



ANNUAL REPORT

Taking the Journey Together

2024



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Letter from Leadership

Dear AAMDSIF Community,

Thank you for your generous financial support of the Aplastic Anemia and MDS International Foundation (AAMDSIF)! Your commitment and generosity help us to deliver on our mission for patients and families living with aplastic anemia, myelodysplastic syndrome (MDS), paroxysmal nocturnal hemoglobinuria (PNH), and related bone marrow failure diseases.

This annual report provides an update to our donors on how we effectively utilize our resources. The theme of 2024's awareness campaigns – **Taking the Journey Together** – reflects our commitment to the community. Our approach advocates, educates, and empowers patients and their loved ones throughout their journey. From diagnosis to survivorship, we offer patients and caregivers answers, support, and hope.

In 2024, the Foundation embarked on a strategic planning process that sought to affirm our strengths and identify opportunities in the work we do to advance our mission. With the input of a wide range of stakeholders, which included

patients, family members, and other caregivers, current and past board members, members of the healthcare provider community such as Medical Advisory Board members and community providers, other patient advocacy groups, donors, and staff, and guided by an external consultant with expertise in the non-profit industry, we have honed in on our strategic priorities and plans that will serve as our north star for the next five years.

AAMDSIF Strategic Plan Priorities

Over the next five years, we will:

- Serve aplastic anemia, MDS, and PNH communities, including related secondary diseases
- Focus on patients who live in areas without ready access to a major medical center, especially within the U.S.
- Serve individuals outside the U.S. who seek the Foundation's help and build our international scope and expertise thoughtfully and as funding allows
- Invest resources and use data to increase access to our programs and services

As a donor and investor in our work, you are a key part of the success of the Foundation. Through the Strategic Planning process, we also updated our Statement of Values, which informs our approach to carrying out our mission and includes:

- Commitment to serving the needs of aplastic anemia, MDS, and PNH patients and their network of support
- Respect for the worth and dignity of all individuals
- Dedication to inclusiveness and diversity
- Commitment to transparency, integrity, and honesty in all dealings
- Responsible stewardship of resources

We are grateful for your continued commitment to the Foundation. Your role as a donor is key to our success, and we hope you see that reflected in the pages of this annual report. Your annual donations and volunteering enable the Foundation to fulfill its mission. We couldn't do all that we do without you.



**Deborah "Debby"
Ziff Cook**
Chair, Board of Directors



Brad A. Wong
CEO & Executive Director

Taking the Journey Together

MISSION STATEMENT

The Aplastic Anemia & MDS International Foundation empowers individuals diagnosed with aplastic anemia, myelodysplastic syndrome (MDS), paroxysmal nocturnal hemoglobinuria (PNH) and their support network by providing timely information, a community of support, improved treatment through health provider education, and hope for a cure through promotion of research.

VISION STATEMENT

Patients, caregivers, and healthcare providers around the world have access to the resources they need so that everyone diagnosed with aplastic anemia, MDS, and PNH can thrive.

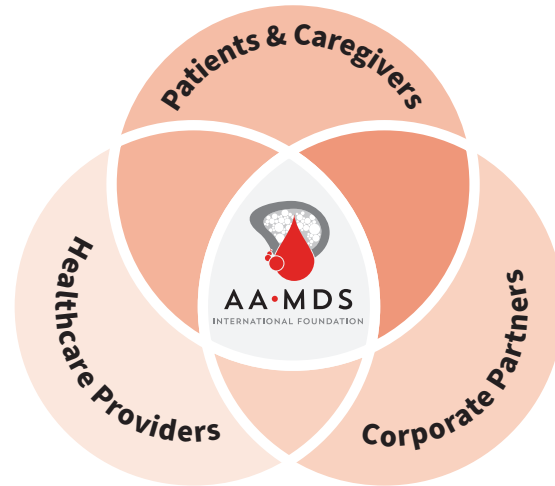
VALUES STATEMENT

In carrying out its mission, the Aplastic Anemia & MDS International Foundation and its Board members, staff and volunteers shall endeavor to act with the following values:

- Commitment to serving the needs of aplastic anemia, myelodysplastic syndrome (MDS), and paroxysmal nocturnal hemoglobinuria (PNH) patients and their network of support
- Respect for individuals
- Dedication to inclusiveness and diversity
- Commitment to transparency, integrity and honesty in all dealings
- Responsible stewardship of resources

At AAMDSIF, being patient-centered is more than a philosophy—it's our promise.

We are here for you at every step of your journey. Throughout the year, we provide trusted resources and fund vital research in rare bone marrow failure diseases to advance scientific knowledge. Our goal is to attract top talent in this important field and build a supportive community where you feel empowered. Being patient-centered means you're not alone. It means we're **taking the journey together**.



Healthcare professionals and corporate partners who share our mission are joining us on this journey. Whether you're a clinician seeking expert collaboration or a corporate partner aiming to better understand and support patients, AAMDSIF is here to connect you with the people, knowledge, and tools to make a meaningful impact.



