

# An Educational Webinar Series for the North American Fabry Community

beginning

**Wednesday, August 11, 2010**

**7:00 - 8:00 pm EDT**

(6:00 pm CT, 4:00 pm PT)



National  
**FABRY DISEASE**  
Foundation

**How Do I Get It? How Do I Give It? What Is it?**  
*The Genetics and Inheritance of Fabry Disease*

canadian**fabry**association  
l'association canadienne de **fabry**

## Panelists will explain

- Fabry as an X-linked genetic disorder
- Different genetic mutations
- How family trees help understand disease
- Current and investigational medicines for Fabry disease

## Speakers include

- Alyson Krokosky, Genetic Counselor, Genetic Alliance
- Jack Johnson, Executive Director, Fabry Support & Information Group
- Christine Eng, M.D., Prof, Dept Molecular and Human Genetics, Baylor College of Medicine
- Moderated by NORD

## To register:

<https://www2.gotomeeting.com/register/734014962>

*Webinar 2: September 16, 2010, 7-8 pm ET*

*Webinar 3: Later in September Watch for details*

This webinar series is a collaboration of the Fabry Support & Information Group, the National Fabry Disease Foundation, the Canadian Fabry Association, Canadian Organization for Rare Disorders, National Organization for Rare Disorders and the Genetic Alliance. It is supported by an educational grant from Amicus Therapeutics Patient Advocacy.



**NORD**  
National Organization for Rare Disorders

