## Location

Chicago Marriott Lincolnshire Resort 10 Marriott Drive Lincolnshire, Illinois

#### **Accommodations**

Book your room by calling 847-634-0100 or by going online https://www.nfed.org/events/2019-family-conference/.

Room block deadline is Tuesday, June 19. Room rate is \$129 plus taxes.

#### **How To Register**

You can register online at www.nfed.org on or before Monday, June 24.

#### Fees

Includes Thursday dinner, Friday breakfast, lunch and dinner; Saturday breakfast and lunch; and Kays' Kids Camp and Teens Program activities. Each paid registration will receive a Conference T-shirt.

Adult (18 and older) - \$200 Children (birth-17) - \$125

Late Registration (Register June 25 or later) Adult (18 and older) - \$275 Children (birth-17) - \$200

Registration Cancellation Policy

If you register but are unable to attend the Family Conference, the NFED will refund your total conference registration fees less a \$30 administration fee. There will be no refund allowed on or after July 1.

### **Bev Meier's Golden Ticket Fund**

Scholarships are available to help you attend the Family Conference. Funding is limited. Apply online by March 1.

Visit www.NFED.org to learn more about all aspects of Family Conference. Or, contact Kelley Atchison at Kelley@nfed.org or 618-566-2020.

#### **Disclaimer**

The NFED Family Conference is not intended as a substitute for medical advice. Only a healthcare provider with a full medical history may determine the proper treatment options. Opinions shared in the sessions do not necessarily reflect those of the NFED. Conference topics are subject to change.

6 Executive Dr.,





Thursday, July 11 • 10 am - 7:30 pm Friday, July 12 • 6:30 am - 8 pm Saturday, July • 13 8 am - 3 pm



# **About**

The National Foundation for Ectodermal Dysplasias (NFED) Family Conference is for everyone affected by ectodermal dysplasias and their loved ones. Come and spend three full days that will enhance your life!

Our caring staff and families will welcome you to the Windy City with open arms and understanding. You will quickly experience-not just know-that you are not alone. You will develop a support network of others affected by ectodermal dysplasias and learn from their practical tips and ideas. Meet medical and dental experts and have the unique opportunity to talk with them one-on-one to ask questions. They are not just knowledgeable but compassionate. Discuss recommended treatment options and ways to cope with the emotional challenges of ectodermal dysplasias. Learn how to advocate for yourself, for your loved one and for the entire ectodermal dysplasias community. You also can share your

insights and story to inspire and help others. Stay in the know by hearing the latest research developments for your type of ectodermal dysplasia. While parents and adults are participating in sessions, Kids and teens are kept busy with activities in Kays' Kids Camp and Teen Program.

View a detailed schedule at https://www. nfed.org/events/2019-family-conference/.

## Kays' Kids Camp and Teens Program

Sponsored by Louis J. and June E. Kay

Kays' Kids Camp is available for children ages infant-12 years old. Teens Program is for children 13-17 years old.

Kays' Kids Camp is available for children ages infant-12 years-old. The Teens Program is for children 13-17 years old. Both programs include fun, age-appropriate workshops, entertainment, arts and crafts, and social time with others who are affected by ectodermal dysplasias or who have siblings that are. Teens who register for the Teens Program will participate in off-site activities.







