



The mission of the 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.

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- Occurs in one in 15,000 live births.
- It is caused by a problem with the UBE3A gene located on the 15th chromosome.
- Symptoms include developmental delays, seizures, inability to speak & sleep problems.
- Individuals with AS are known for their happy demeanor.

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