



CLLPAG UPDATE - ISSUE 6

My CLL Compass

We're pleased to announce the launch of My CLL Compass, a new online tool that helps Canadian patients and caregivers get the information they need and help alleviate anxiety or confusion they may be feeling after a chronic lymphocytic leukemia (CLL) diagnosis. Check out www.cllpag.ca for more information and media interviews with CLLPAG Board Director Rebecca Drotos.

Visit **My CLL Compass** and take the quiz today: www.mycllcompass.ca

Get Involved in CLLPAG

CLLPAG has openings on the Board of Directors for a representative from BC, Quebec and the Maritime provinces. Patients, caregivers and interested persons are welcome to apply. For more information and an application form contact Deborah at :

cllpag.canada@gmail.com . ***CLLPAG is a volunteer-run patient organization that advocates for and provides education about health care that will extend the lives of Canadians affected by Chronic Lymphocytic Leukemia (CLL) and Small Lymphocytic Lymphoma (SLL).*** Directors are expected to participate in 4-5 conference call meetings and one face to face meeting per year. It's an opportunity for you to make a difference for CLL patients in Canada.

Thank you to all who completed our recent patient and caregiver surveys, results of some of the questions are shared on our website www.cllpag.ca . Your completion of the surveys has helped CLL Advocacy groups in Canada, UK and Czechoslovakia provide input to access new CLL treatments.

Check out **RECOMMENDED CLL RESOURCES** at www.cllpag.ca

Have a healthy and safe summer

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