

Canadian **PANS** Registry

Who can join the registry?

Any child or adult in Canada with a diagnosed or suspected case of PANS or PANDAS.

A photograph of a family of five (two adults and three children) running away from the camera through a field of tall grass at sunset. They are holding hands and their silhouettes are backlit by the warm, golden light of the setting sun. The sky is filled with soft, orange-hued clouds.

www.inflamedbrain.org/pans-registry

Share your story, improve **PANS PANDAS** care.

What does the Canadian PANS Registry study involve?

This study will start with screening questions on the homepage, followed by consent and registration. Participants will complete questionnaires on key aspects of their or their child's health, including demographics, symptoms, treatments, access to care, financial impact, and research priorities.

Why participate and what research will be done?

By sharing your or your child's story, you contribute to advancing care standards, understanding quality of life impacts, and fostering new therapies for PANS and PANDAS. Your participation supports the Inflamed Brain Alliance's advocacy efforts, clinical research, and the development of trials and treatments in Canada.



Contact: pansregistry@inflamedbrain.org



National ethics ID: 2025-05-003 | HREBA ID: HREBA.CHC-25-0006

Canadian PANS Registry | Principal Investigator: Mary Noseworthy | January 9, 2025