



Let's Celebrate 30 Years!

As we approach Neutropenia Awareness Day on October 25 (declared in Michigan in 2022), we also celebrate the **30th anniversary of the National Neutropenia Network**. Reflecting on three decades of dedication, we recognize the countless lives we've touched and the incredible advancements in science that have improved care for those living with neutropenia.

Let's take this opportunity to honor those affected by the disease, cherish the memories of those we've lost, and express our gratitude to the healthcare professionals who support and care for us. Together, we can unite in our mission to raise awareness. By doing so, we can advocate for continued advances in treatment, and one day, a cure. Join us in this important celebration!

Community Support



Introducing our Junior Ambassadors

We are excited to introduce 3 new Junior Ambassadors: **Maison, Rylan and Hannah!** In addition, we are excited to have **Jaiya, Tate, Ella Jewell and Mateo** continue their support to the junior ambassador program! These seven ambassadors are using their own journey with neutropenia in a positive light as they spread awareness and do good in the community. We are so proud of

each of them and excited for their year ahead!

As we highlight each of them over the next couple months, we hope their stories will inspire and give hope to all in this community.



Meet Our Senior Ambassadors

Who better to provide support than those who understand neutropenia firsthand—both individuals living with the condition and the caregivers of children affected by it? We extend a warm welcome to Tracy Lemmons, the newest member of our Senior Ambassador group!

A special thank you to Chrissy, Maria, Jason, Lauren, and Heidie for your unwavering commitment and invaluable contributions. Your dedication to engaging our community and raising awareness about neutropenia is truly inspiring. Together, we're making a difference!

To discover more about our ambassadors, our peer support program, or to seek assistance, please visit our website. We're here to help!

[Learn More](#)

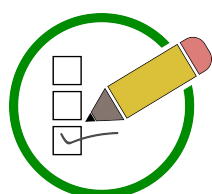
Save the Date



2025 Family Conference

We are thrilled to be planning the 2025 Family Conference! Once again, it will be held in Cincinnati, Ohio at The Graduate Hotel. More details to come but save the date and we hope to see new and familiar faces next year!

Community Feedback



Thank you to all that filled out our survey! But if you missed it, there is still time!

With the 2025 Family Conference in the planning stages, we are keeping the survey open a little longer so we can use the results as a tool in determining topics for the conference as well as upcoming articles and webinars. We

[Click for the Survey](#)

Don't Miss Out! SCNIR Virtual Family Day



SCNIR Family Day is next week, Saturday, Nov. 2, 1-4 pm ET! This will be an opportunity for patients, families, and medical providers to hear information about neutropenia and updates from the Severe Chronic Neutropenia International Registry. This virtual conference will discuss topics important to both pediatric and adult patients with neutropenia and their families. SCNIR investigators will share knowledge learned from the SCNIR and discuss ongoing research to improve the health of people with neutropenia conditions.

Attendees do not need to be a part of the SCNIR to join.

There will be Q&As for patients and families and presentations on topics such as genetic and autoimmune neutropenia, bone marrow transplant, living with chronic neutropenia, and others.

[Learn More and Register Today](#)

Message from our Partner: Probably Genetic Helping Unlock More Answers for Patients

At Probably Genetic, we often witness patient journeys that feature diagnoses like neutropenia but may only be part of an individual's health story. We believe every patient deserves direct, clear, and complete answers. We focus on diagnosing rare diseases that often go unrecognized, providing patients and their families with the clarity they need to pursue effective care. To better help identify the underlying causes of their symptoms, Probably Genetic offers no-cost genetic testing and personalized support through free genetic counseling. These licensed genetic counselors help families understand their test results to make informed treatment decisions and how to discuss a new diagnosis with their care teams. Our free genetic testing services and genetic counselors help patients gain clarity, explore treatments, and plan for the future. Our testing utilizes whole exome sequencing (WES), which is a holistic approach that allows us to identify genetic variants that may be the root cause of a patient's symptoms. **For more information on how Probably Genetic can help your family, visit [our website](#) today.**

The National Neutropenia Network depends on support from the friends and families of those whose lives have been forever changed and challenged by this rare disease. Thank you for your support.

[Donate Today](#)



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Quarterly Newsletter

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