



FRAGILE X RESEARCH
FOUNDATION OF CANADA

The FXRFC is currently seeking new members for the Board of Directors. Experience in communications, fundraising, scientific research, and/or IT expertise are sought after skills. Knowledge of Fragile X syndrome is essential. This is a volunteer position and new Board members can be located anywhere within Canada since the Board of Directors meet electronically.

Please send an expression of interest together with a brief statement as to qualifications to fxrfc@on.aibn.com. The deadline for receipt of applications is **Friday November 20, 2020**.

More about us:

The Fragile X Research Foundation of Canada (FXRFC) is a non-profit, tax-exempt charity run by parents and volunteer professionals that is dedicated to raising awareness of Fragile X syndrome, funding biomedical research for improved treatment and ultimately, finding a cure for this disorder.

Fragile X research is seriously underfunded. In 1997, a group of parents and professionals decided to do something about it. Understanding both the suffering that Fragile X syndrome can cause and, more importantly, the many reasons for optimism and hope through advances in research, they founded the Fragile X Research Foundation of Canada (FXRFC).

- FXRFC is a national non-profit, tax-exempt organization administered by volunteers, all of whom are parents and health professionals.
- FXRFC directly funds promising research aimed at treatment. Applications for research grants and postdoctoral fellowships are accepted on an ongoing basis and reviewed annually.
- FXRFC promotes awareness of Fragile X. We produce a regular newsletter and information materials. We hold fundraising events and organize parent advocacy groups across Canada.