

Location

DoubleTree by Hilton Hotel Bloomington
Minneapolis South
7800 Normandale Boulevard
Minneapolis, Minnesota, 55439 USA

Lodging accommodations can be made on the registration page. Room block deadline is Tuesday, June 17, 2025. Room rate is \$159/night plus taxes and fees.



How to Register

Register online at
www.nfed.org/FC2025

Registration Fees Include

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

Adult (18 and older) — \$215

Children (birth-17) — \$125

Last Call Registration (Register June 1 or later)

Adult (18 and older) — \$400

Children (birth-17) — \$200

Bev Meier's Golden Ticket Scholarship Fund

Travel and lodging scholarships are available to help you attend the Family Conference. Funding is limited. Apply online at time of registration and by Monday, February 17.

Chance Auction

Back by popular demand, the Chance Auction is a fun and exciting way to raise a little money for the NFED. We are looking for donations of items for both children and adults.

Visit nfed.org/FC2025 to learn more about all aspects of Family Conference.

Disclaimer

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

Family Conference events are subject to change based on local and CDC guidelines. Our number one priority is and always will be the safety of you and your family.

nfed
NATIONAL FOUNDATION FOR
ECTODERMAL DYSPLASIAS

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Family conference 2025



Thursday, July 10 through
Saturday, July 12, 2025



**“If you do nothing else, you need to go to
the Family Conference in the summer.”**

— Parent of child affected by XLHED



About

The 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! You will quickly experience—not just know— you are not alone. At the conference, you will:

- Develop a support network of others affected by ectodermal dysplasias.
- Meet medical and dental experts and talk one-on-one.
- Discuss recommended treatment options and how to cope with the emotional challenges.
- Learn how to advocate for yourself, for your loved ones, and for the entire ectodermal dysplasias community.
- Share your story and insights to inspire and help others.
- Stay in the know by hearing the latest research developments for your type of ectodermal dysplasia.
- Have Kays’ Kids Camp and Teens Program for your children under 18 to attend.

View a detailed schedule at www.nfed.org/FC2025

Kays’ Kids Camp and Teens Program

Sponsored by the Louis J. and June E. Kay Foundation

Children ages infant - 8th grade. The Teens Program is for high school students (entering 9th grade or above in the fall of 2025). Programming includes workshops, arts and crafts, and social time with others affected by ectodermal dysplasia or who have siblings that are.

“Family Conference gave Peter a place to belong and that was really important.”

— Peter’s mom



Save Your Spot and Learn More!

www.nfed.org/FC2025



Thursday, July 10 • 2 p.m. — 9 p.m.
Friday, July 11 • 9 a.m. — 9 p.m.
Saturday, July 12 • 8 a.m. — 2 p.m.



“When I felt like I was left out throughout the school year and I felt like I couldn’t connect with my peers, I would go to the NFED conference over the summer and I would connect to these people who understood me for me and didn’t care about a skin condition.”

— Individual affected by Goltz syndrome