



Sharing Hope

A Regional Support Meeting for People Living with PNH

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.

Denver, Colorado

Saturday, August 2, 2008

10 am • Support Meeting

Ilene C. Weitz, MD
Asst. Professor of Clinical Hematology
Keck School of Medicine
University of Southern California

12 pm • Lunch

Hotel Monaco

1717 Champa Street
Denver, CO 80202

Please RSVP by July 30

Ms. Michele Counihan
203-744-0100 ext 252
mcounihan@rarediseases.org

This is a free event. One night hotel accommodations provided if requested.

2008 Regional PNH Patient Support Meetings
Chicago • Denver • Orlando • New York • Los Angeles

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.

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