



HAVE A VOICE AND
**STRENGTHEN CANADA'S
FASD COMMUNITY**

CANFASD IS CANADA'S ONLY NATIONAL FETAL
ALCOHOL SPECTRUM DISORDER RESEARCH NETWORK.



BECOMING AN INDIVIDUAL MEMBER* OF THE CANADA FETAL ALCOHOL SPECTRUM DISORDER RESEARCH NETWORK is a way to have your voice heard, meet others who share your interest and strengthen Canada's FASD community.

Membership is open to anyone including people with FASD, service providers, researchers, professionals and caregivers of people with FASD.



BE HEARD AND SHARE STORIES

Families, caregivers and individuals share stories, compare experiences and find common ground. Researchers learn from individuals and families, while families and individuals stay updated on FASD research. Every story helps build a stronger FASD community across Canada.



CONNECT WITH A COMMUNITY OF SHARED INTERESTS

CanFASD members connect with others across Canada to ask questions, voice concerns and discuss ideas. Members are part of the broader FASD community from coast to coast.



INFLUENCE AND CONTRIBUTE TO FASD RESEARCH

Addressing the complexities of FASD needs ongoing research. Individual CanFASD members support research through membership fees. Members can also take part in research studies, influence policy and comment on plans and issues.



BE THE FIRST TO KNOW AND STAY IN THE KNOW

CanFASD members are first in line to receive newsletters, research summaries, event notifications, stories of interest and project updates. Individual members have easy access to credible, trusted information about FASD.

* Individual CanFASD memberships differ from Jurisdictional Memberships that are offered exclusively to Canadian provinces and territories. Offering memberships to individuals ensures that Canadians from coast to coast to coast can be part of a broader FASD community.



DISCOVER ALL THE BENEFITS OF A CANFASD INDIVIDUAL MEMBERSHIP

1 STAY CONNECTED

- Being a part of the Canada FASD Research Network.
- Invitations to in-person member-only events, such as meet and greets and town hall meetings where you can connect with CanFASD researchers, board members, staff and members from our advisory committees.
- Member-only question and answer sessions with CanFASD staff and researchers.
- First-in-line notices for upcoming events and important announcements.
- A complimentary listing in the CanFASD Membership Directory*.

*This benefit is available exclusively to members who join as a researcher, service providers or professional.

2 SHARE YOUR STORIES

- Opportunities for researchers to seek research participants through CanFASD networks.
- Opportunities for researchers to have their work profiled in the e-newsletter and on CanFASD channels.
- Opportunities for families and service providers to share their stories through our e-newsletter and CanFASD channels.

3 STAY INFORMED

- Bi-weekly e-newsletter updates with exclusive information, including research summaries, invitations to participate in research projects, member-only event notification, professional development opportunities, stories of interest, and more.
- Invitations to member-only webinars that put evidence about FASD in real-world contexts.
- Ongoing research project updates that offer real-time information, not just after the research is published.

4 INFLUENCE RESEARCH

- Opportunities to participate in and contribute to research studies to have your voice heard.
- Member-only surveys that help influence the direction of policy related to FASD.
- Opportunity to provide annual feedback into CanFASD's research agenda and issues of importance.
- Researchers, students, and service providers are eligible to apply for CanFASD research and travel awards.



HAVE A VOICE | STAY CONNECTED

BECOME A CANFASD INDIVIDUAL MEMBER TODAY!

JOIN A COMMUNITY OF CANADIANS FROM COAST TO COAST TO COAST
WITH AN INTEREST IN FASD. VISIT OUR WEBSITE TO FIND OUT MORE.

PROFESSIONAL	PERSONAL
<p>FOR RESEARCHERS, SERVICE PROVIDERS AND PROFESSIONALS</p> <p>Includes a listing in the CanFASD Membership Directory</p> <p>\$129 PER YEAR</p>	<p>FOR INDIVIDUALS AND CAREGIVERS</p> <p>Our preferred rate for people with FASD and caregivers of people with FASD.</p> <p>\$49 PER YEAR</p>