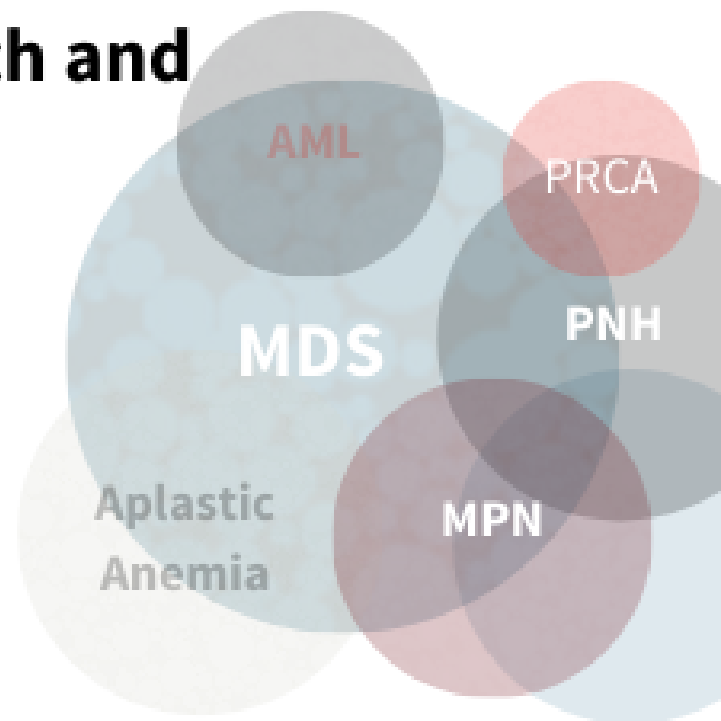


# Understanding PNH: New Research and Patient Opportunities



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## Our Mission

The Aplastic Anemia & MDS International Foundation is the world's leading nonprofit health organization dedicated to supporting patients and families living with aplastic anemia, myelodysplastic syndrome (MDS), paroxysmal nocturnal hemoglobinuria (PNH), and related bone marrow failure diseases. The Foundation provides answers, support, and hope to thousands of patients and their families around the world.

We are a patient-focused, patient-centered organization, serving patients and families throughout the three phases of bone marrow failure diseases:

- the life changing phase of diagnosis
- the life threatening phase of treatment
- the life long phase of living with a chronic disease



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1. Use the Q & A box on the bottom of your screen. Type your question into the box and press **ENTER**.
2. Please do not include private health information about the patient in your question. Our presenters cannot answer specific questions related to your diagnosis or treatment.
3. We will try to answer all questions during the webinar but may not be able to get to everyone.
4. If we do not get to your question, please send it to us via email at [help@aamds.org](mailto:help@aamds.org), by calling the office at (800) 747-2820 x2 or by sending us a message on social media.

## **Aplastic Anemia & MDS International Foundation - Your Resource of Choice for Bone Marrow Failure Disease Education and Support**

- For up-to-date guidance, webinar links, resources and Frequently Asked Questions on COVID-19, visit [www.aamds.org/education/covid-19](http://www.aamds.org/education/covid-19)
- Contact AAMDSIF via e-mail to [help@aamds.org](mailto:help@aamds.org)
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as we have for the past 36 years.*

