Our Vision
A World without WM (Waldenstrom's macroglobulinemia).

Our Mission
Support and educate everyone affected by Waldenstrom's macroglobulinemia (WM) while advancing the search for a cure.

Our Values

- **Community**
  We believe in engaging the WM community by sharing information, providing support, and working together to eradicate the disease.

- **Integrity**
  We care about the WM community and commit to the highest financial and ethical standards, with transparency in all that we do.

- **Teamwork**
  We work as a team that includes all WM stakeholders.

- **Focus**
  We are fiercely dedicated to providing support and education for those affected by WM and supporting research to improve and extend lives and find a cure for the disease.

- **Collaboration**
  We leverage the collective passion and genius of other blood cancer organizations and the medical and research communities.

- **Accountability**
  We are transparent and accountable at every level to the WM community and the general public.
Dear friends of the IWMF,

The global COVID-19 pandemic made 2020 an incredibly challenging and frightening year for Waldenstrom’s Macroglobulinemia (WM) patients and their families. However, due to the hard work, perseverance, ingenuity, and generosity of our staff, our volunteers, the IWMF Board, and the entire WM community, we moved closer to our goal of a “World without WM.” Notable accomplishments in 2020 included:

**Record revenue of over $4,000,000.** up 39% vs. 2019.
For perspective, total health care giving in the US was down 2% in 2020 vs. 2019, while medical research giving was down 11.9% per the Blackbaud Institute. A heart-felt thank you to all our donors for your very generous support!

**Expanding research** support by establishing the Dr. Robert Kyle Career Development Award to support promising young researchers in WM. Since 1999, the IWMF has funded over $20,000,000 in WM-specific research, which has led to better treatments with fewer side effects and a very significant increase in life expectancy for WMers. Since 2015, research has been driven by the IWMF-LLS Strategic Research Roadmap, which evaluates and funds the best research projects on a global basis.

**Pivoting to online support** to ensure that people with WM everywhere had easy access to the information and support they needed about our rare disease. On short notice, our staff moved the in-person Educational Forum planned for Seattle, WA to an easy-to-use online site. Attendance soared from our usual 350-400 in-person attendees to over 1,400 registrants from over 30 countries. The overall rating for the Educational Forum was 4.71 out of 5, equivalent to a grade of 94, proving the willingness of our largely older community to learn new tricks and, ensure they are never alone.
All of our global support groups switched to online Zoom groups. Again, attendance increased as we were able to reach people who couldn’t travel or didn’t live near an existing support group. And we also established a series of online training sessions to help our support group leaders do their very challenging jobs.

**Expanding our global network** by adding new affiliates in Israel, South Africa, Portugal, and Chile and adding new WM expert clinicians in Argentina, China, Greece, Israel, Poland, Russia, and South Africa. All our publications continue to be published in at least seven languages.

**Improving our educational materials** through a new, easier-to-use website launched in November. We also created five global educational webinars that reached over 3,000 people with WM worldwide.

**Shifting the IWMF Board to a strategic board** from an operating board. Consistent with the strategic plan adopted in 2019, the IWMF hired Newton Guerin as President and CEO in February 2020. Newton’s years of experience with the American Liver Foundation, Leukemia & Lymphoma Society, National Multiple Sclerosis Society, the American Cancer Society, and the International Dyslexia Association have provided tremendous value and greatly aided this critical transformation. Peter DeNardis, a long-time IWMF Board member, became chair of the IWMF Board of Trustees in December 2020, and four new Board members were added in the second half of 2020 to help the Board achieve its new goals.

While we are proud of what we were able to accomplish in 2020 despite the pandemic, we are not satisfied, and we never will be until we reach our goal of a “World without WM.” With your continued financial support and emotional support for each other, we will keep expanding and improving. Remember: **With the IWMF, you are never alone.**

Carl Harrington
Chair, IWMF Board of Trustees (2012 - 2020)
Ten years ago at the age of 75, I needed a knee replacement, the result of many years of hard court tennis. When the pathologist did a perfunctory biopsy on the offending knee after it had been removed just to insure that the surgeon had not done unnecessary surgery, a disease was discovered in the knee. Logic said that if the knee was diseased, so must be the body. The original diagnosis was lymphoma, and that was later refined to WM.

