

The first NFED Advocacy Day made history.

Friendships and connections bloomed.

Research continues path to viable treatments and cures.



..... 175

People attended the 1st NFED Advocacy Day on Capitol Hill.



..... 473

People attended our 2017 Family Conference, the largest in NFED history.



..... 3

Toddlers continue evaluation for Dr. Holm Schneider's XLHED research project. They show signs of sweat and tooth buds.



..... 1

Bill to be introduced to U.S. Senate by Senator Tammy Baldwin to cover dental treatments for ectodermal dysplasias.



..... 308

Affected individuals registered with the NFED in 2017.



..... 19

Family members and medical experts met at the NIH campus to discuss a new and improved ectodermal dysplasias classification system.



..... 2nd

NFED Advocacy Day on Capitol Hill on July 18, 2018.



..... 23

People attended the NFED Heartland Families' social meet-up in Iowa.



..... 2

Dr. Maranke Koster continues work on two potential treatments for skin erosions and cornea transplants.



## Our supporters are remarkable.



### Donor spotlight: Becca

"I give to the NFED in loving memory of my mom, who saw her life open up after she found the NFED. With the little money she left behind, she would be elated to know that it went back to the organization that gave her a full breath again. Thank you!"

Volunteers, sponsors, and donors are all vital to the NFED community.

559 active volunteers performed 5,000 volunteer hours.

Thank you, Annual Platinum Sponsors:

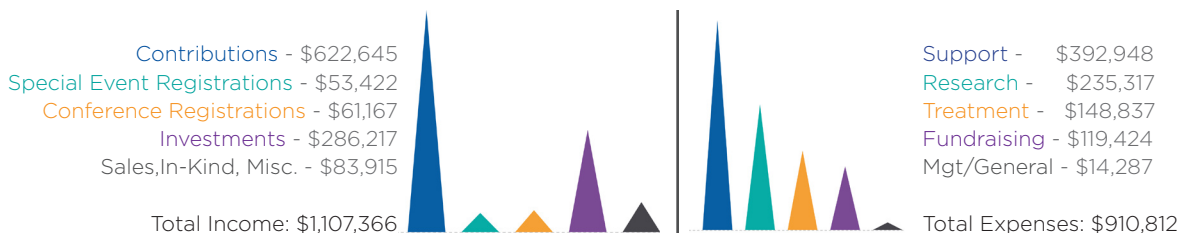
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The Hubbard  
Broadcasting Fdn  
The Louis & Gladyce Foster  
Family Fdn  
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# nfed

Supporting you. Supporting each other.

NATIONAL FOUNDATION FOR  
ECTODERMAL DYSPLASIAS

6 Executive Dr., Ste. 2  
Fairveiw Heights, IL 62208-1360



\*Visit [nfed.org](http://nfed.org) in June to see the completed audit.

### Donor trends:

194 donors increased their giving in 2017, 270 stayed the same, 565 were new donors, 344 donors came back, 37 monthly donors (13% were new), and 40 family fundraising events were held in 2017 (up from 23 in 2016).

*The mission of the NFED is to empower and connect people touched by ectodermal dysplasias through education, support and research.*