

2010-2011 Annual Report



Making sure each person with cutaneous lymphoma gets the best care possible

OUR MISSION

The mission of the Cutaneous Lymphoma Foundation is to support every person with cutaneous lymphoma by promoting awareness and education, advancing patient care and facilitating research.

We exist to make sure that each person with cutaneous lymphoma gets the best care possible.

CONTENTS

3	What Is Cutaneous Lymphoma?
4	President's Letter
5	Chief Executive's Letter
6	Living With Cutaneous Lymphoma: Susan's Story
8	Programs and Services
9	2010-2011 Highlights
10	Financials
11-18	Donor Honor Roll
19	Board of Directors and Staff

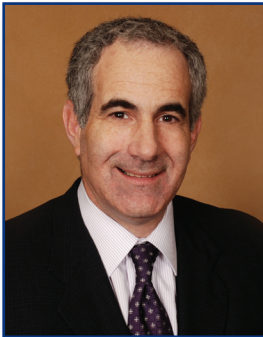
What Is Cutaneous Lymphoma?

Cutaneous lymphomas are cancers of lymphocytes (white blood cells) that primarily involve the skin. Classification is based on lymphocyte type: B-lymphocytes (B-cell) or T-lymphocytes (T-cell).

Cutaneous T-cell lymphoma (CTCL) is the most common type of cutaneous lymphoma that typically presents with red, scaly patches or thickened plaques of skin that often mimic eczema or chronic dermatitis. Progression from limited skin involvement is variable and may be accompanied by tumor formation, ulceration, and exfoliation, complicated by itching and infections. Advanced stages are defined by involvement of lymph nodes, peripheral blood, and internal organs.

Cutaneous lymphomas affect thousands of individuals worldwide. CTCL affects over 30,000 people in the United States and Canada. The incidence of cutaneous T-cell lymphoma in the United States is increasing with approximately 3,000 new cases being diagnosed annually. Due to the difficulty of diagnosing the disease in its early stages and the current lack of an accurate reporting system, prevalence of cutaneous lymphoma is thought to be much higher.

President's Letter



Stuart R. Lessin, M.D.

For the Cutaneous Lymphoma Foundation, the past year was marked by enormous growth and transition. With Judy Jones' recent retirement from her founding roles as Foundation Director and President of the Board of Directors, the new leadership of the Cutaneous Lymphoma Foundation is looking forward to a new and exciting era that will continue Judy's legacy and her vision for the cutaneous lymphoma community: ensuring the best care for every person with cutaneous lymphoma.

As the new President of the Board of Directors of the Cutaneous Lymphoma Foundation, it is my privilege to serve. It is also my pleasure to share with you the accomplishments of the past year and opportunities that lie ahead.

The Board of Directors has grown to include greater representation of patients and expertise in finance, development, event planning, medical affairs and advocacy. The Board is fully engaged in Foundation business.

The Foundation's programs and services continued to grow last year and they now reach more people with cutaneous lymphoma than ever before. The first draft of the Foundation's new *Patient's Guide to Understanding Cutaneous Lymphoma* has been completed with a 2012 publication anticipated. This publication will serve as an authoritative source of information and support for all those diagnosed with cutaneous lymphoma. Plans are underway to have the Guide translated into as many languages as possible so that its impact will be global. The Foundation's 2011 re-launching of its web site (www.clfoundation.org) provides more user-friendly information and will serve as an important platform for expanding services.

As we move forward in the age of globalization, the patient's voice will have an increasingly important role in biomedical research and health care services. Technologies are enabling a more collaborative approach toward discovery and advancement by bringing together scientists, drug developers, government agencies and patients.

The Cutaneous Lymphoma Foundation now represents the largest non-profit organization supporting cutaneous lymphomas and is poised to assume a leadership role in the cutaneous lymphoma community. With the sustaining support from all stakeholders, the Foundation's mission will continue to be realized.

