Registration Now OPEN
Fall Virtual Patient & Family Conference
September 11, 2021

6 Disease Experts - 5 Support Groups - 1 Day

Click Here for More Details

Register NOW

September 11 is the Date!
Thanks to technology - we can support the patients and families served by
AAMDSIF by registering for our Virtual Event or simply by donating.

Register Yourself or Your Team Now!
Have you been wondering how to get involved with PNH research but did not know where to start? Here’s your chance! Check out the Global PNH Patient Registry at pnh.iamrare.org

If you would like to be in a safe place to talk about what you are going through, or if you would like to encourage others, please check out the Virtual Support Groups HERE. Together, we are strong!

Current schedule:

- Wed, 08/18/2021 - 5:00pm (EDT) Transplant Virtual Support Group
- Tue, 08/24/2021 - 7:00pm (PT) Aplastic Anemia Virtual Support Group (Pacific Time)
- Wed, 08/25/2021 - 6:00pm (EDT) Young Adult Virtual Support Group
- Thu, 08/26/2021 - 5:30pm (EDT) Teen Support Group for Bone Marrow Failure Diseases
- Thu, 08/26/2021 - 7:00pm (EDT) Pediatric Parent Virtual Support Group
- Sat, 08/28/2021 - 10:30am (EDT) Aplastic Anemia, MDS, and PNH Virtual Support Group
- Mon, 08/30/2021 - 11:00am (EDT) PNH Virtual Support Group
Find webinars with speakers like these, plus patient panelists!

Here are the next All-New Webinars for you!

All Times EDT-US

Thu, 08/19/2021 - 2:00pm (EDT)  
Participating in a Clinical Trial: Patient Panel

Wed, 08/25/2021 - 10:30am (EDT)  
AAMDSIF COVID-19 Update: Returning to School

Mon, 08/30/2021 - 2:00pm (EDT)  
Living with a Pediatric Bone Marrow Failure Disease: a Parent Panel Discussion

Thu, 09/09/2021 - 3:30pm (EDT)  
Advances in Treating PNH

Mon, 09/13/2021 - 3:00pm (EDT)  
Molecular Genetics of MDS/MPN Overlap Syndromes

Wed, 09/15/2021 - 1:00pm (EDT)  
Cardiac Complications with Bone Marrow Transplant

Fri, 09/24/2021 - 12:00pm (EDT)  
Preparing for Transplant

AUGUST
Minority Donor Awareness Month

Did you know that donating stem cells (like a blood donation) is the way that
most transplants for bone marrow failure patients are done? Ask everyone you know to register today at bethematch.org.

A phase I/II Study of REGN7257 (Anti-interleukin 2 receptor subunit gamma [IL2RG] monoclonal antibody) in patients with severe aplastic anemia that is refractory to or relapsed on immunosuppressive therapy

AAMDSIF does not recommend, endorse, or make any representation about the efficacy, appropriateness or suitability of any clinical trial listed on our website. Pharmaceutical company-sponsored content is highlighted only to give additional information about the trial. All trials are listed on https://clinicaltrials.gov/. Always seek the advice of your physician or other qualified health care provider with any questions you may have regarding a clinical trial, and never disregard professional medical advice or delay in seeking it because of something you have read on our website.

FDA grants Breakthrough Therapy Designation for Venclexta in combination With Azacitidine for the treatment of patients With Myelodysplastic Syndromes - Find out more here.

FDA approves Belumosudil for chronic graft-versus-host disease - Read the announcement here.
AAMDSIF is currently conducting a unique survey aimed at better understanding the emotional and mental health need of MDS patients and caregivers. We'll use this information to improve our resources and support services and we'll also share the findings with patients, families, and health care providers.

If you're interested in participating in this study, send an e-mail to houk@aamds.org to request a link to the survey, or start here. Any information provided in the survey will be treated with the strictest confidence and will remain entirely confidential. The survey will take about 25 minutes to complete and you will be compensated for your time.

The webinar helps patients and caregivers develop a clearer understanding of treatment options, side effects, quality-of-life changes, and financial impacts associated with MDS. Watch HERE.

Imagine being young and thousands of miles away from family.

Now add a diagnosis of MDS.

Discover how Shauna lives each day to the fullest, finding the beauty in every person here.
A final thought:

All of our conferences, webinars, educational opportunities, and patient support are made possible from gifts by people like you, friend. Your donation will help us reach more patients and families.

Yes, I'll Help!

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