Donna, living with MDS, and her granddaughter.



Jillian Stewart, living with PNH.

“AAMDSIF’s mission means that there are people and an organization dedicated to people like me who are afflicted with a rare blood disorder. It means connectivity, new information on treatments, and updates from patients and physicians—all critical to living your best life with this disease.”
— Patient living with PNH

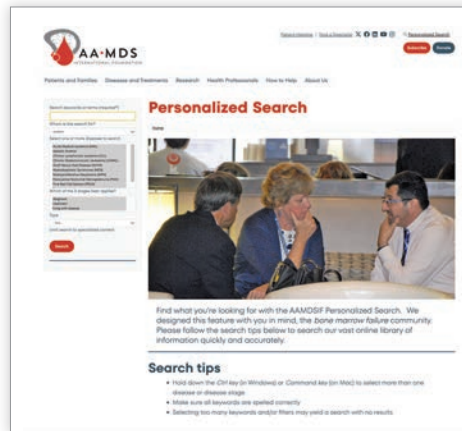
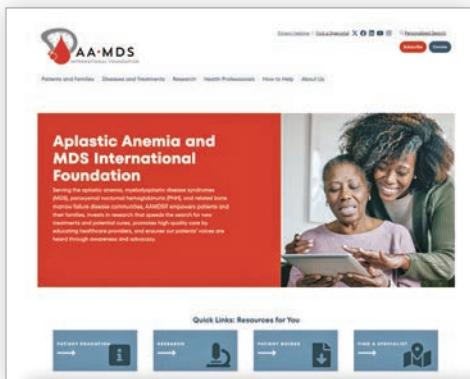
Same Organization, New Look

In 2024, we launched a newly redesigned website—built with you in mind.

We listened to our community and focused on a cleaner, more intuitive design that makes it easier than ever to find the information and support you need.

Guided by best-in-class design standards, user experience expertise, and your feedback, the new AAMDSIF website delivers quicker access to the trusted education, events, research, and resources you’ve come to expect from us.

One of the standout features of the new website is our all-new **Personalized Search tool**. This powerful function allows you to filter and narrow your search results based on your specific needs, putting you in control of your experience. Whether you're newly diagnosed, looking for clinical trial information, or seeking support groups, the search adapts to you—making it easier to locate the information that matters most.

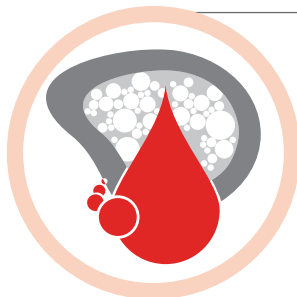


“I receive a wealth of information from the site. Having MDS it’s important to stay on top of new drugs, therapies and trials. I’m confident in the information I glean from this site.”

“My healthcare team referred me to the website as the #1 place for information about aplastic anemia. It is the gold standard for patients. It’s where I learned many things.”

This redesign is a part of our ongoing commitment to being truly patient-centered. It’s about more than just a website—it’s about creating a free, accessible space where our community feels supported, informed, and empowered.

This new digital experience is just one more way we’re **taking the journey together**—every step of the way.



Favorite Resources

- Immunosuppressive Therapy
- Iron Chelation
- Stories of Hope
- What are the symptoms of low blood counts?
- Nutrition

AAMDSIF is Powered by Your Support

GIVING TUESDAY

Every year, we're inspired by the generosity of our supporters on Giving Tuesday—but 2024 was truly exceptional. Thanks to a remarkable \$30,000 matching gift from two anonymous donors and the heartfelt contributions of 122 donors, AAMDSIF raised over \$68,000, setting a new record for the organization!

To learn more about how you can support the Foundation's fundraising efforts, please visit aamds.org/donate, where you can find links to our *Ways to Give* and *Fundraise* pages.

The Foundation's work is only possible with support from people like you and me, people who believe in their mission. I'm proud to be an AAMDSIF donor. I'm able to donate annually because my friends and family support me through a golf outing that I host called "Fighting PNH... Rare but Real."
— Brianne

At AAMDSIF, everything we do—from providing trusted information and emotional support to funding life-changing research—is made possible by the generosity of our community. Your donations support our mission to serve patients, families, and caregivers affected by bone marrow failure diseases, and healthcare professionals who treat them.

Opportunities for Impact

Our fundraising efforts take many forms: individual donations, monthly donations, donations from foundations, community-driven grassroots events, corporate partnerships, and planned giving. Every contribution has a meaningful impact.

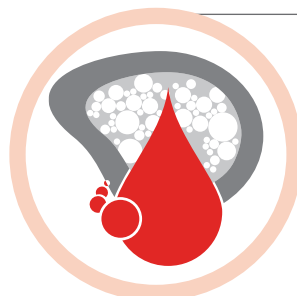
The Power of Grassroots Fundraising

Grassroots fundraising efforts demonstrate that every individual has the power to make a difference. Many supporters create personal fundraising pages to rally their communities, raise critical funds, and raise awareness of the challenges faced by those living with bone marrow failure diseases.

