Mission

CTNNB1 Connect and Cure is a 501(c)(3) nonprofit organization dedicated to finding treatment options and a cure for CTNNB1 Syndrome while improving the lives of our patients and community.

Vision

To create a world where every family affected by CTNNB1 Syndrome has access to safe and effective treatments and a supportive community.

Core Values

Transparency
Teamwork
Integrity

www.curectnnb1.org

info@curectnnb1.org

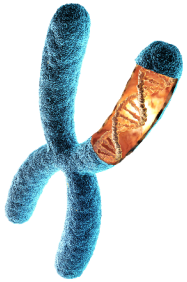
CTNNB1 Connect and Cure, Inc.
EIN: 83-4541448

CTNNB1
CONNECT AND CURE

Accelerating *Research*
Connecting *Families*
Raising *Awareness*

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