emma Robbins Family & Friends
Carlene Willis Family & Friends

Connecticut
Coginchaug High School Dress Down Friday
Connecticut Support Group
Kim Martino Family & Friends
McDonald Family & Friends

Delaware
Debora Pellicano Family & Friends

Florida
John Chapin Family & Friends
Parikh Family & Friends

Illinois
Ken & Jennifer Anderson Family & Friends
Chicago Support Group
Gardner Elementary School
Levi, Ray & Shoup
Cheri Walczak Family & Friends

Indiana
Maia Campbell Family & Friends
Nettles Family & Friends

Iowa
Friends of Aplington-Parkersburg Schools

Kansas
Rebecca Lading Family & Friends

Kentucky
Emilee Lockwood Family & Friends

Louisiana
South Beauregard Elementary School
Abby Ter Haar Family & Friends

Maryland
Zachary Sents Family & Friends

Massachusetts
Carla Couilliard Family & Friends
Westminster Elementary School

Michigan
Linda Cauley Family & Friends
Community Programs Jeans
Mallory Crown Family & Friends
Venicia Focht Family & Friends
Cassie McElwain Family & Friends

Minnesota
Rose Gallagher Family & Friends

Missouri
Rogers Middle School

North Carolina
Samuel Lanning Family & Friends

New Jersey
Central School
Lilliana Hakim Family & Friends
Christina Manuola Family & Friends
Christopher Passarela Family & Friends

New York
Anonymous
Harrison Goldberg Family & Friends
Lise Hernandez Family & Friends
Lord & Taylor
Erica Meilhede Family & Friends
Margaret Staib Family & Friends
Lexi Steinman Family & Friends
Madison Urraro Family & Friends

Pennsylvania
Delta Phi Epsilon
Friends of The Philadelphia 76ers
     Awareness Fun Day
Glass Family & Friends
Hair Direct
Theresa Hibbs Family & Friends
Macy Matzner Family & Friends
Amanda Shannon Family & Friends
Coleen Van Osten Family & Friends

Tennessee
Rebecca Hibbs Family & Friends

Texas
Brandi Battenfield Family & Friends
First Command
Mackenzie Hibert Family & Friends
Michael Maddux Pitching Clinic

Utah
Cassie Lindsay Family & Friends

Virginia
Courtney Parker Family & Friends

Washington
Jacqui Meaker Family & Friends

Wisconsin
Kiah Lang Family & Friends
• NAAF along with the Coalition of Skin Diseases continued to advocate for the importance of skin disease research so that all skin diseases can benefit.

• NAAF took a very active role in educating medical professionals about alopecia areata, hosting informational exhibits at key medical and scientific conferences, including the American Academy of Dermatology, the Dermatology Nurses’ Association, the Society for Investigative Dermatology, and the American Academy of Pediatrics.
NAAF’s Treatment Development Program is on track with the timeline initiated in 2010 to accelerate our quest for a viable treatment. NAAF serves as the ultimate concierge, leveraging all of our available research resources and clinical partnerships. Our strategic goal is to produce safe, effective, affordable treatments useful to millions of people with alopecia areata. Recent additions and developments to this critical program include the following actions:

- NAAF funded a study in the Eisenbarth Laboratory at the University of Colorado’s Barbara Davis Center for Childhood Diabetes to find the autoantibody related to alopecia areata. This research seeks to quickly build upon Dr. Angela Christiano’s genetic research by utilizing the well-established laboratory of a related autoimmune disease and their well-developed technologies.

- NAAF funded another laboratory at the University of Colorado to utilize another method to find the antigen targeted by the autoantibody related to alopecia areata. We want to see if it generates the same result as the study at the first laboratory.

- NAAF also coordinated the study of 210 serum samples from the National Alopecia Areata Registry sent to the laboratory of Dr. Yehuda Shoenfeld in Israel to find antibodies to infectious agents and to investigate antibodies similar to those for other autoimmune diseases.

- NAAF organized an Alopecia Areata Clinical Research Summit to identify and address issues and opportunities to facilitate the testing of new treatment candidates. All recommendations from that summit were implemented in 2011.