Community-based events are another example of meaningful community engagement. Take Brianne, a dedicated long-time supporter of AAMDSIF, who recently hosted her ninth annual fundraising event. Her continued commitment is a testament to the strength and heart of this community—and a reminder that when people come together, incredible things happen.



"Fighting PNH...Rare but Real" golf event chair Brianne Sandborn (seated third from the left) and event participants.



AAMDSIF relies on volunteers to deliver on our mission. To learn more about volunteer opportunities, please visit aamds.org/volunteer

Taking the Journey Together

2024 Impact



1,146

people registered
for AAMDSIF
support groups



121

support group
meetings



46

adult and pediatric
sessions held at our
Patient & Family
Conferences



4

Patient & Family
Conferences



94

trained Peer Support
Network volunteers



352

Patient & Family
Conference
registrants



521

HelpLine calls
answered



4,360

podcast
listeners
in 2024

Patient-Centered Programs

Providing Answers

Finding reliable answers is essential for patients and their support networks as they navigate life with a rare diagnosis. AAMDSIF is committed to **taking the journey** with you and providing you with high-quality resources to help you better understand your diagnosis and the associated challenges.

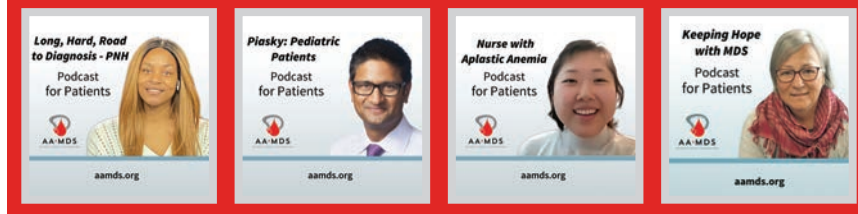
More Comprehensive Disease Management Tools

The Foundation provides free educational materials on aplastic anemia, MDS, and PNH, with the most current information—all developed with the needs of our patients and their support system in mind.

Our patient toolkits provide the latest disease and treatment information, and in 2024, we added information on treatment adherence, finding a specialist, and building a strong healthcare team. These additions will help empower patients to take control of their care.



Podcasts for Our Community by Our Community



In 2024, the treatment landscape continued to evolve with the introduction of newly approved therapies to improve patients' quality of life. Our patient podcast series offers a quick and effective way to share information about these new treatments. When a new treatment is approved, we promptly release special podcast episodes because we know patients must make informed, real-time decisions about their care.

In response to the community's need to hear from fellow patients, we have increased the number of patient stories featured in the Foundation's podcast series. These stories provide personal insights into the challenges of obtaining a diagnosis, making treatment decisions, and the lessons learned along the way, highlighting their triumphs as well.



Live Access to Experts

The AAMDSIF patient webinar series covers various topics, with live presentations by medical experts. This series is designed to ensure that anyone, regardless of their location, can access these experts. All webinars are posted on our website. In 2024, AAMDSIF webinars covered the following topics:

- New research findings
- Strategies for managing fatigue
- Nutrition
- Blood transfusions
- Iron overload
- Palliative care

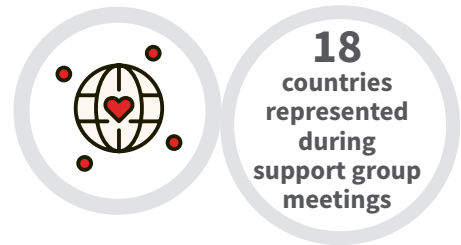
The webinars also included patient panels, providing an opportunity for patients to share their experiences.



Patient-Centered Programs

Providing Support

Our strong, vibrant community is our greatest asset. Throughout 2024, the Foundation's commitment to **taking the journey together** facilitated meaningful opportunities for our community to connect, share, and grow.



Nationwide Patient & Family Conferences

AAMDSIF's free, one-day Patient & Family conferences provide patients and their families with tailored support and education opportunities. These regional conferences offer access to some of the world's leading medical experts on aplastic anemia, MDS, and PNH. Attendees have the chance to meet other patients in person, fostering a sense of community and reducing feelings of isolation.

This year, we introduced a "Patient Story" session at each conference. These sessions featured a patient courageously sharing their personal story of diagnosis, treatment, and living well. This powerful narrative sets a positive tone for the day and helps attendees feel connected and understood.

At our conference in Los Angeles, we included a track focused on bone marrow transplants, which featured a session on addressing cardiovascular concerns post-transplant. In Chicago, the conference offered two interactive sessions: one on managing fatigue through gentle yoga and another discussion led by Dr. Swapna Thota about quality of life after diagnosis.

Additionally, each conference featured selected virtual sessions available as webinars for those unable to attend due to health reasons.

Global Community: Virtual Support Groups

The Foundation's monthly virtual support groups offer frequent and flexible opportunities for community members to connect. This highlights our strong commitment to ensuring that all patients and their support networks, regardless of their location, receive the necessary support and resources. In 2024, AAMDSIF hosted more than 1,100 registrants for over 120 support group meetings.