Stuart

Chief Executive's Letter



Jennifer Viano

Being diagnosed with cutaneous lymphoma presents so many challenges for patients and their loved ones. Cutaneous lymphoma is a complicated disease with many different forms and treatment options. And, many patients find it difficult to obtain an early, definitive diagnosis.

It's not surprising then that our recent needs assessment revealed that patients ranked learning about cutaneous lymphoma, learning about treatment options available and locating treatment specialists among their top needs. And, one message came through loud and clear: Patients want and need more information, resources and support to help them along their journey.

At the Cutaneous Lymphoma Foundation, we're playing an important role in helping patients, loved ones, medical professionals and all those affected by cutaneous lymphoma through comprehensive programs and services that deliver positive outcomes and help improve the quality of life of patients.

During the 2010-2011 fiscal year, we were able to build upon the extraordinary success of the Foundation's first 14 years by expanding and enhancing programs, connecting with more patients, growing our Board of Directors, and much more. I hope you enjoy reading more about all that we accomplished on page 9.

Without the help of our generous supporters, none of our work would be possible. The incredible generosity of our individual and corporate supporters is humbling. We are deeply grateful to each person and company who contributed the vital funds necessary to fulfill our mission.

Our 2010-2011 fiscal year was one of impressive progress. But, together, we have so much more to do to make sure that each person with cutaneous lymphoma gets the best care possible.

A handwritten signature in black ink, appearing to read 'Jen', with a large loop at the bottom.

Living With Cutaneous Lymphoma: Susan's Story

When Susan Ferguson first turned up at her doctor's office with an itchy, flaky rash in 2003, she had no idea what "CTCL" meant, but her dermatologist had a suspicion, and he requested a biopsy to exclude it. The report came back "equivocal."

By 2005, the rash had spread and could not be ignored. The prescription ointments and creams she tried did not work. She then researched "CTCL" -- cutaneous T-cell lymphoma? T-cells? Lymphoma? Susan was frightened. A new biopsy and this time the diagnosis was unambiguous: patch-stage mycosis fungoides (MF). Stage 1B.

Luckily for Susan, she lives in Greenwich Village and was only a taxi ride away from top specialists at NYU Langone Medical Center. Nevertheless, every night she was on the Internet searching for facts. She found the Cutaneous Lymphoma Foundation and combed the archives. She found Judy Jones, founder of the Foundation. She joined the CTCL-MF Listserv. Now she wasn't alone.

After some initial nasty encounters with her insurance company, Susan started an effective treatment plan and attended every patient event held by the Foundation in New York. "I felt so grateful to be a part of this caring, compassionate community dedicated to helping patients," she shared.

Recently, Susan decided to write a new will and has made a generous provision in support of the Cutaneous Lymphoma Foundation. She commented, "Finally I've found a way to demonstrate my gratitude to the Foundation and my wish to do something good for my fellow patients."

Susan continues to grapple with her CTCL, but she is committed to being diligent about her treatment, care and overall health. She also uses her experiences to help others by contributing to the CTCL-MF Listserv and Foundation communications. Susan is passionate about what the Foundation has meant to her: "The programs, resources and support made available by the Cutaneous Lymphoma Foundation have been a source of great comfort to me. With the help of the Foundation, I know I am not walking this path alone."



“The programs, resources and support made available by the Cutaneous Lymphoma Foundation have been a source of great comfort to me. With the help of the Foundation, I know I am not walking this path alone.”

Susan Ferguson, Patient

Programs and Services

The Cutaneous Lymphoma Foundation offers comprehensive programs and services that provide relevant, timely information and resources that deliver life-changing, positive outcomes for patients, caregivers, loved ones, medical professionals and others affected by cutaneous lymphoma. These programs include:

Patient Educational Forums

These day-long events held throughout the year in the United States and Canada provide exceptional opportunities for people with cutaneous lymphoma to receive accurate information about the disease, treatment options, access experts in the field, and connect with other people who share in similar experiences.

Website

Our website at www.clfoundation.org is one of the most important resources we offer. At anytime, comprehensive information can be found on our website about the many types of cutaneous lymphoma, treatment options, treatment centers and specialists, programs and services, research and advocacy efforts, ways to become involved in the Foundation's work and much more.

