



The Bow Foundation

2022 YEAR IN REVIEW



THANK YOU!

As we close 2022, we want to send a special note to thank you again for your continued support of the Bow Foundation. Since inception in 2017, the Bow Foundation has raised over \$1.1 million to advance our mission of building a better tomorrow for GNAO1 patients and their families. More than 95% of that funding has gone directly to medical research into the rare disease.

As a volunteer-run Foundation, our sole focus is building a better tomorrow for GNAO1 patients and their families. Thank you again for your ongoing support of our mission. We're looking forward to 2023 and know that none of this would be possible without you.

Thank you for believing in our mission!

Sincerely,

The Bow Foundation Team
Emily and Stephen Bell
Alice and Duke Fox

2022 ACCOMPLISHMENTS

- Continued funding four revolutionary research projects focused on expanding our knowledge of GNAO1. Read more about these exciting projects here.

- Hosted the GNAO1 Medical and Family Conference in June in St. Louis, Missouri.

More than 25 families connected, heard research updates and received advice from experts on topics including caregiver wellness, living a more inclusive life and more. If you're interested, click below to watch the conference recording.

- Organized a virtual GNAO1 International Conference focused on connecting medical experts around the globe and identifying common objectives and opportunities to advance shared research into the rare disease.

More than 50 medical researchers and clinicians participated in the conference and roundtable. Experts from the Netherlands, Italy, Switzerland, Israel & the USA led the presentations. If you missed those presentations, you can watch a recording in English, Italian, or Spanish by clicking the link below.

- Elevated GNAO1 awareness with our fourth annual GNAO1 International Awareness Day on October 1, 2022, raising an additional \$50,000+ to support more GNAO1 research.



emily@bowfoundation.org



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www.gnao1.org