While I have kept pretty current about treatment options, I have been fortunate to not have any serious symptoms, so I have not considered any treatment options at this time. Thanks to the resources available through the IWMF, I'll be ready when the time comes.

I usually hesitate before I give too much advice to other people whether they are family, friends, or people that I don't know. But I'll tell you what I think: **Waldenstrom's is not a death sentence.** A large number of those afflicted will live long enough to die of something else rather than WM. It is, for many, a situation that we merely need to be aware of, and for others an affliction that needs to be managed. For some, there are life modifications that need to be made, although I am one who has not needed to do that. Fortunately, there are very worthwhile treatment options that can be utilized when necessary. Outstanding research is being conducted, and eventually there will be a cure; In the meantime there will be better and better treatment available. Optimism is important when it comes to Waldenstrom's, as it is in most aspects of life.

Waldenstrom's has changed me and my life in one important way: I get more out of every day than ever before. I have been diagnosed with Waldenstrom's, a serious and potentially fatal disease. As a result, I have a realization that I need to redouble my efforts to get more out of every day, to enjoy every moment, and to tell those that I care about how much they mean to me.
After ten years since my original diagnosis, I have lived some of the best years of my life, and, to a great extent, I have Waldenstrom's to thank for that.

My wife, Joanne, and I are very active. We play golf three times a week, and I look forward to doing that for as long as possible. We love to travel. I look forward to getting back to exploring the world. I love time spent with my kids and grandkids, and when their busy schedules permit, I get great joy from that. I have always been involved with some charitable endeavors. I do not want to be so presumptuous as to tell others what to do, but, for me, I would say enjoying life to the fullest is a good prescription for happiness. WM has given me an added incentive for that.
My dad was diagnosed with Waldenstrom’s macroglobulinemia in 2011.

It’s going to be a familiar story to anyone reading this, no doubt, united as we all are by our experience with loved ones suffering from this illness, or by our own affliction. We only found out because he was hospitalized for something entirely unrelated.

My father has always been athletic and a sports lover; he was an avid tennis player throughout most of his life. But the human body can only take so much, and every single step, however considered, was a blow to his knees. By 2011, the damage was so great that his tennis days were over, and he needed a complete knee replacement. But once his surgeons got in there, they discovered something terribly wrong with his blood. It took something as dramatic as wrecking his knee to discover the indolent illness spreading throughout his blood by degrees.

With his diagnosis of Waldenstrom’s came a five-year prognosis, and we prepared for the worst. I’m forever thankful that the worst never came, and he’s long since outlived his initial prognosis—a feat that that many have achieved thanks to the incredible strides in research made possible by the IWMF. Just as much as the research the IWMF supports, I’m grateful for the support I’ve found here, and the support I’ve in turn been able to offer.

If there’s anything I’ve learned from my dad over the course of my life, it’s perspective. From childhood lessons on independence, self-reliance, and positivity to his lifelong commitment to fitness and health, to his chin-up response to his diagnosis, he’s always had an attitude that, while easily mistaken for “always look on the bright side of life,” is much richer than that. It’s an attitude of what I’ll call “optimistic resignation.” Your fate, in short, is what you make of the circumstances you’re given. Wealthy or poor, healthy or sick, your life is ultimately your own.
At the end of the day, we are the captains of our fate; we live and we die by our own decisions, and the mark we leave on the world is ours to choose. It’s why I, after learning of the IWMF in early 2017 (a full six years after my father’s diagnosis, and a year after his life was supposed to be over) began donating $500,000 a year to its work, and why I continue to do so. The money given through the Elizabeth Elting Foundation, and through our family, is a fraction of what my father has given me in my life. It was the upbringing my father offered that made my success possible. I’m proud to be able to support the IWMF’s work. The goal isn’t just philanthropy as an abstract good; it’s to participate meaningfully in something that affects not only my father, but countless others.

