

SHARING HOPE

A Regional Meeting for People Living with PNH

NORD

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, April 18, 2009

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 Round Table Discussion

GUEST SPEAKER

Monica Bessler, MD, PhD

Professor of Medicine
Professor of Molecular Biology & Pharmacology
Washington University School of Medicine St. Louis

LOCATION

Hotel Monaco Chicago

225 North Wabash Avenue
Chicago, IL 60601

Please RSVP by April 15, 2009

Ms. Michele Counihan

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



*Dedicated to Helping People
with Orphan Diseases*