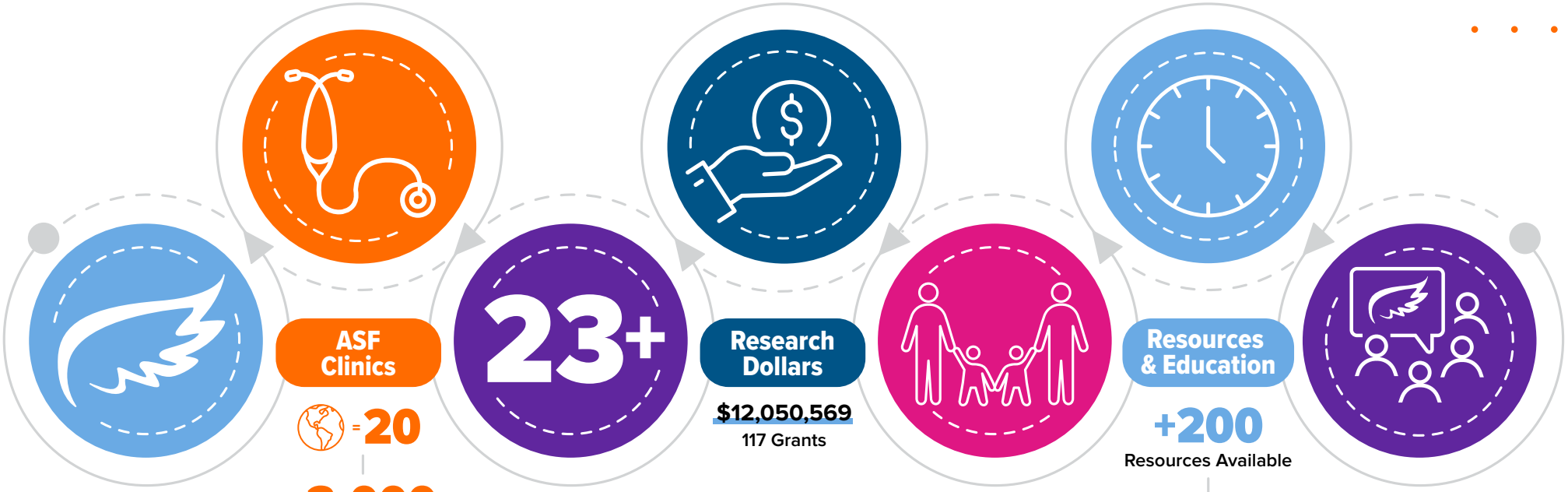
ANGELMAN SYNDROME FOUNDATION

Angelman syndrome (AS) is a rare neuro-genetic disorder that occurs in one in 15,000 live births, or 500,000 people worldwide. It is caused by a loss of function of the UBE3A gene in the 15th chromosome derived from the mother. People with AS suffer from seizures, sleep problems, lack of speech, walking and balance issues—just to name a few. In 1992, ASF was formed to provide support for the AS community, as well as fund vital research to find better treatments and (ultimately) a cure.



1992
Angelman Syndrome Foundation was formed

1996
First Clinical Trial funded

ASF Clinics
= 20

+3,000
Patients Seen

23+
Therapeutic Approaches in the Pipeline

Research Dollars
$12,050,569
117 Grants

ASF Family Fund

2019
ASF Launched The Family Fund

$393,190
Total Awarded (348 Awards)

Resources & Education

+200
Resources Available

+3,000
Hours of One-On-One Support

+211,000
Views on Educational & Training Videos

Family Conferences & Research Symposia

+10k
Attendees Since 1993

angelman SYNDROME FOUNDATION

The mission of Angelman Syndrome Foundation is to advance the awareness and treatment of Angelman syndrome through education and information, research and support for individuals with Angelman syndrome, their families and other concerned parties. We exist to give all of them a reason to smile, with the ultimate goal of finding a cure.

**as of March 2022*

Angelman.org