Library of Printed/Electronic Education Materials

We offer a variety of printed and electronic education materials including fact sheets, archived teleconferences, newsletters and more. You can download or order these materials online at www.clfoundation.org or by contacting us at info@clfoundation.org or 248.644.9014.

Physician and Treatment Center Referral Network

Finding the best possible treatment and care is vital for every cutaneous lymphoma patient. Our website at www.clfoundation.org lists top cutaneous lymphoma treatment centers and specialists.

Personal Assistance via Phone and Email

Personal assistance is available at the Foundation via phone or email. If you are not able to get your questions answered via our website – or if you just want to talk with someone about anything related to your unique experiences and how you are affected by cutaneous lymphoma – please contact us via phone at 248.644.9014 or via email at info@clfoundation.org.

Research Funding

In recent years, more attention, funding and science has been devoted to cutaneous lymphoma research than ever before. The Foundation has been a strong partner to researchers who are a driving force behind this change, and we are working to increase the number of dollars, scientists and research projects devoted to cutaneous lymphoma.

The Foundation has funded important epidemiology and quality of life studies, and it provides funding to early investigators through its ongoing Young Investigator Awards. In addition, the Foundation is developing a Research Awards Program that would provide funding to investigators interested in conducting innovative cutaneous lymphoma research designed to improve treatments, clinical care and quality of life for patients.

Advocacy

At the Cutaneous Lymphoma Foundation, we advocate on behalf of people affected by cutaneous lymphomas to drive positive change on many issues that are important to a patient's overall care and quality of life. These issues include increasing funding of research, ensuring access to high quality medical care and medication reimbursement.

For more information about the Cutaneous Lymphoma Foundation and all of its programs and services, visit www.clfoundation.org.

2010-2011 Highlights

During the past fiscal year (July 1, 2010 – June 30, 2011), the Cutaneous Lymphoma Foundation accomplished much progress on many fronts including:

- Thousands of patients, loved ones and others served through live or electronic educational opportunities including Patient Educational Forums, cutaneous T-cell lymphoma sessions held at workshops organized by the Lymphoma Research Foundation and The Leukemia & Lymphoma Society, and teleconferences.
- Re-launch of our website at www.clfoundation.org, which went live in late 2011.
- Needs assessment conducted providing feedback about programs that are most important to cutaneous lymphoma patients.
- New *Patient's Guide to Understanding Cutaneous Lymphoma* begun with publication date projected in early 2012. This comprehensive educational resource will include helpful information about understanding the disease, treatments available, tips for managing symptoms and side effects, finding resources and support, and much more.
- Thousands of patients and others served through publication of our printed and electronic newsletters.
- Hundreds of patients, caregivers, medical professionals and others served through personal assistance via phone, email and literature mailings.
- Recommendations were initiated for a new Research Awards Program, which would provide funding to investigators interested in conducting innovative cutaneous lymphoma research designed to improve treatments, clinical care and quality of life for patients.
- Board of Directors expanded with six new members added in 2011.

For additional and updated information on all of the Foundation's ongoing work helping patients, visit www.clfoundation.org.

Financials

Statement of Financial Position • Year Ending June 30

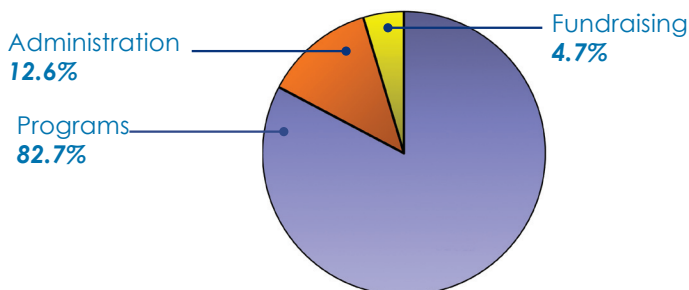
Assets	2011	2010
Current Assets		
Cash and cash equivalents	\$374,112	\$268,174
Money market accounts	\$882,241	\$842,731
Investments	\$5,973	-
Promise to give	\$35,775	\$74,500
Inventory	\$3,802	\$2,266
Total Current Assets	\$1,301,903	\$1,187,671
Liabilities and Net Assets		
Current Liabilities		
Accounts payable	\$19,630	\$4,211
Total liabilities	\$19,630	\$4,211
Net Assets		
Unrestricted net assets	\$660,749	\$591,647
Temporarily restricted net assets	\$621,524	\$591,813
Total Net Assets	\$1,282,273	\$1,183,460
Total liabilities and net assets	\$1,301,903	\$1,187,671

