



Meet Moriah

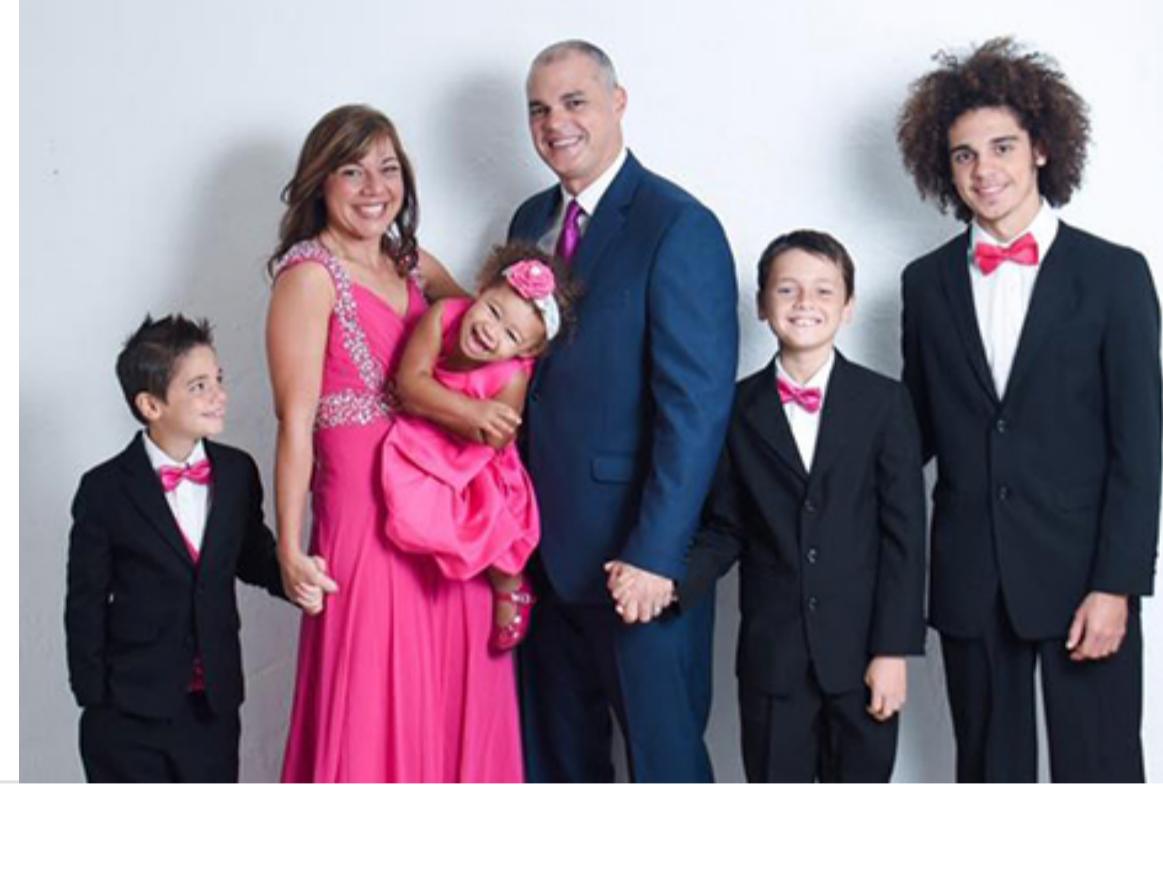
Moriah Joy was diagnosed with a rare chromosome abnormality when she was 16 months old. Xtraordinary Joy was founded to provide advocacy and funding for X chromosome deletion research that advances lifelong health, behavior and learning for

[DONATE](#)

Our Joy

In 2014 we adopted Moriah Joy. Our story is one of patience, joy and unexpected discovery.

Established in 2016, our vision for Xtraordinary Joy, Inc. is to raise funds that will promote clinical interventions, therapies and care opportunities for children afflicted by X chromosome deletions.

[LEARN MORE](#)

Our Next Event

Join us for our 2nd Human Foosball Event!

When: April 9th

Where: Trinity United Methodist Church

Find out how to be an [X Caliber sponsor here.](#)

[SIGN UP](#)

Research & Resources

XQ27.3-Q28 DELETION

Fragile X Syndrome
fragilex.org

Hunter Syndrome
mpssociety.org/mps/mps-ii/

POST DIAGNOSIS

National Organization for Rare Disorders NORD
rarediseases.org

Unique
rarechromo.org

Chromosome Disorder Outreach
chromodisorder.org

SCIENTIFIC RESEARCH PLAN

Center for Cellular Reprogramming
ccr.med.ufl.edu

Department of Neuroscience
neuroscience.ufl.edu