

SHARING HOPE



2010 Regional PNH Patient Support Meetings

Baltimore - Portland - Cincinnati

Boston - Denver - Chicago

Los Angeles - Nashville - Ann Arbor

New York City - Atlanta - Phoenix

The National Organization for Rare Disorders is a unique federation of voluntary health organizations dedicated to helping people with rare "orphan" diseases and assisting the organizations that serve them. NORD is committed to the identification, treatment, and cure of rare disorders through programs of education, advocacy, research, and service.

NORD / PNH Regional Meeting

Baltimore

March 13, 2010



NORD

National Organization for Rare Disorders

55 Kenosia Avenue, P.O. Box 1968

Danbury, CT 06813-1968

www.rarediseases.org

*Supported by an educational grant from
Alexion Pharmaceuticals*

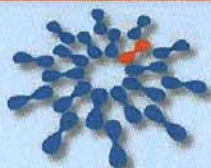


NORD

National Organization for Rare Disorders

SHARING HOPE

A Regional Meeting for People Living with PNH



NORD

National Organization for Rare Disorders

The National Organization for Rare Disorders invites patients and families affected by Paroxysmal Nocturnal Hemoglobinuria (PNH) to participate in a special event to meet other patients and speak with a medical expert.

This meeting will create an opportunity for us to come together in a small group, and gain the encouragement and hope that comes from meeting face-to-face with other PNH patients.

Everyone will be able to share personal experiences with each other, gain a better understanding of this orphan disease, and help others cope with the effects of their illness.

NORD will support the efforts of the PNH community by sharing accurate and useful information, and will explore ideas for future assistance.

DEDICATED TO HELPING PEOPLE
WITH ORPHAN DISEASES



Saturday, March 13, 2010

AGENDA

- 9:00 AM – 10:00 AM Breakfast
- 10:00 AM – 12:00 PM Support Meeting with PNH Expert
- 12:00 PM – 1:00 PM Lunch
- 1:00 PM – 3:00 PM Resources & Round Table Discussion

GUEST SPEAKER

Martin Weltz, DO, MPH, MBA, FACP
Hematology Oncology Associates
Greenbelt, MD

LOCATION

Baltimore Marriott Waterfront
700 Aliceanna Street
Baltimore, MD 21202

Please RSVP by March 9, 2010

Ms. Susan Olivo
203-744-0100
solivo@rarediseases.org

*This is a free event.
Accommodations provided if requested.*