What I hope to do, more than anything, is to help people see that WM isn’t a death sentence. That has been the greatest lesson my dad has taught me these last few years. You can look at it as a sword dangling over your head or as a daily reminder to live your life on your own terms. By supporting research that helps doctors identify and treat WM, we—you and I, together—can make those years as meaningful as possible, allowing those with WM to live their lives on their own terms. Remember that we only get one go-round in this world and to spend it living, loving, and doing good. The IWMF makes it possible for so many to live their own lives, not under the sword of WM, but in the liberating light of life.
Dear friends of the IWMF,

In December 2020, I assumed the role of Chair of the IWMF Board of Trustees. This position has a long history of volunteer patients who have each, in their own way, successfully propelled the IWMF further in its mission to “support and education everyone affected by Waldenstrom’s macroglobulinemia (WM) while advancing the search for a cure.”

Naturally, first and foremost, my objective is to do justice to those who came before me, and to do my best to follow in their footsteps.

Looking ahead, the IWMF will continue to sharpen its focus on education and research by pursuing such innovative strategies as:

Create a global collaboration for WM with other international cancer organizations. The intent is to ensure that, no matter where one happens to end up when searching for digital information about WM, the IWMF will have already developed and provided the background material for that information. As a result, everyone wins – patients and caregivers are provided with the most accurate, up-to-date details regarding diagnosis of, treatment for, and living with WM.

Focus on providing opportunities for young investigators new to the field of WM so that they can expand their knowledge and understanding of our disease and, in turn, become WM experts who can propel the next generation of treatments for us.

Provide mentorship/fellowship opportunities to clinicians and researchers so that they can become more conversant about WM and establish stronger relationships with current WM experts around the world.
Further expand medical research activities around the world by providing opportunities and incentives to researchers to work in the WM disease space.

Enhance educational content for the global WM community, ensuring that the information is updated based on the latest research findings and treatment recommendations, and capitalizing on the newest mechanisms available for disseminating that information.

Engage with the global WM community by seeking out new ways to interact and by taking advantage of social media tools and technology.

Elevate the visibility of the IWMF so that any person around the world that is newly diagnosed will automatically get connected with the informational, educational, and support services we provide.

Reach out to currently underserved communities of WM patients and caregivers by expanding the IWMF affiliate and support group network (if you’re interested in setting up an affiliate and/or support group in your geographic area, please contact the IWMF home office).

The IWMF is a patient/caregiver-led organization, with a relatively small staff of highly talented and committed individuals. Still, we can’t accomplish all our objectives nor provide the most comprehensive services and support to the global community of WM patients and caregivers without financial support from that community, and from pharmaceutical sponsors. Towards the objectives of further expanding WM-specific research activities, and of taking educational and support services to the next level, we are embarking on an ambitious fundraising effort over the next five years. Stay tuned for some exciting new giving opportunities where everyone can participate (no matter how large or small the donation) and engage with people with WM around the world.

Best of health to all!

Peter DeNardis
Chair, IWMF Board of Trustees (2020 - Present)
In fiscal year 2020, IWMF raised $4.080 million to support our mission and operations. We spent $2.263 million or 87.15% of our expenses on specific programs designed to find new treatments for WM and improve the lives of patients and their families.

The IWMF has achieved a Charity Navigator overall score of 95.47 on a scale 100, and a 100 rating on accountability and transparency. This is the IWMF’s fourth consecutive 4-star rating from Charity Navigator. Only 21% of the organizations reviewed by Charity Navigator have received this ranking four years in a row. This outside confirmation of what the IWMF has done should be a strong signal that donating to the IWMF is a good investment.