Statement of Activities • Year Ended June 30, 2011

Revenue	Unrestricted	Temporarily Restricted	Total
Contributions and grants	\$400,928	\$271,023	\$671,951
Summit proceedings advertising	\$73,500	-	\$73,500
Merchant sales	\$440	-	\$440
Investment income	\$1,342	-	\$1,342
Total support and revenue	\$476,210	\$271,023	\$747,233
Net assets released from temporary restrictions	\$241,312	(241,312)	-
	\$717,522	\$29,711	\$747,233
Expenses			
Program	\$458,013	-	\$458,013
Management and general	\$69,914	-	\$69,914
Fundraising	\$25,775	-	\$25,775
Total functional expenses	\$553,702	-	\$553,702
Change in Net Assets			
Net assets beginning of the year	\$163,820	\$29,711	\$193,531
Prior period adjustment	\$591,647	\$591,813	\$1,183,460
	(94,718)	-	(94,718)
Net assets beginning of the year, as restated	\$496,929	\$591,813	\$1,088,742
Net assets - end of year	\$660,749	\$621,524	\$1,282,273

Expense Ratio

The Foundation's programs expense ratio in fiscal year 2011 was approximately 83 percent. These funds were used to deliver vital programs and services including Patient Educational Forums, our website, teleconferences, educational newsletters and e-newsletters, personal assistance, advocacy and research efforts, and much more.



Donors

The Cutaneous Lymphoma Foundation is deeply grateful to all of the donors who supported our mission through financial contributions during our 2010-2011 fiscal year (July 1, 2010 – June 30, 2011). Thank you so much for your support. Your gifts have enabled us to make sure each person with cutaneous lymphoma gets the best care possible.

Recognition Society Donors

Beginning with our 2010-2011 fiscal year, we created new recognition societies in conjunction with our Annual Fund Campaign that provide leadership donors with benefits that include special recognition in our annual report. We are pleased to acknowledge these donors below.

Visionary of Care Society

\$10,000+ Annual Fund Campaign Giving

Ceptaris Therapeutics
Therakos

Partner in Care Society

\$5,000 - \$9,999 Annual Fund Campaign Giving

Ruth Aschkenasy
Suzanne Gylfe
Glenn Jaffe
Florence Seligman

Friend in Care Society

\$1,000 - \$4,999 Annual Fund Campaign Giving

Kathy Bromage
Claudia Day and Carl Prieser
David Feinbloom
Mike Johnson
Judy Jones
Dr. Stuart R. Lessin
Chris and Melody Malachowsky
Mary McCarthy
Deborah McNelly
David Rumph
Mark Rusley
Robert Shipp
John Steinbrunner
University Physicians, Inc.
John Webb

All Donors

The following listing includes all individual, corporate and foundation donors who contributed to our Annual Fund Campaign or supported other programs and services.