"I attended my first group and was so happy to connect with other people. I learned so much and look forward to attending other meetings"
—Debra M.

"Meeting other people is one of the greatest values of attending a conference in person. The conferences provide an opportunity for you to connect with others in your community for support." — Lisa U.



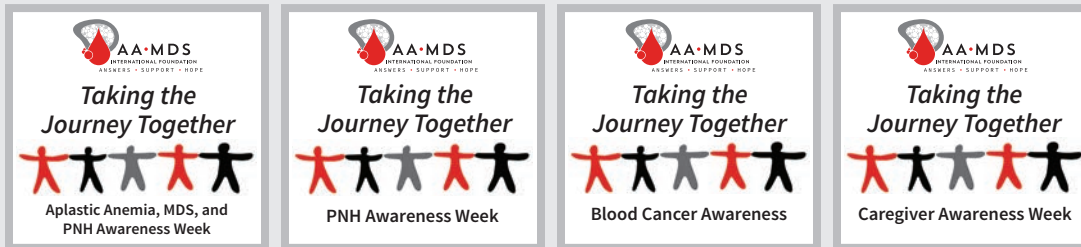
The Foundation's Patient and Family HelpLine staff is ready to answer your questions and provide you with the support you need.

To speak with an information specialist, please call (800) 747-2820, option 2, or email help@aamds.org.

Patient-Centered Programs

Providing Hope

Taking the Journey Together: Building Awareness



Raising awareness through designated days or weeks is a way to shine a spotlight on diseases and people that need championing. Throughout 2024, AAMDSIF hosted four awareness weeks (above)—all centered around the theme of **“Taking the Journey Together.”** These activities included live webinars featuring medical experts, interactive patient panels, and virtual support groups that connected patients with their support networks.

A Home for Hope

In 2024, we collected “Stories of Hope” from community members to share on our website as inspiration! Please enjoy some excerpts from three of these inspiring stories. You can visit our website to read their complete stories and find other stories of hope we shared in 2024.

Shauna, MDS patient



Shauna was once told her life expectancy with MDS was only two years. She told her doctor, “If I’ve only got two years, I’m gonna make it the best two years of my life!” That was in 2015, nearly 10 years ago at the time of this posting. What should other patients remember after diagnosis? When telling her own story, Shauna embraces living with no regrets. She’s proud of herself, wanting to make her family proud. She refined what was important to her. Knowing she is in remission but not cured, she continues to make each day count. She says it’s never too late to turn your life around, but most importantly, “Never lose faith!”

Xisto, Aplastic Anemia patient



By the time he was 10 years old, Xisto had earned a Black Belt in taekwondo. Yet he was tired enough to fall asleep at unusual times. He was diagnosed with aplastic anemia and assigned to a pediatric specialty care unit, with wonderful nurses and caring doctors. Now, Xisto is in college studying for his Registered Nursing degree and license. He is teaching taekwondo and working as a nurse in pediatric home health—two extra jobs while he studies! Xisto has kind words for pediatric patients: “The people around you will get you through it.”

Lisa, PNH and Aplastic Anemia patient



Lisa loved her work as a teacher. Then Lisa faced a dual diagnosis of aplastic anemia and PNH. “I thought I was invincible—work was my life. If people knew I was ill, they would feel sorry for me and I didn’t want that. The mental side of the illness was way tougher than the physical treatment and symptoms.” Things are looking up for Lisa after a successful stem cell transplant. She is indebted to her donor and medical team for her second chance at life and feels so blessed to be here. Her mantra remains “Look for the light in every day” and Lisa now devotes as much time as she can to helping others find their light too.

Creating a Lasting Legacy

The Foundation’s planned giving program offers a powerful and meaningful way to make a lasting impact. Through gifts made in wills, insurance policies, retirement accounts, or trusts, these generous donors thoughtfully included AAMDSIF in their estate planning.

Their foresight and generosity extend their legacy of support and provide a foundation of hope for those we serve—now and in the years to come.

With gratitude, we recognize the following individuals who have made this extraordinary commitment to our community.

Considering leaving a lasting legacy? Contact us at (301) 279-7202 or at development@aamds.org to learn how planned gifts can benefit AAMDSIF.



Estate of Florentine Camenisch



Estate of Virginia O’Allen



Nancy Upp Potter Charitable Trust



Estate of Mark Allen Thompson



Estate of Howard Ziff



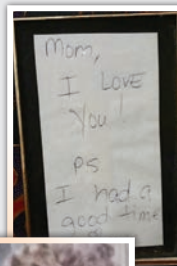
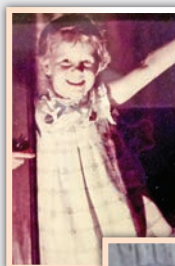
In 1988, my daughter Annie began to experience constant nose bleeds and bruising. We initially thought she had Leukemia. Within a difficult and stressful two weeks we found out she had Aplastic Anemia.

