



NORD[®]
National Organization
for Rare Disorders

SHARING HOPE

 *meetings*

FOR PATIENTS & CAREGIVER EDUCATION

Paroxysmal Nocturnal Hemoglobinuria (PNH)

Each year, NORD hosts meetings around the country to help bring together hundreds of patients and families impacted by Paroxysmal Nocturnal Hemoglobinuria (PNH).

These meetings provide opportunities for new and familiar faces to come together, share experiences, learn from each other and experts, and to offer one another support and hope. Meet staff from the PNH Foundation and hear what services they have to support you and your family.

NORD is inviting ALL patients diagnosed with PNH and their caregivers to join us this year as we continue to empower a strong and knowledgeable community of PNH advocates!

2017 Dates & Locations

March 24-25 • Sterling, VA

April 7-8 • Miami, FL

April 28-29 • Charleston, SC

June 2-3 • Dallas, TX

July 14-15 • Denver, CO

September 15-16 • Detroit, MI

October 27-28 • Providence, RI

November 10-11 • Irvine, CA

**Alone we are rare.
Together we are strong.[®]**

NORD, a 501(c)(3) organization, is a patient advocacy organization dedicated to improving the lives of individuals and families impacted by a rare disease. NORD, along with its more than 260 patient organization members, is committed to the identification, treatment, and cure of rare disorders through programs of education, advocacy, research, and patient services.

Register Today! This is a FREE event. Contact: Elizabeth Hoey at NORD

Email: ehoey@rarediseases.org Online Registration: www.rarediseases.org/PNH-2017

Learn more about NORD by visiting:
rarediseases.org

SHARING HOPE



meetings

FOR PATIENTS & CAREGIVER EDUCATION

Paroxysmal (PNH)
Nocturnal
Hemoglobinuria

A Meeting to Connect, Learn and Share Hope Among Others Living with PNH

“ I look forward to [the events] every year, and meeting new people and seeing the familiar faces from past meetings... It really does a lot for me to be able to talk to people with PNH. I feel that we have an instant connection. ”

— Lisa from NORD's PNH Meeting



“ I cannot emphasize enough how important these patient meetings are. Whether you've been to 1, 2 or 10 meetings in the past, you are going to continue to learn new things. ”

— Jill from NORD's PNH Meeting

Connecting

Each event is another opportunity to meet new patients and families or reconnect with familiar faces. A welcome reception Friday evening helps attendees to break the ice.

Learning

NORD invites expert physicians in the diagnosis and treatment of PNH to present new information and share updates to help you better understand your disease.

Sharing Hope

Gain the hope and support that comes from sharing experiences in an open and safe environment with others who understand your journey as a patient or caregiver.

Register Today! This is a FREE event. Contact: Elizabeth Hoey at NORD

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Online Registration: www.rarediseases.org/PNH-2017

*Small stipends are provided to each patient to help defray the cost of travel. A hotel room will be provided to each patient for them and their family for one (1) night, by request only. These meetings are supported by a sponsorship from Alexion Pharmaceuticals, Inc. NORD does not endorse or recommend any specific medical treatments. NORD's patient programs are for informational purposes only. Patients must rely on the personal medical advice of their qualified health care professionals before seeking any information related to their particular diagnosis, cure or treatment of a condition or disorder.

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