sat b2 gene foundation





The SATB2 Gene Foundation, Inc. was established to enrich the lives of individuals with SATB2-Associated Syndrome (SAS) through support, research, and education.

♥ FACTS ABOUT SAS ♥

- De novo genetic condition, also known as Glass Syndrome, that results in an alteration (i.e., misspelling, deletion, duplication) of the SATB2 Gene
- Because the SATB2 protein plays an important role in brain and facial development, impacts of the syndrome can include:
  - Global developmental delay, including severe speech and language impairment
  - Intellectual disability
  - Craniofacial and teeth abnormalities
  - Sleep and behavioral difficulties
  - Medical complications, including the potential for seizures and low bone density

FOR FAMILY AND FRIENDS

VISIT WWW.SATB2GENE.ORG
FOLLOW US ON
CLOSED FACEBOOK GROUP: SATB2 SYNDROME (2Q33.1)

750 SATB2 Individuals
50 Countries

FOR RESEARCHERS

VISIT WWW.SATB2GENE.ORG/RESEARCH FOR SCIENTIFIC AND RESEARCH INFORMATION

"Our Smiles Make The World Go Round"

"It has given me a place to find peace in our journey and know that we are not alone. I'm grateful for the community of families that understand."
– Danielle Duran
Mom to Dane

Our Impact

"I'LL NEVER FORGET WALKING IN TO OUR VERY FIRST FAMILY CONFERENCE AND REALIZING THESE WERE MY PEOPLE! THEY UNDERSTOOD THE HARD. THEY UNDERSTOOD THE FRUSTRATIONS. THEY GOT IT. AND I HOPE THAT THOSE OF US THAT HAVE "BEEN AROUND" CAN CONTINUE TO COME ALONGSIDE THOSE NEW TO THE DIAGNOSIS AND OFFER HOPE."
– ALICE WOLFE
MOM TO PAUL AND WILL

"HAVING A COMMUNITY THAT NOT ONLY KNOWS WHAT YOU'RE GOING THROUGH, KNOWS YOUR STRUGGLES, CAN CELEBRATE THE SMALL MILESTONES, AND CAN OFFER ADVICE AND SUPPORT HAS BEEN AMAZING! I AM SO FORTUNATE TO HAVE HAD THE EXPERIENCE OF ATTENDING NOT ONLY THE SATB2 CLINIC BUT THE FAMILY CONFERENCE AND AM SO THANKFUL FOR THIS COMMUNITY."
– RACHEL WARD
MOM TO LINCOLN

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