Ken Abrams	Margaret Baum	Jonathan and Amy Britz
Dr. Donald J. Adler	Alan and Rina Beach	Kathy Bromage
John Adler	Norbert Beauchemin	David Brown
Beverly Albert	Peggie Beaudoin	Dawn Brown
Allos Therapeutics	Judy Beck	Diane Brown
Dortha Allred	Carolyn Beebe	June Brown
Richard Allwes	Mary Ellen Behm	Keith Brown
Marla Almeida	Teri Behm	Ivan Budisin
L. Ambio	Carlene Belanger	Judith Burgstein
Dr. Thomas Anderson	Arnold Bell	Carolyn Burrous
Thomas Andrews	Avis Bent	William Button
Ari-Zev Anolic	Anne Benvenuto	Jeannine Byrne
Raymond Arel	Dorothy Berchtold	Ellen Cameron
Dr. Frank Arena	Natalie Berger	Chris Campos
Bob Arnold	Linda Bernstein	Elaine Caplan
Kathy Arnold	Debby Beveridge	Joe Carey
Howard I. Aronson	Frank Birney	Ray Carroll
Ruth Aschkenasy	William Blackburn	Anna Carullo
Susan Aucoin	Eric Block	Mary Carvell
Judith Axelrod	Andrew Bloom	Mary Casale
John Baier	Karen Blount	Warren Casner
Timothy and Pamela Bardy	Pastor Dan and Patricia	Phyllis Cavan
Gail Baril	Bodine	Celgene
Nicole Baril	Rev. James Bogardus, Jr.	Ceptaris Therapeutics
Judith Barker	Barbara Bolton-Budlong	Nancy Chandler
Sandra Barker	Richard Bradlow	The Earl M. and Margery C.
Beth Barnett	Allen Brandt	Chapman Foundation
Barr-Hirschlag	Sally Branson	Leslie Chappel
Geraldine and Robert Barton	Phil Braverman	Carol Charles
Richard and Laura Bass	Henry Brehm	Patricia Chason
Elizabeth Bassan	Renate Brenneke	Philip Church, Jr.
H.L. Bauer	Linda Brewer	Cleveland Museum of Art Docent Association

All Donors Continued

Patricia Clowdus	Marie deVegvar	Kate Ferreira
Anne Clune	Dr. Neil Dicker	Barbara Fizzano Uebler
Avery Cohen	Sally Dietz	Priscilla Flanz
Noal Cohen	Joseph DiSena	Alan L. Fletcher
Wallace M. Cohen	Richard Dobrin	Martin Forman
Kent Colberg	Barry Dootson	Larry France
Donna Cole	Donald Dorer	Samuel Frank
Denise Collins	Chery Downen	Stacy Frank
Paris Collins	Joan Dresner	Thomas Frank
Fred and Gail Colvin	Martha Drilling	Lowell Frederick
Jeanne Conner	Mark Drusin	Jackie and Bud Freedman
Ellen Connors	Steven Dubin	Melvin Freeman
Dr. Kevin D. Cooper	Robert Dubois	Tiffany Freiband
David Copelin	Ben Dussan	Joan Friedman
Cord Blood Registry	Patricia and Ronald Eamich	Rich Friedman
Steven Cordova	Frank Eastburn	Norma Friel
Julie Costa	Amy Edney	Dr. Ramsey Frist
Monica Coté	Guri Edoff	Frank Fritchman
Judi Cotter	Eisai Inc.	Steven Froberg
Stephen Cringle	Ann Eisenmenger	Marguerite Fumano
Michael B. Critchley	David Elefant	William Fuson
Helen Dagostino	Roland Elefant	Joseph and Ida Gabay
Dahab Associates, Inc.	Dave Elliott	Pat Gagne-Asiel
Faith Daiak	Ed Engel	Cathy Galloway-Davis
Deb D'Angelo	Kathy Erickson	Mary Gary
Denise and Scott Davis	Catherine Eskenazi	Fred Geer
Roger Davis	Jeanette Estremera	Rosalie Gershon
Walter Davis	Bill Evans	Dr. Francesca Giancotti
Charles Day	Robin Everett	Edward Gilbert
Thomas Day	Dr. James E. Faris	Linda Gilboord
Laurie Dean	David Feinbloom	Leann Gogel
Ron Decker	Sharon Felcoski	Frederick Goldberger
Donald Derda	Fellowship Place	Alice and Donald Goldsmith
Patricia Deuser	Susan Ferguson	Deb Goll

