ATTACHMENT 3



GBS/CIDP Foundation of Canada

Guillain-Barré Syndrome/Chronic Inflammatory Demyelinating Polyneuropathy Support, Education, Research, Advocacy

Honorary Board

Larry Brenneman (deceased)
Tom Feasby, MD
Angelika Hahn, MD
Susan Keast, Founder
Serge Payer
Kenneth Shonk, MD

Executive Director

Donna Hartlen

Officers

Darryl Bedford
President
Pamela Stoikopoulos
Vice President (Interim)
Ayman Kafall
Treasurer

Board of Directors

Darryl Bedford
Holly Helm
Alexandre Grant
Ayman Kafal
Holly Longstaff
Akash Purewal
Pamela Stoikopoulos
Demetrios Strongolos
Ron Van Holst

Medical Advisory Board

Steven Baker, MD Katherine Beadon, MD Timothy Benstead, MD Pierre Bourque, MD Vera Bril, MD Colin Chalk, MD Kristine Chapman, MD Angela Genge, MD Hans Katzberg, MD Kurt Kimpinski, MD Sandrine Larue, MD Rami Massie, MD Elizabeth Pringle, MD Zaeem Siddigi, MD Jiri Vajsar, MD Chris White, MD Douglas Zochodne, MD February 13, 2025

Office of the Mayor The of City Vaughan 2141 Major Mackenzie Dr. Vaughan, ON L6A 1P7

Dear Mayor Steven Del Duca:

We are writing to respectfully request that the city of Vaughan proclaim May 2025 as GBS/CIDP Awareness Month. The Month of May, has been internationally designated as "GBS and CIDP Awareness Month" to educate the public and to focus attention on these rare conditions. The GBS/CIDP Foundation of Canada is a national, not for profit patient organization that supports patients and families afflicted with Guillain-Barré Syndrome, Chronic Inflammatory Demyelinating Polyneuropathy, and variants such as Multifocal Motor Neuropathy. We serve patients through support, education, research and advocacy.

GBS has a sudden onset of symptoms, which can cause complete paralysis within a day. Recovery can be unpredictable, patients and their families face an uncertain future, usually requiring months of hospital care without knowing if or when they will recover, or whether they will face long-term disabilities. Earlier diagnosis, treatment, and access to rehabilitation services can improve the chances of avoiding permanent lifelong residual damage of the nerves. The cause of these conditions is unknown, and can develop in any person, regardless of age, gender or ethnic background

We are working to raise awareness of these rare conditions so that future patients get help sooner and will know that our organization is here to provide hope. We provide support and information through trained volunteers, our website, materials provided to hospitals, and organize local and online peer-to- peer support group meetings. We provide patient educational events and build awareness within the medical community. We advocate for access to diagnosis and appropriate treatment, and also support Canadian research that aims to improve the quality of life of GBS, CIDP, and variant patients.

While considered rare diseases, in Canada GBS affects 2 in 100,000, and in Zika outbreaks 9-24 in 100,000. CIDP effects 5-7 in 100,000. For more information: www.gbscidp.ca Please let us know if there is anything further you require for our proclamation request to be approved for this year and for subsequent years.

Please see the attached email or application from Nancy Galaski. Nancy is an Ontario liaison her husband is a GBS survivor. We currently support 8 patients in Vaughan.

Thank you for your consideration.

Kim Brooks

Patient Advocate & Volunteer Coordinator

kbrooks@gbscidp.ca 1-403-510-3170

www.gbscidp.ca

Canadian charity registration number: 887327906RR0001