**Consolidated Statement of Activities for the year ended December 31, 2020 (US dollars)**

<table>
<thead>
<tr>
<th>Revenue:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Campaign Contributions</td>
<td>$ 3,989,770</td>
</tr>
<tr>
<td>Other Income</td>
<td>$ 82,850</td>
</tr>
<tr>
<td>Net interest and dividend Income</td>
<td>$ 7,809</td>
</tr>
<tr>
<td><strong>Total Revenue:</strong></td>
<td><strong>$ 4,080,429</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Expenses:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Research Grants</td>
<td>$ 1,457,166</td>
</tr>
<tr>
<td>Information, Education &amp; Support</td>
<td>$ 805,623</td>
</tr>
<tr>
<td><strong>Total Program Services</strong></td>
<td><strong>$ 2,262,789</strong></td>
</tr>
</tbody>
</table>

Management and General                         | $ 163,279 |
Fundraising                                     | $ 170,306 |
**Total Expenses:**                             | **$ 2,596,374** |

**Change in Net Assets**                        | **$ 1,484,055**

**The IWMF Board of Directors have approved multi-year agreements for research grants totaling $1,950,815. These grant payments are conditional, and are not included as liabilities on this consolidated balance sheet.**

**Consolidated Balance Sheet as of December 31, 2020**

<table>
<thead>
<tr>
<th>Assets:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Cash and cash equivalents</td>
<td>$ 4,606,674</td>
</tr>
<tr>
<td>Prepaid Expenses and other assets</td>
<td>$ 73,220</td>
</tr>
<tr>
<td>Contributions and other receivables, net</td>
<td>$ 547,508</td>
</tr>
<tr>
<td>Fixed Assets, net</td>
<td>$ 90,530</td>
</tr>
<tr>
<td><strong>Total Assets:</strong></td>
<td><strong>$ 5,317,932</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Liabilities:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Accounts payable &amp; accrued Expenses</td>
<td>$ 77,171</td>
</tr>
<tr>
<td><strong>Total liabilities</strong></td>
<td><strong>$ 77,171</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Net assets:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Without donor restriction</td>
<td>$ 2,410,402</td>
</tr>
<tr>
<td>With donor restriction</td>
<td>$ 2,830,359</td>
</tr>
<tr>
<td><strong>Total net assets</strong></td>
<td><strong>$ 5,240,761</strong></td>
</tr>
<tr>
<td><strong>total liabilities and net assets</strong></td>
<td><strong>$ 5,317,932</strong></td>
</tr>
</tbody>
</table>
BEN RUDE HERITAGE SOCIETY

The Ben Rude Heritage Society recognizes those who have made provisions for a future gift to the IWMF, such as a bequest, listing the IWMF as a beneficiary for a life insurance policy or qualified planned asset (such as a 401k or IRA), or a life income agreement, such as a Charitable Remainder Trust. Legacy gifts represent an important component of the IWMF’s financial future. There are many ways to support the IWMF through a planned gift, but a bequest is perhaps the easiest and most tangible way to leave a lasting impact. The following supporters are members of the Ben Rude Heritage Society:

ANONYMOUS (4)
JEAN-MARC AUDIBERT*
MARTIN BAER*
C. EDWIN BAKER*
JACK BAKER
JANET BAUSser*
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ROGER* AND BARBARA ROBINETE
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LAURIE RUDE-BETTS ◊
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RAY SOBOROWICZ*
KEN AND LINDA SOLOW
CORDELIA AND PETER STEARNS
JUDITH LEE STERLING AND
LESLIE CHARLES WILSON
JOHN AND DIANE TIPLADY
RUBEN TROSS◊
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TAMARA VINCENT
DR. JACOB WEINTRAUB
VIRGINIA A. WILLIAMS*
LISA WISE
PENNI WISNER
JAMES P. YEAGER*
GREGORY FITZWATER AND
M ARILYN ZOLLNER-FITZWATER
RALPH ZUCKERMAN*

