



Scan to Register



## Schedule:



Monday, Nov 3: Advocacy Training + Awareness Walk.



Tuesday, Nov 4: Full Day of Legislative Meetings.

## Lodging & Travel



Hyatt Place Washington, D.C. National Mall \$155/night  
Reservation deadline is October 10, 2025.

Questions: [becky@nfed.org](mailto:becky@nfed.org)

## Calling All Advocates! Join us on Capitol Hill!

Join families and advocates from across the country on Capitol Hill to share our stories and push for the Ensuring Lasting Smiles Act (ELSA).

**Your voice has the power to create lasting change!  
Come to make it heard!**

## Why you should attend:

- ✓ Meet face-to-face with your U.S. Senators and Representatives and/or their staff.
- ✓ Share your story or stories on behalf of your patients as to why ELSA matters to families with ectodermal dysplasias and congenital anomalies.
- ✓ NFED provides training, talking points, and schedules all meetings.
- ✓ No experience is needed!

[www.nfed.org](http://www.nfed.org)



ELSA would require a group health plan and a health insurance issuer offering group or individual health insurance coverage to provide coverage for outpatient and inpatient items and services related to the diagnosis and treatment of a congenital anomaly or birth defect that primarily impacts the appearance or function of the eyes, ears, teeth, mouth, or jaw. Ensuring access to the medically necessary items and services necessary to functionally improve, repair, or restore bodily function or approximate a normal appearance due to a congenital anomaly, that primarily impacts the appearance or function of the eyes, ears, teeth, mouth, or jaw, such as cleft lip and palate, maxillofacial abnormalities, microtia, hypodontia, and congenital aphakia.