

HISTIOCYTOSIS  ASSOCIATION™
A Rare Community

(<https://histio.org/>)

Thank You!

Thank you for all you have done this year, in any and every way, to uplift and support the histiocytosis community. We are so thankful to serve alongside you and strive to provide the support, resources, and connections you need along every step of this journey. Histiocytosis is a club no one wants to be a part of, but **YOU** keep making it more bearable and hopeful.

Click the button below to see the impact YOU have made in patients and families lives this year.

From all of us at the Association, happy holidays and have a wonderful new year.

We have big goals for 2024!

**Your Impact
(<https://histio.org/wp-content/uploads/2023/11/2023-Histiocytosis-Association-Impact-Report-compressed.pdf>)**

Three Podcast Episodes this Month! It's Our Gift to You!



Episode 059:
HopeCast Podcast Edit



Episode 060:
Sharing Difficult
Diagnoses w/ Children
w/ Dr. Lana Harder

This month we have released 3 new Beyond the Diagnosis episodes.

The first episode is episode 58 with Ambassadors Ana, Nate and Melinda discussing their experience with the various Global Genes Rare conferences held in September.

The next episode, episode 59, is the audio-only version of the HopeCast. If you haven't seen the HopeCast yet, but want to be able to listen to it in the background, this is the episode for you.

background, and is the episode for you.

Finally, our final episode, released today, is episode 60 with Dr. Lana Harder, discussing how to share difficult diagnoses with children.

Listen Below!

Listen Now
(<https://histio.org/beyond-the-diagnosis-podcast/>)

Resource of the Month

DOLLAR FOR 

Dollar For is a resource we partner with who cuts the high costs for healthcare for those in need. They breakdown the system of Charity Care with patients and help them get costs deducted or forgiven if they qualify.

Click below to learn more and see if you are eligible!

Learn More
(<https://dollarfor.org/>)

Family Toolkit Webinar



Two weeks ago, we hosted a webinar partnering with the organization Got Transition to walk histio patients and families through some of the challenges associated with transitioning from pediatric to adult care and how to work through those challenges using their family toolkit.

In case you missed it, click below to view the recording. Visit our webinars page for this talk and others <https://histio.org/resource-overview/webinars/>. (<https://histio.org/resource-overview/webinars/>)

Watch Now!
(<https://youtu.be/511mSPz6OnQ>)

New Study Opportunity for LCH



(<https://youtu.be/1HYNyEg6hYc>)

The purpose of this study is to see if treatment with mirdametinib in patients with LCH or other histiocytic disorders will be better than current treatments and with fewer side effects. The study is open to all patients 2 or older (yes, children and adults) with any histiocytic disorder (LCH, JXG, ECD), newly diagnosed or recurrent/relapsed. Learn more about the study and eligibility criteria, here: <https://clinicaltrials.gov/study/NCT06153173?intr=mirdametinib&rank=1> (<https://clinicaltrials.gov/study/NCT06153173?intr=mirdametinib&rank=1>) or on our website at <https://histio.org/research/clinical-trials/> (<https://histio.org/research/clinical-trials/>).

Watch Here (<https://youtu.be/1HYNyEg6hYc>)

Connect With Us

UNITE
MATCH
CARE

Bridging the Gap
Transplant Care
PEDIATRIC TO ADULT



Sarah Rusch
NMDP/ Be the Match

Join us for a live webinar
TUESDAY, January 14th
@ 11:00 AM EST

Hosted by:



Next month, on **January 14th**, we will be hosting a webinar to guide you through best practices and gaps in transition of care for bone marrow transplant patients and how we can best fill them.

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The talk will be hosted by Sarah Rusch, NMDP/Be the Match. The talk will cover all blood-based conditions including leukemia, lymphoma, and bone marrow transplant (BMT).

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The information will be beneficial for patients with multiple myeloma and other blood-based conditions. Please share this opportunity with family and friends to help them learn how to receive and advocate for transition of care following BMT.

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<http://www.facebook.com/histiocytosis>

This webinar is free, but registration is required.

Register Here
(<https://us02web.zoom.us/j/81234567890>)
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SUH

<http://www.twitter.com/histiocytosis>

http://www.instagram.com/histiocytosis_association

<http://www.youtube.com/histiocytosisassoc>

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Stay Connected to the Histo Community

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