

Ensuring Lasting Smiles Act (HR.1916, S.754 in the 117th Congress)

Please cosponsor and support the bipartisan, bicameral *Ensuring Lasting Smiles Act (ELSA)*, which will be reintroduced by Senate leads, Senator Tammy Baldwin (D-WI) and Senator Joni Ernst (R-IA), and House leads, Congresswoman Anna Eshoo (D-CA-18) and Congressman Drew Ferguson (R-GA-3), later this year for the 118th Congress.

In the 117th Congress, this bill gained strong bipartisan support from 62 U.S. Senate cosponsors, 317 U.S. House cosponsors, and passed in the U.S. House of Representatives with a 310-110 vote. In preparation for ELSA's reintroduction in the 118th Congress, the bill leads and a broad coalition of ELSA supporting organizations have worked together to incorporate feedback and received several rounds of technical assistance to strengthen ELSA's language. ELSA is endorsed by a broad coalition of 70+ national health care professional and patient advocacy organizations (see back).

Summary: This bipartisan legislation 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, including any medically necessary item or service to achieve normal body functioning and clarifies coverage for adjunctive dental, orthodontic, or prosthodontic support. *The bill explicitly excludes coverage of cosmetic surgeries.*

Background: About four percent of children in the U.S. are born with congenital anomalies that affect the way they look, develop, or function. Many born with congenital anomalies suffer from severe oral defects (such as cleft lip or palate, hypodontia, enamel hypoplasia), skeletal defects (such as craniosynostosis), vision defects (such as congenital cataracts or aphakia), hearing defects (such as microtia), or other loss of bodily functions. Individuals who do not receive timely, continuous care for their congenital anomalies face long-term physical and psychological injuries.

Problem: Most group and individual health plans include coverages for congenital anomalies, and all 50 states require health insurers to provide coverage for the treatment of congenital anomalies. Despite this, health plans systematically and routinely deny or delay claims and appeals for treatment of congenital anomalies by wrongfully categorizing certain treatments or body parts as cosmetic or not medically necessary. This is a common practice that leaves families with the burden of paying 100% of the cost for their child's medically necessary treatments, despite maintaining private health plans.

Solution: The Ensuring Lasting Smiles Act will close loopholes and ensure that group and individual health plans do not deny or delay medically necessary treatments of congenital anomalies, therefore improving the health of countless Americans.

Contact Information: To learn more or cosponsor, please contact **(Senate)** Sen. Baldwin's or Sen. Ernst's office or **(House)** Rep. Eshoo's or Rep. Ferguson's office.

Academy of General Dentistry

American Academy of Dermatology Association

American Academy of Facial Plastic and

Reconstructive Surgery

American Academy of Neurology

American Academy of Ophthalmology

American Academy of Oral and Maxillofacial

<u>Pathology</u>

American Academy of Oral and Maxillofacial

Radiology

American Academy of Pediatrics

American Academy of Pediatric Dentistry

American Association for Dental, Oral, and Craniofacial

<u>Research</u>

American Association of Oral and Maxillofacial Surgeons

American Association for Pediatric Ophthalmology and

<u>Strabismus</u>

American Association of Orthodontists

American Association of Women Dentists

American Behcet's Disease Association

American Cleft Palate-Craniofacial Association

American College of Prosthodontists

American College of Surgeons

<u>American Dental Association</u>

American Prosthodontic Society

American Society for Dermatologic Surgery Association

American Society of Dentist Anesthesiologists

<u>American Society of Maxillofacial Surgeons</u>

American Society of Pediatric Otolaryngology

American Society of Plastic Surgeons

Association of Dental Support Organizations

Barth Syndrome Foundation

Born a Hero Research Foundation

Bridge the Gap - SYNGAP Education and Research

<u>Foundation</u>

CCD Smiles

CDH International

Children's Hospital of Wisconsin

<u>Colorado Rare</u>

Costello Syndrome Family Network

Crane Dental Laboratory

Dermatology Nurses' Association

Derma Care Access Network

Ear Community, Inc.

EveryLife Foundation for Rare Diseases

Face-To-Face Colorado

FACES: The National Craniofacial Association

FD/MAS Alliance

Foundation for Ichthyosis and Related Skin Types,

Inc. (FIRST)

Genetic Alliance

Georgia Prosthodontics

Gillette Children's Specialty Healthcare

International Pemphigus and Pemphigoid

Foundation

Karen S McAndrew DMD, MS, PLC

Lawrence C. Wright Craniofacial Center at John R.

Oishei Children's Hospital

Lupus and Allied Diseases Association

Lymphedema Advocacy Group

M-CM Network

March of Dimes

Moebius Syndrome Foundation

myFace

National Association of Dental Laboratories

National Foundation for Ectodermal Dysplasias

National Organization for Rare Disorders

Noah's Hope - Hope4Bridget

Operation Smile

Ozark Prosthodontics

Pathways for Rare and Orphan Studies

Project Accessible Oral Health

Rare & Undiagnosed Network

rareLife solutions

Smile Train

Soft Bones: The US Hypophosphatasia Foundation

SunnyStrong

The APS Type 1 Foundation Inc.

The Marfan Foundation

The XLH Network, Inc.

The Sturge-Weber Foundation

Usher 1F Collaborative

Virginia Council of Nurse Practitioners

Wisconsin Speech-Language Pathology and

Audiology Association (WSHA)