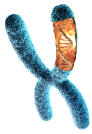
CTNNB1
CONNECT AND CURE

Accelerating Research | Raising Awareness | Connecting Families

#1 Genetic Cause of Cerebral Palsy



What is CTNNB1 Syndrome?

A rare disease caused by variants of the CTNNB1 gene that result in decreased levels and/or function of the beta-catenin protein.

Estimated Prevalence:

3 in 100,000

Symptoms Can Include

- Cognitive impairment
- Microcephaly
- Behavioral challenges
- Sleep disturbances
- Limited speech / nonspeaking
- Epilepsy

- Truncal hypotonia
- Peripheral spasticity
- Dystonia

- Exudative vitreoretinopathy
- Strabismus
- Refractive errors

- Tethered spinal cord
- Congenital heart defects
- Osteopenia
- Scoliosis
- Feeding difficulties
- Gastrointestinal problems

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