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What can we do for you?

Earlier this year, the FamilieSCN2A Foundation began building a 3-year strategic plan. We have identified a range of goals that will further our mission to accelerate research, build community, and advocate to improve the lives of those affected by SCN2A-related disorders around the world.

Now we need to hear from you!

Please take 15 minutes to rank YOUR priorities for the Foundation by taking the survey below. There are no wrong answers! We exist to fulfill the vision of a world with effective treatments and cures for all SCN2A-related disorders. We can only do that by serving you, our patient/family community. **Your input is invaluable.**

In the spirit of our core values of URGENCY, INTEGRITY, COLLABORATION, and INCLUSION we hope that every one of you will take the survey and provide any additional feedback in a timely manner.

Last day to complete survey is Tuesday, JULY 19. We appreciate all of the responses received so far. Your voice matters!

While the survey does ask for an email address, it is only to verify that you are a member of the community and to avoid duplicate entries. Your answers will remain anonymous.

TAKE THE SURVEY



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