

SHARING HOPE

meetings

FOR PATIENTS & CAREGIVER EDUCATION

Atypical Hemolytic Uremic Syndrome

Each year, NORD hosts meetings around the country to help bring together hundreds of patients and families impacted by Atypical Hemolytic Uremic Syndrome (aHUS).

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

(aHUS)

support and hope. Meet staff from the aHUS Foundation and hear what services they have to support you and your family.

NORD is inviting ALL patients diagnosed with aHUS and their caregivers to join us this year as we continue to empower a strong and knowledgeable community of aHUS advocates!

FREE EVENT: August 26-27*

InterContinental Mark Hopkins
999 California Street • San Francisco, CA 94108



Speaker:
Dr. Ilene Weitz
Associate Professor of Clinical Medicine
Keck School of Medicine USC

Fri., Aug. 26th: Welcome reception from 7-9pm (light buffet served)

Sat., Aug. 27th: Registration & Educational Meeting 8am - 2:30pm
(breakfast at 8am, lunch from 11:45am-12:30pm)

Connecting

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.

Learning

NORD invites expert physicians in the diagnosis and treatment of aHUS to present new information and share updates to help you better understand your disease.

Sharing Hope

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.

Register Today! Contact: Susan Olivo at NORD

Phone: (203) 304-7263 Email: solivo@rarediseases.org
Online Registration: www.rarediseases.org/ahus_2016/



NORD[®]
National Organization
for Rare Disorders

August 26-27, 2016
San Francisco,
California

“ 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 aHUS. I feel that we have an instant connection. ”

— Patient from NORD's aHUS Meeting

Alone we are rare.
Together we are strong.®

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, cure or treatment of a condition or disorder.

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