#### Compassion, Connection, Community.

Stronger, together. Since 1996.



### Let's Connect



@Chromosome22Central



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Chromosome 22 Central



c22central.blogspot.com/



Chromosome 22 Central





C22C is a global collection of parents and people affected by chromosome 22 disorders who connect, advocate and grow, together. Our mission to is to spread awareness, and offer support and information on all chromosome 22 disorders.

### **Contact Us**

Website www.c22c.org

#### Address

1129 Carolina Gardens Ave., Fuquay-Varina, North Carolina, 27526 USA

**Email** c22central@gmail.com

**Phone** 919-762-7979



## Chromosome 22 Central

# Support for disorders of chromosome 22

A 501(c)(3) Corporation (USA)

www.c22c.org

## What else is C22C about?

- We maintain a website that allows people to find others.
- We help researchers by posting their studies.
- We mail out parent guides on Emanuel syndrome, at no cost to families.
- We have held family conferences and plan for more in the future.
- We ensure our group is easy to find so families can find the support they need by registering with other larger umbrella organizations and connecting with other groups like ours.
- We try to fill the gaps where we find there isn't enough support or information for families.
- Our families come together to find connections around the world.



#### SUPPORTING ALL CHROMOSOME 22 DISORDERS

- 🖄 22q11.2 Deletion
- 済 (Velocardiofacial and DiGeorge
  - Syndrome)
- 22q11 Microduplication Syndrome
- 🖄 Emanuel Syndrome
- $\stackrel{\scriptstyle <}{\searrow}$  11/22 translocation
- 🖄 Cat Eye Syndrome
- ₩ Mosaic Trisomy 22
- K Complete Trisomy 22
- 🖄 Phelan-McDermid Syndrome
- 🖄 (22q13 deletion, Ring 22)
- Unique chromosome 22 conditions.

## More about us

Our website is a hub for people to find immediate connections with others through our various social media platforms, and to find basic information and links to other resources.

Our existence allows families to feel part of a community so they don't feel alone on this journey.

We work hard to spread awareness of chromosome 22 disorders.

We offer comfort in the support of each other, validation, strength and community.

We raise funds to support activities that are important to us, but we don't charge membership fees.

We are thousands of members in over 40 countries, and waiting to welcome you to the family.