- NAAF, along with Dr. Christiano, coordinated the submission of clinical study applications to the National Institute of Arthritis and Musculoskeletal and Skin Diseases (NIAMS) so there aren’t competing alopecia areata grants.

- Dr. Christiano was awarded a grant by the National Institutes of Health to identify biomarkers for alopecia areata—one of the goals of the Alopecia Areata Clinical Research Summit.

- NAAF continues to negotiate with several companies to establish partnerships and conduct clinical trials on possible treatments for alopecia areata.

- NAAF coordinated an update of alopecia areata prevalence and incidence studies. This data determines the scope of the problem, which is of interest to commercial partners and to the Food and Drug Administration.

- NAAF coordinated the transfer of funding of the National Alopecia Areata Registry from the National Institute of Arthritis and Musculoskeletal and Skin Diseases (NIAMS, a part of the NIH) to NAAF. This is a huge financial commitment for our Foundation but this Registry is a critical resource for future research, particularly clinical trials.
The AAD, the Society for Investigative Dermatology (SID) and the Coalition of Skin Diseases (CSD) join together to sponsor the First Annual Skin Disease Research Day, which is dedicated to advocating for increased federal funding for research.

NAAF funded Quality of Life Studies to objectively demonstrate the burden of alopecia areata. This data will also be presented to the Food and Drug Administration when we find a treatment.

NAAF organized and conducted a scientific briefing for companies with product development interests in treatments for autoimmune diseases, dermatological conditions, and hair growth or restoration. NAAF encouraged healthcare companies to develop treatments for alopecia areata.

NAAF continued to update the plan for the Treatment Development Program and monitor whether changes are needed to advance to our ultimate goal more quickly.

The National Alopecia Areata Foundation had an exhibit booth to attract new research investigators to study our disease during the Society for Investigative Dermatology Meeting in Phoenix, Arizona, in May 2011. In addition, there were separate meetings for alopecia areata immunology researchers, National Alopecia Areata Registry researchers, and the Clinical Research Task Force. All these meetings advance the Alopecia Areata Treatment Development Program.

RESEARCH GRANTS

Daisy Dai, PhD  |  Dennis Roop, MD, PhD  |  University of Colorado School of Medicine, Denver, Colorado

*Using the sera of alopecia areata patients to identify antigens that are targets of the autoantibodies of alopecia areata patients*

This study identified genes of proteins to which there are autoantibodies in alopecia areata. The proposed project was designed to identify antigens recognized by antibodies produced by patients with alopecia areata. This study will likely produce candidate targets of the host immune system.

Tito R. Mendoza, MS, MEd, PhD  |  The University of Texas MD Anderson Cancer Center, Houston, Texas

*Quality of life in patients with alopecia areata*

At the NAAF Corporate Leadership Council meeting, the pharmaceutical companies requested that we have Quality of Life studies completed and published before they bring drugs to be approved by the Food and Drug Administration (FDA). So this study is a priority for NAAF.

The investigators developed a National Alopecia Areata Registry Symptom Scale (NAARRSS), following standard psychometric procedures, to perform secondary analysis of the National Alopecia Areata Registry (NAAR) data to describe the quality of life of patients with alopecia areata.

Researchers tabulated mean data values and summarized content validity index (CVI) from clinicians' ratings of the importance of each item as they relate to concerns of patients, drafted and tested a pilot symptom questionnaire, and then refined it for use with research subjects.

Sultan Mirzoyev, Medical Fellow  |  Rochelle Torgerson, MD, PhD  |  Mark Davis, MD  |  Mayo Clinic, Rochester, Minnesota

*Incidence of alopecia areata in Olmsted County, Minnesota*

The objective of this study is to assess the incidence and natural history of alopecia areata among unselected patients from the community. The last study that was published from the data in Olmsted County, Minnesota, was for the years 1975 to 1989. This is a follow-up to the previous study.

Alina Markova, MD  |  Alpert Medical School of Brown University, Pawtucket, Rhode Island

James A. Solomon, MD, PhD, Director of Ameriderm Research, was coordinating this study

*Incidence of alopecia areata in dermatology practice with 1.3 million patients*

Data from 69 offices of Advances Dermatology and Cosmetic Surgery and 72 offices of Ameriderm (141 offices in total) across the United States was being studied to determine incidence of alopecia areata.

