

EVENT SPONSORSHIP

Imagine receiving a diagnosis of a rare and life-threatening disease that affects around 200 individuals worldwide.

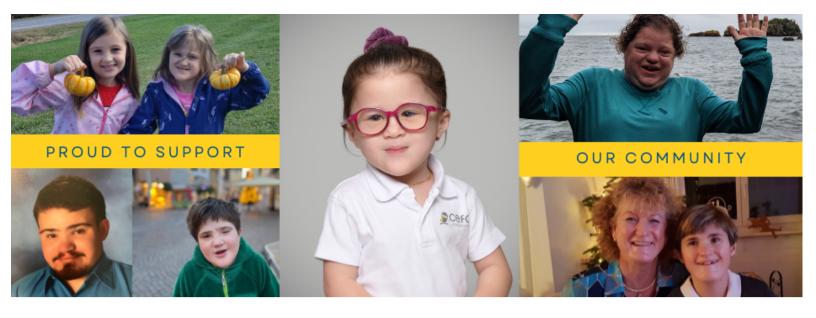
Envision a reality where you've never crossed paths with someone who truly comprehends your challenges.

With your support, we will hold the inaugural Myhre Syndrome Foundation conference, which will unite individuals affected by Myhre syndrome.









ABOUT THE EVENT

We'd love for your business to partner with us and benefit from the following:

- Logo placement across all event comms
- A newsletter takeover in the month of your choice
- · Social media shout-outs
- The love and thanks from a community whose lives you will be changing.

You can be a partial or full sponsor of the following:

The Art Room - \$2,750

During the conference, we'll have a team of childcare professionals supporting the needs of our young Myhre community. It means parents and caregivers have the time to attend the conference, knowing that, just next door, their children and young adults are safe and engaged in creating an art piece they get to take home and cherish.

Family Travel Fund - \$5,000

Can you help us bring families to the conference who otherwise couldn't attend? We're supporting families with travel and hotel expenditures so they can meet Myhre doctors for the first time, which will significantly impact the quality of care they, or their child, receive.

The Celebratory Dinner - \$10,000

Following a full day of learning and fun, we want all our families to relax and enjoy a wonderful evening of celebration. We'd love to treat our families to a 3-course dinner, with dancing and entertainment that will put smiles on the faces of everyone.

Please join us as a sponsor by December 2023 so you can benefit from all of the event communications and the wonderful build-up to this special event.

MYHRE CONFERENCE & DINNER

Saturday, 27 July, 2024

Expert talks from the international Myhre specialists followed by a celebratory dinner.

DOCTOR MEET & GREET

Sunday, 28 July, 2024

The chance to sit down with doctors and ask questions about Myhre syndrome.