All Donors Continued

Susan Gombos	Bernice Henry	Mike Johnson
Dan Goodman	Karen Hepting	Pattie Johnson
Betty Gordon	Roseann Herman	Brian Jones
Elaine Gordon	Bruce and Linda Hermes	Helen Jones
Amy Gorman	Carol Herzlich	Judy Jones
Eda Gowdy	Maureen Higgins	Thomas Jones
Brian Gracey	Thomas Higgins	Rita Jungk
Dr. Lisa Grandinetti	Robert Hildebrandt	Ghassan Kabbara
Judy Grant	Margaret Hine	Deborah Karl
Scott Grant	Judith J. Hobensack	Ellen Katz Trust
Harrison Green-Fishback	Gretchen Hofer	Howard Katz
Donald Greenhall	Zita and Mel Holden	Irving Katz
Jennifer Greer-Glanville	Ann Hollis	Rona Katz
Roger Grekin	Barbara Hollis	Herbert Keating
Malvin N. Groce	John Holter	Terrence Keefe
Shirley Grossman	Helen Homulka	Allan Keiller
Suzanne M. Gylfe	Thomas Homulka	Sara Kellerman
Phyllis Haas	Diane Hooper	Fred Kiesner
Deberah Haferkamp	Marcia Houchens	Brad and Amanda Kik
Christine Halder	Susan Howell	Gertraude King
Jeanne F. Hall	Carol Howells	Esther Kirkpatrick
Dorothy L. Hammond	Mary Humphrey	Melvin Klaff
Roberta Hamovitz	Dr. Sam Hwang	Sharon Kleinberg
Emogene Hanvey	Mirry Hwang	Marlene Klotz
Arlen and Lorraine Harbaugh	David Ingber	Dr. Elizabeth H. Knobler
Georgann Harbert	David Jackman	Kenneth Kobarg
Mary Harkins	Brian Jackson	Deborah S. Koch
John Harmon	Joy Jacobs	Benjamin Kohler
John Harrell, Jr.	Glenn Jaffe	Richard Kolster
Ray Hassard	Gerald Jaffee	Nabil Korban
Kathy Hausman	Tina Janichek	KPMG LLP
HBO, Inc.	Jacquelyn Jensen	Brace Krag
William Heffernan	Seta Jessourian	Elaine Krasnick
Sharon Heiman	Carolyn Johnson	Samuel Krimm

All Donors Continued

Ronald Krolak	Bari Lissner	Carl McDowell
Kenneth Kulpa	Sandra Littman	Patricia McFillin
Peg Kunz	Roger Lloyd	Dr. Chauncey A. McHargue
Mark Kushnir	Susan Loesser	Charles J. McIntosh
Dorothy Kuzneski	Carol Loewy	Jean McIntosh
Nancy Kvapil	Carmel Lowenthal	Joyce McKee
L.E.K. Consulting LLC	Leora Lowenthal	Kim McKenzie-Campbell
Richard LaFrate	Reinhard Ludin	Martha McLaughlin
Amelia Laing	Rosario Lugay	Diane McManis
David S. Lamb	Jeff MacFarland	Scott McNeil
Stanford Lamberg	The Mack Foundation	Deborah McNelly
Gregory Latino	Chris and Melody	Kathleen Meagher
Dolores Lawrence	Malachowsky	Susanne Mease
Anthony Lawson	Cordella Malloy	Jan Medved
Randy Lawson	Roberta Manchester	Kathy Meier
Alan Lazarus	Juliann Mancino	Peter Mercier III
Elizabeth Lazenby	Sid Mandelberger	Merck Oncology
Giselle Leary	Fred Mangione	Merck Partnership For Giving
Tanya Lebovici	John Mangusi	Barbara Merwitzer
Dr. Chao Te Lee	Allan and Audrey Marrus	Jack Meschino
William Lee	Linda Marshall-Smith	Loretta Miccariello
Kenneth Leeds	Noreen Martin	Walter Milchanoski
Christine Lenart-Mireau	Rainer Martin	Lauren Miller
Albert Lenny	Harris Maslansky	Mark Miller
Lenox Hill Hospital	Carolyn Masters	Celia Millington-Wyckoff
Leroy Jefferson Williams, Jr. Foundation	Ann Mary Masterson	Bruce Millrood
Dr. Stuart R. Lessin	Theresa Matsey	Jennifer J. Mills
The Leukemia & Lymphoma Society	Margie Matthews	Marlene Miner
Carly Leung	Katherine P. Matthews	Bruce F. and Barbara Minkoff
Trudi Lewkowitz	Helen Mattox	Ronald Misccko
Dr. Henry Lim	Charles Matuszak	Dorothy Mize
Dolores Lipiro	Barbara McCahill	Margaret Montgomery
	Mary McCarthy	
	Sharon McCarron	