2004

NAAF advocacy and science result in multimillion dollar funding from the NIH for the National Alopecia Areata Registry.

2000

NAAF embarked on Alopecia Areata Immunology Research Initiative to draft Immunology Research Program Development Plan.

2009

NAAF embarks on Alopecia Areata Immunology Research Initiative to draft Immunology Research Program Development Plan.

2010

Led by NAAF Scientific Advisory Council member Dr. Angela Christiano and using cases from the National Alopecia Areata Registry, a team of investigators at Columbia University Medical Center discover eight genes that contribute to alopecia areata, including one that plays a possible role in the onset of the disease.
## Statement of Financial Position
### December 31, 2011

<table>
<thead>
<tr>
<th>ASSETS</th>
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<tbody>
<tr>
<td><strong>Current assets:</strong></td>
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<tr>
<td>Cash and cash equivalents</td>
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<tr>
<td>Marketable securities</td>
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<tr>
<td>Accounts receivable</td>
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<tr>
<td>Prepaid expenses and other current assets</td>
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<tr>
<td><strong>Total current assets</strong></td>
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<tr>
<td>Fixed assets, net of accumulated depreciation</td>
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<td><strong>Total assets</strong></td>
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<tr>
<th>LIABILITIES AND NET ASSETS</th>
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<tbody>
<tr>
<td><strong>Current liabilities:</strong></td>
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<tr>
<td>Accounts payable</td>
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<tr>
<td>Accrued vacation payable</td>
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<tr>
<td><strong>Total current liabilities</strong></td>
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<tr>
<td><strong>Net assets:</strong></td>
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<tr>
<td>Unrestricted</td>
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<tr>
<td>Temporarily restricted</td>
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<tr>
<td><strong>Total net assets</strong></td>
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<tr>
<td><strong>Total liabilities and net assets</strong></td>
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## Statement of Activities for the Year Ended
### December 31, 2011

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<th>REVENUES</th>
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<tbody>
<tr>
<td><strong>Donations – general</strong></td>
</tr>
<tr>
<td><strong>Donations – stock and major donors</strong></td>
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<tr>
<td><strong>Restricted grants and donations</strong></td>
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<tr>
<td><strong>Conference</strong></td>
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<tr>
<td><strong>Special events – Links for Locks and Tortoise and Hair</strong></td>
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<tr>
<td><strong>Prior year grant reimbursements</strong></td>
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<tr>
<td><strong>Newsletter income</strong></td>
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<tr>
<td><strong>Alopecia Areata Marketplace</strong></td>
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<tr>
<td><strong>Brochures and videos</strong></td>
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<tr>
<td><strong>Interest and dividend income</strong></td>
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<tr>
<td><strong>Net realized and unrealized gains on marketable securities</strong></td>
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<tr>
<td><strong>Net assets released from restrictions</strong></td>
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<td><strong>Total revenues</strong></td>
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<tr>
<th>EXPENSES</th>
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<tbody>
<tr>
<td><strong>Program services:</strong></td>
</tr>
<tr>
<td>Research</td>
</tr>
<tr>
<td>Education and support</td>
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<tr>
<td><strong>Supportive services:</strong></td>
</tr>
<tr>
<td>Fundraising</td>
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<tr>
<td>Management and general</td>
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<tr>
<td><strong>Total expenses</strong></td>
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<tr>
<td><strong>Change in net assets</strong></td>
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<td><strong>Net assets, beginning of period</strong></td>
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<td><strong>Net assets, end of period</strong></td>
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<table>
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<tr>
<th>Spending Percentage</th>
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<tbody>
<tr>
<td><strong>Program, Support, Research and Awareness</strong></td>
</tr>
<tr>
<td><strong>Fundraising</strong></td>
</tr>
<tr>
<td><strong>Management</strong></td>
</tr>
</tbody>
</table>

