

Participants Needed for Consensus Group

We are looking for parents of children or adolescents with congenital heart disease (CHD), and for youths (13-25 years old) with CHD to participate in the elaboration of guidelines to improve the developmental follow-up of children and adolescents with CHD.

What is the study about?

The purpose of this study is to develop recommendations to improve the identification of developmental delays in children and adolescents with CHD across Canada.

What's involved?

Your participation in this research study involves participating in one online consensus group that will last approximately 3 hours. During the consensus group, you will be asked to generate and select recommendations on specific topics based on the information that was presented and your knowledge and experience of the Canadian health care system.

Are there benefits to participating?

You may not directly benefit from participating in this research. However, we hope that the study results will contribute to improving developmental follow-up for children and adolescents with CHDs.

Participants will be given a gift certificate in recognition of their contribution.

Study Title: BRIGHT Pathways: Recommendations to Improve the Developmental Follow-up of Children with CHD in Canada

Principal Investigator: Dr. Mike Seed

Version 2: 30-05-2025

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**If you are interested in participating or have questions on the study,
contact us at bright.pathways@sickkids.ca (416-813-7230)
or scan the QR code.**