All Donors Continued

Anne Moore	Kathi Palacios	LLC
Darryl Moore	Erica Pallaron	Daniel Ray
Rosemary Moore	William Panella	Johnnie Ray
Richard Morris	Dr. Dennis Parenti	Carol Reigle
Paul Morrison	John Parham	Elaine Reinbold Pitkin
Sydelle Morrison	Suchin Parihar	Georganna Reinbold
Sam Moschella	James Parkman	Kimberly Reinbold
William D. Mosher	Gabriel and Margaret Passero	Martha A. Reinbold
Mary Mourkas	Douglas Payne	Howard Reiss
Mary Muldoon	Jo Pembroke	Joy Reliford
Paul Muldoon	Marilyn Perlotto	Megan Reynolds
Jane Mullen	Tracy Perry	Mike Reynolds
Gary Muzzelo	Douglas Peters	Roger F. Reynolds
Michael Myers	Wanda Peters	Deb Richards
Ronald Nappi	Shirley Petrie	Betty Richmond
National Association of Insurance Commissioners	Sherry Pickett	Dr. Gerald M. Rittenberg
Gregory Nemetz	Paul Pietri	Richard Rivard
Dr. Aharon Netzer	Mark Pipal	Rita Roach
Jim and Judie Neville	Paula Pirrotta	Thelma Roach
Barbara Newton-Holmes	Pete Pistell	Robert Roberts
Mary Ney	John Plank	William Roberts
Jonathan and Chris	Gary R. Posner	Joshua Robin
Nieuwsma	Lynn Price	William Robins
Paula Ninerell	Holly Priebe	Nancy Rocamora
Northwestern University	Eleanor Prieser	Richard Rocamora
Iris Noya	Una Provenzano	David Rogoff
Paul Nunez	The Public Health Laboratory Staff	Gary Rose
Tim and Nita Oesch	Seymour Putterman	Richard Rose
Kathleen O'Keefe	Myrna W. Pyke	Maria Rossi
Dr. Elise Olsen	Nancy and Sheldon Rachman	Mary Claire Rossi
Barbara Ostberg	Carolyn Ransom	Gordon Roston
William Owens	Walter Rapchinski	The Rotary Club of Rocky Point
Louis Paillex	RAS Management Advisors,	John Roth