* Deceased
◊ Founding Member
For a commitment of $50,000 per year for a minimum of two years, or a lump sum of $100,000 or more, you can become a research partner supporting a specific IWMF research project approved by the IWMF's Scientific Advisory and Research Committees. Research Partners will have an opportunity to be kept informed of the progress of the research project and will be formally acknowledged by the investigators in their report of the project as well as in any resulting publications. Generally 10 to 12 research projects are underway with new projects under consideration each year. The following funds support current IWMF research:

- The David and Janet Bingham Research Fund of the IWMF
- The Elting Family Research Fund of the IWMF
- The Lynn M. Fischer Research Fund of the IWMF
- The K. Edward Jacobi Research Fund of the IWMF
- The Carolyn Morris Research Fund of the IWMF
- The Ed and Toni Saboe Research Fund of the IWMF
- The Paul and Ronnie Siegel Family Research Fund of the IWMF
- The Robert and Nadeline White Family Research Fund of the IWMF
- The Marcia Wierda Memorial Fund of the IWMF
- The Yang Family Research Fund of the IWMF
NAMED GIFT FUNDS

For a commitment of $10,000 per year for five years, or a lump sum of $50,000 or more, you can establish a named fund at the IWMF in your own name or in the name of someone you wish to honor. The following funds support information, education, mission programs, research, or a combination of each:

**Baker Family**  
Research Fund of the IWMF

**Yoshiko Button**  
Mission Support Fund of the IWMF

**Friedlander-Scherer Family**  
Research Fund of the IWMF

**Gary Green**  
Research Fund of the IWMF

**Dr. Robert A. Kyle and Charlene M. Kyle**  
IWMF Career Development Research Fund

**Lynn Martin and Carrie Wells**  
Research Fund of the IWMF

**Dennis and Gail Mathisen**  
Research Fund of the IWMF

**Gail Murdough**  
Mission Support and Research Fund of the IWMF

**Sesnowitz Family**  
Research Fund of the IWMF

**Donald and Alison Weiss and Family**  
Research Fund of the IWMF

**Donald and Kathryn Wolgemuth**  
Research Fund of the IWMF

**Joseph and Maureen L. Janda**  
Research Fund of the IWMF

If you have discretionary giving power and would like to help move our research program forward in a special way, we invite you to join those listed above. For more information about Research Partners and Named Gift Fund opportunities and potential gifting options that might make that possible, please contact Director of Development and Communications Jeremy Dictor at JDictor@iwmf.com or 941-927-4963.
OUR SUPPORTERS

The work of the IWMF would not be possible without the generosity and support of people like you. On behalf of the entire WM community, we offer our sincerest thanks for your support in 2020:

$1,000,000+

The Lynn M. Fischer Research Fund of the IWMF

$500,000 - $999,999

Elizabeth Elting

$100,000 - $499,999

BeiGene, Ltd.
Catherine Howell and Michael D. Ernst
Janssen Oncology
Pharmacyclics

$50,000 - $99,999

Maynard Morris
Howard Prass
Edward and Toni Saboe
The Patterson Foundation
Robert White
Kenneth Wierda

$25,000 - $49,999

Gina Maya and Richard Capelouto
Stanley Kaufman
Leukaemia Foundation - South Australia
Waldenstrom's Macroglobulinemia Foundation of Canada
Ralph Zuckerman
$10,000 - $24,999

Annette and Elias Aburdene
Frederick and Almie Baker
Dr. H. David and Lois Baru
John Ciesielka and Dale Bottoms
Dorothy Bryan
Anita and Bruce Buyer
Cellectar Biosciences, Inc.
Gia Colosi
Marilyn Friedlander
Dieter and Margarita Haussmann
Timothy and Sidney Hoesch
Josephine Hughes Sterling Foundation
Roy Langhans
Jayne McMellen
Gail and Sam Murdough
Dr. and Mrs. Stephen and Karen J. Schange
Meryl and Robert Selig
Michael and Carol Sesnowitz
Daniel and Robin Stern
The Treadway Foundation
Weingart Foundation
Donald J. Weiss
X4 Pharmaceuticals
Willard R. Young III
Zivin Family Foundation