Annie spent the following two years in and out

of the hospital. She was transfusion dependent and developed other significant problems.

She spent the last two years of high school in the hospital or home schooled. Luckily, she was able to go to prom and graduation and had a wonderful support system with her friends. When she turned 18, she started making her own decisions about her treatment. I wasn’t pleased, but I respected her decisions. She even went as far as signing a DNR.

She decided to receive a bone marrow transplant. The day we got the news of the match was the same day the Berlin Wall came down. She rushed into the house that night yelling at me if I had heard the news? I told her “Of course; you already told me we have a match.”



She promptly told me “NO NOT THAT... the Berlin Wall is coming down!” It was so like her to be so excited to witness the wall coming down that night on the news. We proceeded to watch the celebration all evening.

Preparation for the transplant started which involved receiving chemotherapy to destroy the existing bone marrow, shaving her hair and then side effects. Her attitude remained positive, but there were bad days too. It was hard for me to stay strong for her as well.

Within a few months things started going downhill. She was going into liver failure. She was back in isolation and in a coma. With her room full of friends and family, we just prayed, cried and told “Annie” stories. On the early morning of October 15th, 1990, she took her last breath, and she was at peace.

I was fortunate to have a wonderful daughter. In her memory I have included the Aplastic Anemia & MDS International Foundation in my estate planning. Including the Foundation in my will ensures that Annie’s legacy lives on not only in the memories of those who loved her but also helps to fund programs that help future patients and families. I encourage you to do the same in memory of our families.

Barbara Wiens, mother of Ann Marie “Annie” Vermersch

March for Marrow: Walking for a Cure



“I think the wonderful thing about the Walk for PNH is that it gives patients and families an opportunity to get together and talk about the journey and embrace their rare disease community.”
— Erin Olivier



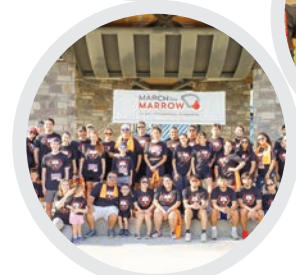
Erin Olivier, PNH Walk Founder (right) with Marlena Connor, PNH Walk Co-Chair.



The Foundation’s annual March for Marrow events are a cornerstone of our fundraising and awareness efforts. In 2024, patients, families, friends, healthcare professionals, and corporate partners gathered in communities nationwide—united by a shared mission and a spirit of hope.

Thanks to the incredible dedication of our volunteers, participants, and generous sponsors, we raised more than \$216,000 to support AAMDSIF’s programs and services. These funds significantly impact the lives of people affected by bone marrow failure diseases, while these events offer meaningful gathering opportunities for community members.

We would like to extend our special thanks to our March for Marrow volunteer leaders who inspire the community to take action in raising funds and awareness for rare bone marrow failures diseases. A heartfelt acknowledgment goes to Stephanie Hamm, the Chair of the Texas and California March for Marrow events; Debby Cook, the Chair of the Virginia March for Marrow; and the co-chairs of the Walk for PNH: Erin Olivier, Marlena Connor, and Hetal Soni.



“I’ve been participating in the walk for the better part of the last six years. The walks are a great experience to connect with other patients, make friends, get the latest treatment information from providers, and, of course, help drive donations and support to find a cure and alternative treatments”
— Dario Spina



Stephanie Hamm, Chair of Texas and California March for Marrow Events.

Convening Experts to Improve Patient Outcomes

AAMDSIF continues to be a trusted partner for clinicians and researchers. Together, we strive to identify knowledge gaps in the diagnosis and treatment of bone marrow failure and work collaboratively on research priorities to further our mission.

Ninth International Bone Marrow Failure Disease Scientific Symposium



Attendees of the 2024 Scientific Symposium learning and networking.

AAMDSIF hosted our ninth International Bone Marrow Failure Disease Scientific Symposium, bringing together 149 leading physicians, researchers, and clinical investigators specializing in bone marrow failure diseases from over nine countries. The Symposium was chaired by Richard Stone, MD and Neal Young, MD—both members of our Medical Advisory Board.

Dr. Guillermo Garcia-Manero received the AAMDSIF Leadership in Science Award for his significant contributions to treating bone marrow failure diseases and his dedication to AAMDSIF's mission. He is a professor in the Department of Leukemia at The University of Texas MD Anderson Cancer Center and serves as the Chair of the Section of Myelodysplastic Syndromes (MDS).



His research aims to improve patient outcomes by studying the molecular basis of leukemia and MDS and developing new therapies. He directs the MDS/AML Moon Shot program, which accelerates scientific discoveries and facilitates the transition of new treatments into clinical practice, resulting in over 500 publications and numerous clinical trials. Dr. Garcia-Manero is also committed to professional and patient education through his long-standing role on the AAMDSIF Medical Advisory Board since 2007.

