

# SHARING HOPE



## 2010 Regional PNH Patient Support Meetings

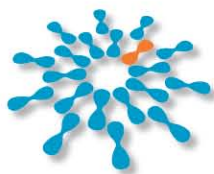
Baltimore - Portland - Cincinnati

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

National Organization for Rare Disorders

55 Kenosia Avenue, P.O. Box 1968

Danbury, CT 06813-1968

[www.rarediseases.org](http://www.rarediseases.org)

*Supported by an educational grant from  
Alexion Pharmaceuticals*

## NORD / PNH Regional Meeting

### Los Angeles

July 24, 2010



## 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.

**Saturday, July 24, 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

---

**Ilene Weitz, MD**  
Assistant Professor of Clinical Hematology  
Keck School of Medicine  
University of Southern California

## LOCATION

---

**Marina del Rey Marriott**  
4100 Admiralty Way  
Marina del Rey, CA 90292

**Please RSVP by July 20, 2010**

---

**Ms. Susan Olivo**  
203-744-0100  
[solivo@rarediseases.org](mailto:solivo@rarediseases.org)

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

DEDICATED TO HELPING PEOPLE  
WITH ORPHAN DISEASES