The Statement of Financial Position of the National Alopecia Areata Foundation as of December 31, 2011, was audited by R. J. Riccardi, Certified Public Accountant. A copy of the complete report, including notes, is available for public review from the National Alopecia Areata Foundation, 14 Mitchell Boulevard, San Rafael, CA 94903, or on the website www.naaf.org.
重大捐赠者—$2,500 TO $4,999
Dr. Rex & Johnnie Amonette • Maria Beckett • Andrea & Robert Berger • Claire Chardinet • Arleen & Richard Glenn • Marcia & Jerry Gross • David & Ami Handler • Matthew & Fred Levin • Leslie Dumont • Shirley Levitt • Jessica & Steve Libowitz • Emily Lockwood • Terri McGee • family & Friends • Michael & Christine Moscare • Scott Mroz & Claudia Held • Rita & Robert O’Connor • Kate Paley • Marcia Rimm & family • Richard & Linda Riesing • Dr. Joseph & Patricia Russ • Jennifer Nordstrom • family & Friends • Clay & Sharon McGee • T J Morris • Rose & Henry Moskowitz • Gary & Camie Murphy • Miry Myer & family • Gary & Marla Nathanson • Susan Amanda Shannan • family & Friends • Ranjit Thaker • Colleen & James Van Oster • Ed & Mary Wojtowicz

企业
HarDirect, Bainbridge, PA • Johnson & Johnson • AML & Company (Cincinnati, OH) • Bosley Medical, Beverly Hills, CA

非营利组织
Hedsker Foundation for Children, NY

银级捐赠者—$10,000 TO $24,999
Anonymous • Caroline & David Brown • Elizabeth Catto-Shaw • Richard & Rosa Dest • Brian & Hana Eldred • Robert & Anna Flint • Guy & Marilyn Elizabeth Catto-Shaw • Richard & Rosa Dest • Brian & Hana Eldred • Robert & Anna Flint • Guy & Marilyn

非营利组织
Café, Houston, TX • The Richard S Klingenstein Family Foundation
GRAND PATRONS

$250 TO $499

Dana & Mary Leonard & Mary Lyne Albis
Avenue Schenone • Tom & Jackie Ahrens
Candace Akerson • Akerson Jeremy • Jon &
Kathleen Andry • Carolyn & Dean Anderson
• Andrea Andre • Tamar Brandon • Glenn Auze
• Jim & Lucy Baker • Lori Ball • William Bass
• Stephen & Rosemary Bedard • Olga Benda
• Melissa Berg • Tami & Kevin Best • Fran
Blanck • Kathleen Bock • Jennifer Booth
• Ray & Betty Braun • Margaret & Dennis Brennan
• John W. Buckley • Sandor • Andrea Buchus
• Dennis & Catherine Bye • Laura & Robert Byreu
• Laurel & John Campbell • Stephen & Christine
Campbell • Ralph Carr • III • Mary & Anthony
Cefalu • Donald & Patricia Chambers • John •
Suzanne Chartrand • Michael Chetko • Mindy
Chrytal • Nancy & Bob Clark • Susan & Roy
Cleveland • Michael Colpoy • Art Cooper
• Carla Coiledd • Family • Friends • Ian & Matilda
Dab & William Dab • Wilinda • William Dab •
• Darin B. Darwin, Jr. • Hochi De Los Santos • Diane
• Jennifer & DeFreece • Donald Deines • Celeste &
Octavi Delgado • Celeste Descotau • B uncis Dinczuk
• George E. Ding • Cheryl Ding • Craig Ellingsworth
• Nicole Erb • Jackie Erisc-Acadel • James Garcia
Eveloff • Jonathan Fein • Mark Ferreira • Luann
• Nicole Erb • Jackie Erisc-Acadel • James Garcia
Eveloff • Jonathan Fein • Mark Ferreira • Luann
Ferrante • Brendan Fischler • Ramsob • Debbi
Fuller • Jeff Fuller • Kevin & Karen Ggem •
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Kathy & Lee Gause • Diana & David Gerland
• Stephen Gevaiz • Gary & Doreen Gittings • Fran
• Kevin Golvich • Craig Goff • David Goldin
• Craig Grai • Dan Gronke • David Gronke
• Ernest Guenzburger • Ernst Rantube • Albert
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Hammond • Barbara Hansen • Harold Harms Jr.
• Craig Grai • Dan Gronke • David Gronke
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MISSION
The National Alopecia Areata Foundation (NAAF) supports research to find a cure or acceptable treatment for alopecia areata, supports those with the disease, and educates the public about alopecia areata.