Providing Equitable and Inclusive Care: 3rd Annual Diversity, Equity, and Inclusion Community Symposium

For the third year in a row, the Foundation remained committed to championing equitable and inclusive care for patients living with bone marrow failure disease. This hybrid event, organized in partnership with the University of Miami and chaired by Namrata Chandhok, MD, showcased this dedication. It brought together health professionals focused on addressing the existing gaps and barriers in patient care for individuals with bone marrow failure.

Highlights from the Diversity, Equity, and Inclusion Community Symposium

In addition to sessions on significant disease treatment updates, the symposium featured panel discussions on health equity issues impacting patient outcomes. These included:

- Innovative Approaches to Address Unmet Needs of Patients
- Making Clinical Trial Participation Diverse and Accessible
- The Spectrum of Supportive Care for Bone Marrow Failure Patients

Noteworthy takeaways

Shaanan Beg, MD

National Cancer Institute

Dr. Beg emphasized that true progress in patient care demands collaboration. From improving access to clinical trials to streamlining how we capture and share data—the focus is on creating a connected and diverse patient ecosystem. When we standardize and share data openly, we empower every voice in the fight against bone marrow failure disease.

Ingrid Barrera, PsyD

University of Miami

Holistic care means seeing the person, not just the disease. Dr. Barrera brought forward a powerful reminder: patient care isn't complete without comprehensive psychosocial support. This includes access to mental health professionals, patient and caregiver education, support groups and peer connections, stress management techniques, and family-centered care.

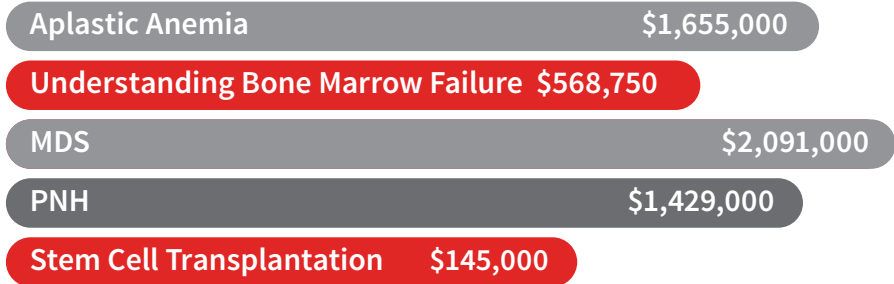
Investing in Research to Find a Cure

Promoting research to find a cure is central to the Foundation’s mission. Since 1989, the Foundation has invested over \$5.9 million in research initiatives, focusing on funding projects that show the greatest potential for developing treatments and cures for patients with aplastic anemia, MDS, and PNH. Each year, AAMDSIF awards two-year grants for research projects to attract investigators to this field and improve the understanding of these rare diseases.

35 Years of Investing in Cures

107 grants Total money invested: **\$5,918,750**

Total money invested by disease:



Many past grantees, empowered by the Foundation’s initial investment in their innovative research, have risen to leadership positions in their fields. By validating their approach with the AAMDSIF grant, these grantees have not only secured significant research support from the NIH, private foundations, and their institutions, but they also now lead their own research labs.

“These four new grantees have demonstrated outstanding dedication, innovation, and potential for making significant contributions, now and in the future, to the advancement of medical knowledge and patient care. Our organization is proud to support their research and looks forward to witnessing the positive impact their work will have on the bone marrow failure community,”

— Brad A. Wong, CEO & Executive Director of the AAMDSIF

Catching Up with a Past Grantee

Joseph Oved, MD

Dr. Oved from Memorial Sloan Kettering Cancer Center, was awarded an AAMDSIF research grant in 2017. This grant played a crucial role in his study aimed at identifying which pediatric patients are most suitable for treatment with immunosuppressive therapy versus bone marrow transplantation. The findings from this research have contributed to the development of Emapalumab, a potential new treatment for pediatric aplastic anemia.



Meet Our 2024 Research Grantees

Brian Ball, MD

Assistant Professor, City of Hope, Duarte, CA

Grant generously funded by the family and friends of Hadi Abdur Rasheed

Impact of recipient somatic mutations on outcomes after transplant in pediatric and adolescent/young adult patients with aplastic anemia or Inherited Bone Marrow Failure Syndromes



Gloria Gerber, MD

Assistant Professor, Johns Hopkins University School of Medicine

Optimizing Complement Inhibitor Selection and Monitoring in PNH



Parveen Shiraz, MD

Instructor, Stanford University School of Medicine

Evaluating Thymic Stromal Lymphopoietin as a Biomarker for Acute Graft versus Host Disease



Ganesh Kumar Viswanathan, DM

Associate Professor, All India Institute of Medical Sciences New Delhi

Structural variations & copy number alterations in adult immune aplastic anemia



Pivotal Year for Global PNH Registry

In partnership with the National Organization for Rare Disorders (NORD®), AA-MDSIF launched the Global PNH Registry in 2021 to advance PNH patient care. Your support helps the Foundation continue its recruitment efforts to expand the registry's reach and amplify the data.

