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!

FREE EVENT: June 24-25*
Baltimore Marriott Waterfront
700 Aliceanna Street • Baltimore, MD 21202

Fri., June 24th: Welcome reception from 7-9pm (light buffet served)
Sat., June 25th: Registration & Educational Meeting 8am - 2:30pm (breakfast at 8am, lunch from 11:45am-12:30pm)

Speaker: Dr. Elizabeth Griffiths
Associate Professor
Roswell Cancer Center

“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.”
— Patient from NORD’s PNH Meeting

2016 Dates & Locations

June 24-25 • Baltimore, MD
Sept. 16-17 • New Orleans, LA
Sept. 30-Oct. 1 • Seattle, WA
Nov. 4-5 • San Antonio, TX
Dec. 2-3 • Boston, MA

Connecting
Learning
Sharing Hope

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.

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.

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.

For over 30 years, the National Organization for Rare Disorders (NORD) has led efforts to improve the lives of individuals and families impacted by rare disease through partnership programs of education, advocacy, research and financial assistance. Learn more about NORD by visiting: rarediseases.org

*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 an educational grant 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, care or treatment of a condition or disorder.

(NORD is a registered trademark of the National Organization for Rare Disorders. NORD is a registered 501(c)(3) charity.)