A Rare Community

The Histiocytosis Association is a global nonprofit organization, based in Pitman New Jersey, that is dedicated to addressing the unique needs of patients, families, and physicians impacted by histiocytic disorders. Founded in 1986, the Association provides educational and emotional support, raises awareness, and funds research leading to advancements in treatment and cures.

The Association's website includes an interactive physician directory, educational videos & webinars, registry and clinical trial resources, and opportunities for peer connection. We also monitor our phone line and emails regularly as we seek to connect with every patient and family member to provide individualized support.

Histiocytosis Association
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New York 2023 Patient & Family Conference on Histiocytic Disorders

MEETING AGENDA

Saturday, September 23rd: 9am - 3pm EST

8:00 - 9:00am	Check-In & Breakfast
9:00 - 9:30am	Welcome/Opening Remark
	Patient and Family Introductions
9:30 - 10:00am	Diagnosis Under the Microscope
10:00 - 11:00am	HLH Session
10:00 - 11:00am	Pediatric LCH Overview
	Adult Histiocytosis Overview
11:00 - 11:30am	Survivorship and Long-term consequences
11:30 - 12:00pm	Open Q & A
12:00 - 12:45pm	Lunch
12:45 - 1:30pm	Living with Histiocytosis: Psychological Aspects of Care
1:30 - 1:50pm	Update on Clinical Trials
1:50 - 2:00pm	Closing Remarks
2:00 - 3:00pm	Patient and Family Workshop

12:45 - 1:30pm

PRESENTING SPONSOR

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