Publication of the First State of the Registry Report

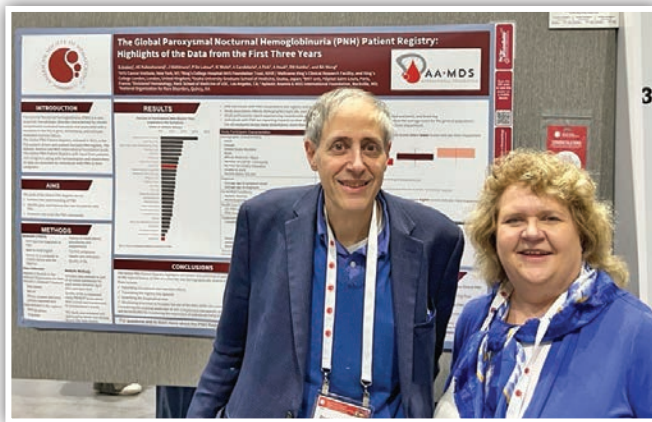
This comprehensive report summarizes the first three years of the Registry data collection and is now available to current sponsors. This report is a key tool in enhancing our understanding of PNH, improving patient care, and empowering and uniting the PNH community.

Highlights of the first three years presented at the American Society of Hematology (ASH) Annual Meeting

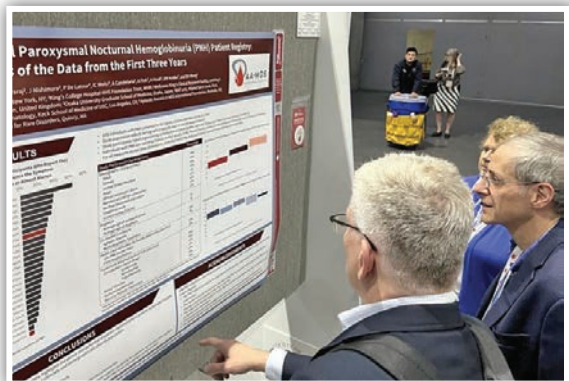
Summary of key data collected since the Registry was initiated in April 2021 up to April 2024 was accepted for presentation at the 2024 ASH Annual Meeting, the premier medical meeting in hematology. This recognition highlights the significance and validation of the Registry's efforts.

Spanish Translation

We are proud to announce that all PNH Registry surveys and related materials have been translated into Spanish, a significant step in our commitment to diversity and inclusion.



Medical Advisory Council Chair, Dr. David Araten (left), and Principal Investigator, Alice Houk (right), presenting data from the registry and interacting with ASH attendees.



The Registry is governed by a Medical Advisory Committee of international PNH experts:

David Araten, MD (Chair) –
New York University

Austin Kulasekararaj, MBBS, MD,
MRCP, FRCPath – Kings College
London, UK

Jaroslawn Maciejewski, MD, PhD –
Cleveland Clinic

Jun-ichi Nishimura, MD, PhD –
Osaka University, Japan

Christopher Patriquin, MD, MSc,
FRCPC – University of Toronto,
Canada

Regis Peffault deLatour, MD, PhD –
Hôpital Saint-Louis, Paris, France

Antonio Risitano, MD, PhD –
AORN Moscati, Avellino, Italy

Jeff Szer BMedSc, MB, BS –
Royal Melbourne Hospital, Australia

Ilene Weitz, MD – University of
Southern California

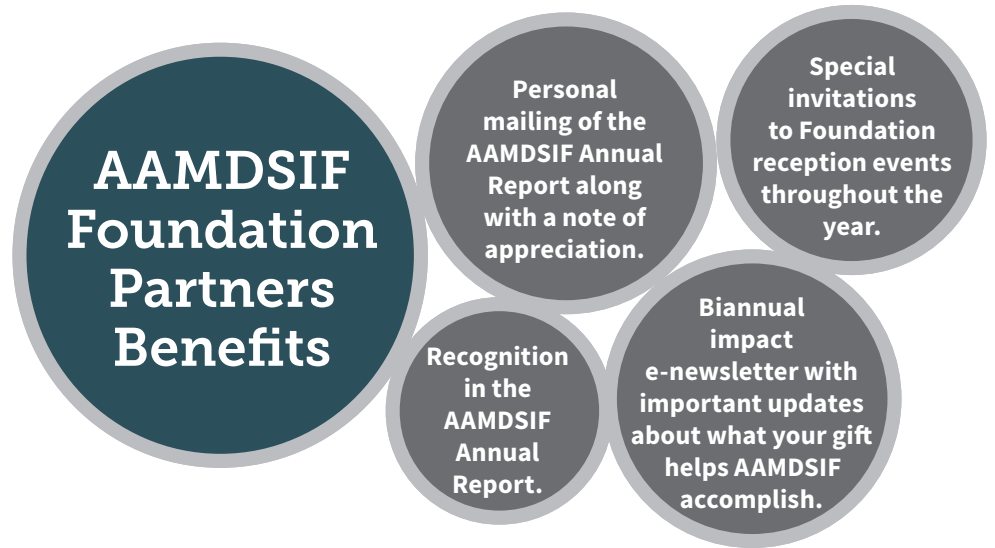
We are grateful to the generous current sponsors of the Global PNH Patient Registry: Apellis Pharmaceuticals, Genentech, Inc., and Novartis Pharmaceuticals.



