



CLL SOCIETY

Smart Patients Get Smart Care™

Brian Koffman, MDCM
Co-Founder, Executive Vice
President, Chief Medical Officer

Patricia Koffman
Co-Founder, Communications
Director

Carly Boos, MEd, MBA
Executive Director

Liza Avruch, MPA
Program Director

Terry Evans
Director, CLL Support Network

Brad Hedrick, MS
Director, Information Technology

Nancy Day
Program Coordinator

Medical Advisory Board

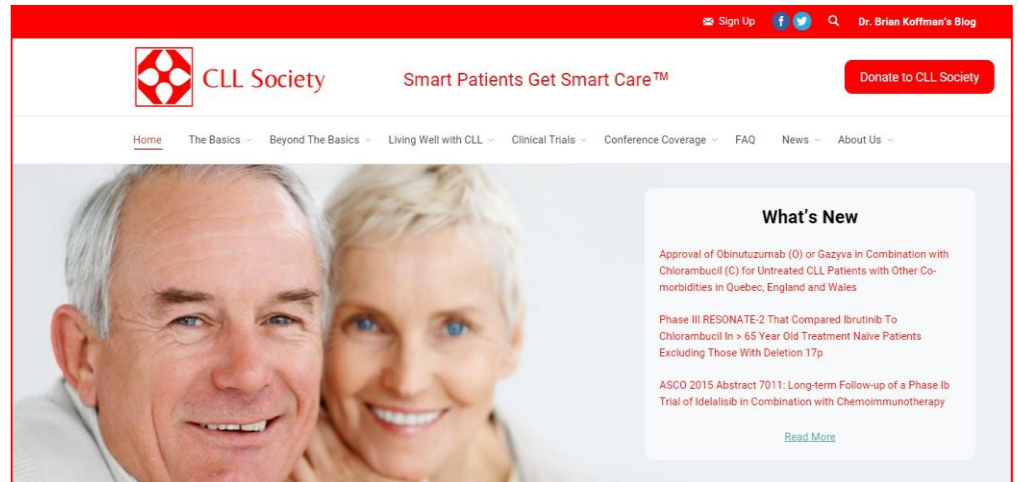
Farrukh T. Awan, MD
Danielle M. Brander, MD
John Byrd, MD
Alexey V. Danilov, MD, PhD
Matthew S. Davids, MD, MMSc
Ian Flinn, MD, PhD
Richard Furman, MD
Thomas Henry III, MBA, RPh, CPh
Brian Hill, MD, PhD
Peter Hillmen, PhD, MB
Kaitlin Kennard, BSN, RN
Thomas Kipps, MD, PhD
Brian Koffman, MDCM, MS ED
Nicole Lamanna, MD
Susan J. Leclair, PhD, CLS (NCA)
Anthony Mato, MD, MSCE
Susan M. O'Brien, MD
John M. Pagel, MD, PhD
Sameer Parikh, MBBS, MD
Guilherme Perini, MD
Lindsey Roeker, MD
Steven T. Rosen, MD
John Seymour, MB, PhD
Mayzar Shadman, MD, MPH
Sanjay Sharma, MD
Tanya Siddiqi, MD
Alan Skarbnik, MD
Kostas Stamatopoulos, MD, PhD
Deborah Stephens, DO
Chris Stewart
William G. Wierda, MD, PhD
Adrian Wiestner, MD, PhD

Patient Advisory Board

Patient advocacy and education
leaders from around the world,
representing diverse perspectives
and experiences with CLL.

CLL Society

The CLL Society Inc. is a patient–centric, physician–curated, nonprofit 501c3 focused on education, research, advocacy and support for patients with chronic lymphocytic leukemia (CLL). Dedicated to addressing the unmet needs of the CLL and related blood cancer communities, we develop real, recent and relevant information and are the “go to place” for CLL patients and caregivers.



CLL Society Website: This is a place where patients, caregivers, and HCPs find relevant, reliable information, in patient-friendly language. Includes fundamental information, deep-dives into specific topics, video interviews with top CLL researchers, real-world advice, clinical trial information, and much more.

The CLL Tribune: Our Quarterly online newsletter contains features such as *Ask the Doctor*, *Ask the Pharmacist*, and *Ask the Laboratory Scientist*, as well as first-hand accounts of patient and caregiver experiences.

Patient Support Groups: The CLL Society has created or supported over 30 CLL-specific support groups across the US and Canada, where patients and their caregivers learn and share in a supportive and educational peer-to-peer environment.

Patient & Caregiver Educational Forums: In-depth, half-day forums in 12 locations across the country annually.

CLL Toolkit: The CLL Toolkit is a binder-formatted collection of materials which provides hematologists and other HCPs with a broad range of CLL-specific educational materials for distribution to patients, promoting well-timed learning, dialogue, and patient empowerment.

Expert Access™ Program: CLL patients can connect with a CLL expert physician via video consultation in this HIPAA compliant, second opinion program.

CLL Webinars: CLL Society offers quarterly webinars series with the theme *Living Well with CLL*. Audience interaction is encouraged with polling and audience Q&A.

Test Before Treat™: CLL Society's Test Before Treat Campaign raises awareness of the importance of testing before *each* treatment, to facilitate patient-HCP dialogue to empower patients to insist on testing. Resources include a one-page informational sheet about testing and red, double-sided Test Before Treat wristbands that serve as a reminder.

Partnerships: In order to avoid a duplication of efforts and to ensure maximum impact of its work, the CLL Society collaborates with other lymphoma and leukemia organizations around the world.

Visit **CLLSociety.org** for reliable information and support.