$5,000 - $9,999

Anonymous
Michael and Laurie Berg
Bernard A. Egan Foundation, Inc.
Bernard F Graham Charitable Foundation
Laurie Betts
Paul Bodin
Ronald and Mary Jane Branscome
Ann F. Collins
Michael and Judith Crutcher
Joseph and Allyn Dunn
Cathy and Don Hutton
David and Penny B. Kirby Jr.
Murilyn Koutstaal
Ellen C. Kreiger Rubin
Robert Kyle MD
Robert and Janet LeGrand

Don and Sandra Livinghouse
Lynn and Carrie B. Martin
Brenda and Stuart McCroskey
Linda Nelson
Lynn and Joyce Overboe
Frances Owen
Michael and Dianna Poh
Harvey and Joyce Rose
Jane E. Rosen
Jeffrey Rubenstein
Kenneth and Linda Solow
Julia and Stanley Stasch
Jeffrey and Andrea M. Stewart
Waldenstrom France
Thomas and Deborah White
Laurence and Susan Wolfe
$1,000 - $4,999

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Sylvia Almstadt  
Robert Altomare  
Amazon Smile  
Harold and Colleen Ambauen  
Judith Aucoin  
Susan and Robert Ayres  
Gayle and John Backmeyer RN, NP  
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Laura and Richard Bass  
Thomas Battle  
Michael and Maggie Bayer  
John Bearss  
Lisa Beeby  
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Bryan Bier  
James C. Black  
Clinton and Susan Blandford  
William Block  
Diane Borreson  
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Nancy and Vince DiGirolamo  
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Hubert and Jacqueline Edfors  
Deirdre and Christopher Edgerton  
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Dennis and Judith Fiehler  
Marjorie Fields  
Fingerhut Family Foundation  
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William and Susan Friedlieb  
Brian and Louise Frumkin  
Eleanor Fuller  
John and Frances P. Gallagher  
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David and Catherine Grossman  
Newton Guerin  
George Guttman  
David Hamberg  
Donald Handal  
Carl Harrington  
Robert and Lynn Hawkey  
Kathryn and David T. Heiser  
Garith Helm  
Jane and Ralph Hendrickson  
Roxanne Hetzel-Bunger  
Mike Heymann  
Kevin and Kathleen Hogan  
Teresa and Wayne Holimon  
Mary Ann Houser  
William and Kristine Howanski  
John and Lee Howard
$1,000 - $4,999