All Donors Continued

Louise C. Roth	Laurie Scott	Libby Sorkin Routman
Pat Roth	Robert Scott	Bill Souveroff
Robert and Eileen Roth	Lorraine Sebastiao	Paul Spann
David Rothman	Mary Anne Sedney	Lloyd Sparks
David Rumph	Linda Seibert	Tim Spencer
Kaye Ruskey	Florence Seligman	Linda Spinks
Mark Rusley	Jerry and Darlene Shaffer	Michael Spivak
Bill Rusu	Alan Shanoff	Dr. Robert Steele
Frederick Ruvkun	Lawrence Shapiro	Jeffrey Stein
Carol Ryan	Judith Shea	John Steinbrunner
Patrick Ryan	Gratia Sheppard	Tracy Stetson-Zejer
Linda Sage	Jianfeng Shi	Anne B. Stevenson
Marc Saleh	Christopher Shipp	Cathy Stewart
William Jack Salmon, Jr.	Robert Shipp	Richard Stieglitz
Gilda Sampson	Cynthia Shortall	Ann Stinson
Sherri Sampson	Wes Shouse	Richard Storch
George H. Sams	Amanda Shurgin	Howard Strauss
Judith Sapol	Sharon Siegel	Ted and Anne Suess
James Sater	Silvia Sikora	William Sullivan
Alvin Sauer	Deborah Simmons	Paul Sutton
Irving Scharff	Joyce Simmons	Katrina Swan
Gerald Scheffler	Yvette Simmons	Sandor Szombathy
Ernie Schell	Joanne M. Simonson	Barry and Helen Szoszun
Amy Schlesinger	Mary Singleton	Dr. John Szydelko
Beatrice Schlesinger	Ginger and David Smerling	Chris Talbott
Caryn Schlesinger	Gregory Smith	Ted Tarbet
Sandy Scholnick	Joyce Smith	Julia Tartt
Philomena Schorza	Steve Snider	John Tauzell
Teresa Schruben	Sidney Socolar	Marianne Tawa
Marie Audrey Schultz	Connie Soeder	James Taylor
Barbara Schwartz	Robert Solimano	Karl Tepfer
Ellen Schwartz Siegel	Caren Solomon Bharwani	Ian Tepielow
Robert Schwartz	Catherine Songster	Robert Tetreault
Herman Schwartzman	Sons of the American Legion	John Thayer

All Donors Continued

Ellen Theg	Bridget Wackerly	Jeanette Wind
Therakos	Kylie Wade	Richard Wingate
Bill Thiel	Terry Walker	Donna Workman
Andrew Thomas	Joyce Walkwitz	Bette Wozobski
Ray Thomas	Dr. Roberta Walsh	Paul Wrezel
Nikki Thomason	Charles Walter	Allen Wyke
Susan Thornton	Ralph Warren	Michael Yang
Gail Tietz	Sue Warren	Stanley Youngs
Barbara Titcher	Irwin Warsaw	Alice Yu
Karen Toepper	Alice Watson	Leslie Zarnowski
Brian Tomlinson	Susan Watson	Joni and Michael Zavis
Ruth Townley	Alice Watts	William Zeisset
Carmen Tozzo	Georgiana Weaver	Dr. John A. Zic
Wanda Trapp	John Webb	Andrea Zigler
Mike and Mary Treworgy	Richard Webman	Lewis Ziman
Nick Tucci	Michael Weinberg	Donald Zittel
Norma A. Turner	Rochelle Weinberg	
Laura Lee Underwood	Leila Weinburg	
University Physicians, Inc.	Borrah Weiner	
Jim Van der Kloot	Jeremy Weinstein	
Joseph Vandenberg	Carl F. Weiss	
Brian Vaughn	Ann T. Welch	
Tom Vaughn	David Westermoe	
Laura Veazey	Colette Wharton	
Artemio Ventura-Gomez	Barbara Whistler	
Jennifer Viano	Edward Whitman	
Joan Vlahakos	Kacie Wiersma	
J. Richard and Maryann Volpe	Linda Wilford	
	Annette Williams	
Dr. Eric C. Vonderheid	Janelle Williams	

We gratefully acknowledge our generous donors and please accept our apologies if we have made any errors or omissions with your name or gift. If your listing requires correction, please contact us at info@clfoundation.org or 248.644.9014. We will correct our records immediately if necessary.

Board Of Directors

Officers

Stuart R. Lessin, M.D.
President

Marianne Tawa, RN, MSN, ANP
Vice President

Jeff Ward
Treasurer

Laurel Carlson
Secretary

Directors

Brace Krag

David Lamb

Pierluigi Porcu, M.D.

Christopher Shipp

Michael Young

Jennifer A. Viano

Staff

Holly Priebe
Director of Operations and Finance

Jennifer A. Viano
Chief Executive Officer

Cutaneous Lymphoma Foundation

PO Box 374
Birmingham, MI 48012
Phone: 248.644.9014
Fax: 248.644.9014
Email: info@clfoundation.org
Web: www.clfoundation.org