**288
Enrolled
Participants**

Foundation Partners Program

Our **Foundation Partners** Program celebrates the powerful generosity of individual supporters who donate or raise \$1,000 or more annually to AAMDSIF. Their generous contributions enable AAMDSIF to provide answers, support, and hope to bone marrow failure disease patients and their families.



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Special Thanks to These Generous Funders

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We're deeply grateful for the support of our corporate and foundation funders, who recognize AAMDSIF as the world's leading nonprofit organization dedicated to supporting patients and families living with aplastic anemia, MDS, and PNH. We appreciate your commitment to the patients and families we serve.

Foundation Funders

The Amy Beck Charitable Foundation	James A. Reep Family Foundation
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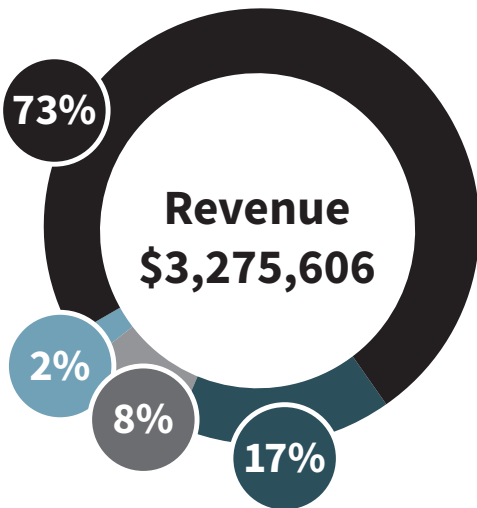


Monthly Donations:

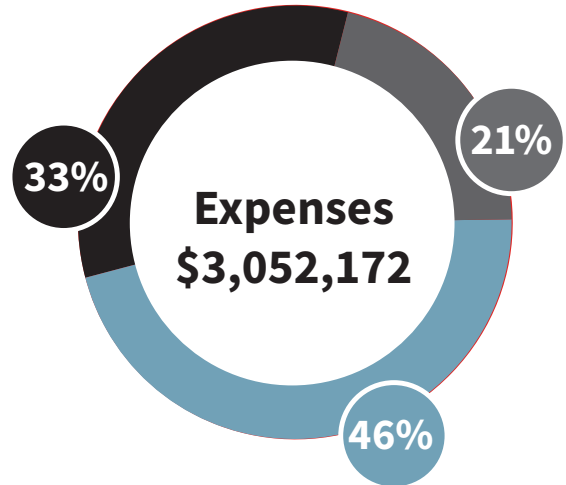
We are incredibly grateful to our monthly donors. A monthly donation helps deliver our services consistently throughout the year.

2024 Financials

The Aplastic Anemia and MDS International Foundation



Corporations: \$2,390,095
Individuals: \$540,328
Other Revenue: \$263,333
Family Foundations: \$81,850



Program Services: \$1,407,832
Research: \$995,224
Fundraising & General: \$649,116

2024 Balance Sheet

Assets

Cash & cash equivalents	\$1,560,518
Contributions and grant receivable	722,213
Investments	4,157,306
Fixed assets, net	235,035
Right-of-use assets, net	524,507
Other assets	28,127
Charitable Remainder Trust	236,972

Total assets **\$7,464,678**

Liabilities

Accounts payable and accrued expenses	\$166,702
Lease liabilities	580,417

Total Liabilities **\$747,119**

Net Assets

Without donor restriction	\$5,302,941
With donor restriction	1,414,618

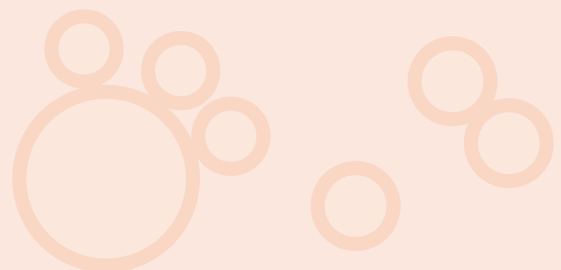
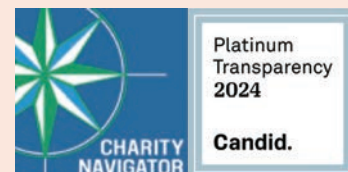
Total net assets **\$6,717,559**

Total liabilities & net assets **\$7,464,678**

**Please visit aamds.org/financials for full financial reports.*

2024 Financials

The Aplastic Anemia and MDS International Foundation is recognized as a fiscally responsible nonprofit dedicated to financial transparency, earning a Four-Star Rating from Charity Navigator and a platinum-level GuideStar Transparency rating.



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MDS International Foundation**

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