Deh-i Hsiung
IWMF California
Madeleine L. Jackson
Dr. Dustin G. James MD
Jane C. MacElree Foundation
John and Ann P. Jenkinson
Cheryl Jones
Dan and Elizabeth Junius
William R. Kaye
Linda and Clyde Kenneaster
J. Kevin and Ann Kilbane
Jennifer and Daniel Killam
Kathryn and Luther Kissam
Dr. Lois Lenarz and Robert Kolasa
Warren Kozak
Mary Kresky
Paul M Kuveke Jr.
Thomas and Beverly Lacey
James A. Larkin
Rowland and Tobbie Leep
Frederick Leonard MD
LLS Leukemia & Lymphoma Society
Ann Lipkowitz
Carl Lisman
Joseph and Yvonne LoRe
Thomas and Alice Lovely
Raymond and Jo Lucas
Ralph Lunan
John Thomas and Jan Lund
Bonnie MacMaster Anderson
Elena Malunis
Dr. Barbara and John Manousso MD
R. Stephen Manty
Joseph Marcianite
Margaret P. Stevenson Foundation
Margaret Stevenson Foundation
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Ms. Antonella Marotti
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David E. Muth Jr.
Linda Nelson
Noble Drilling Services, Inc.
Barbara and Michael Noonberg
Nordson Corporation
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Financial and other information about The International Waldenstrom’s Macroglobulinemia Foundation, Inc. can be obtained by writing the Foundation at 6144 Clark Center Avenue, Sarasota, FL 34238. In addition, several states where The International Waldenstrom’s Macroglobulinemia Foundation, Inc. is required to file financial information each year also require the following disclosures: Colorado: Colorado residents may obtain copies of registration and financial documents from the office of the Secretary of State, (303) 894-2680, http://www.sos.state.co.us/. Florida: Registration No. CH33403. A COPY OF THE OFFICIAL REGISTRATION AND FINANCIAL INFORMATION MAY BE OBTAINED FROM THE DIVISION OF CONSUMER SERVICES BY CALLING TOLL-FREE WITHIN THE STATE 1-800-HELP-FLA OR VIA THE INTERNET AT http://www.FloridaConsumerHelp.com. Georgia: A full and fair description of the programs and activities of The International Waldenstrom’s Macroglobulinemia Foundation, Inc. and its financial statements are available upon request at the address indicated above. Maryland: For the cost of postage and copying, documents and information filed under the Maryland charitable solicitation law can be obtained from the Secretary of State, Charitable Division, State House, Annapolis, MD 21401, (800) 825-4510. Michigan: MICS No. 45029. Mississippi: The official registration and financial information of The International Waldenstrom’s Macroglobulinemia Foundation, Inc. may be obtained from the Mississippi Secretary of State’s Office by calling 1-888-236-6167. Registration with the Secretary of State does not imply endorsement by the Secretary of State. New Jersey: INFORMATION FILED WITH THE ATTORNEY GENERAL CONCERNING THIS CHARITABLE SOLICITATION AND THE PERCENTAGE OF CONTRIBUTIONS RECEIVED BY THE CHARITY DURING THE LAST REPORTING PERIOD THAT WERE DEDICATED TO THE CHARITABLE PURPOSE MAY BE OBTAINED FROM THE ATTORNEY GENERAL BY CALLING (973) 504-6215 AND IS AVAILABLE ON THE INTERNET AT www.njconsumeraffairs.gov/ocp.htm#charity. REGISTRATION WITH THE ATTORNEY GENERAL DOES NOT IMPLY ENDORSEMENT. New York: A copy of the latest annual report can be obtained from the organization or from the Office of the Attorney General by writing the Charities Bureau, 120 Broadway, New York, NY 10271. North Carolina: Financial information about this organization and a copy of its license are available from the State Solicitation Licensing Branch at 1-888-830-4989 (within North Carolina) or 919-807-2214 (outside of North Carolina). The license is not an endorsement by the State. Pennsylvania: The official registration and financial information of The International Waldenstrom’s Macroglobulinemia Foundation, Inc. may be obtained from the Pennsylvania Department of State by calling toll-free, within Pennsylvania, 1-800-732-0999. Registration does not imply endorsement. Virginia: Financial statements are available from the State Office of Consumer Affairs, P.O. Box 1163, Richmond, VA 23218. Washington: The notice of solicitation required by the Charitable Solicitation Act is on file with the Washington Secretary of State. Information relating to financial affairs of The International Waldenstrom’s Macroglobulinemia Foundation, Inc. is available from the Secretary of State, and the toll-free number for Washington residents is 1-800-332-4483. West Virginia: West Virginia residents may obtain a summary of the registration and financial documents from the Secretary of State, State Capitol, Charleston, WV 25305. REGISTRATION IN THE ABOVE STATES DOES NOT IMPLY ENDORSEMENT, APPROVAL OR RECOMMENDATION OF THE INTERNATIONAL WALDENSTROM’S MACROGLOBULINEMIA FOUNDATION, INC. BY THE STATE.
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