# 2020 Annual Report To Community





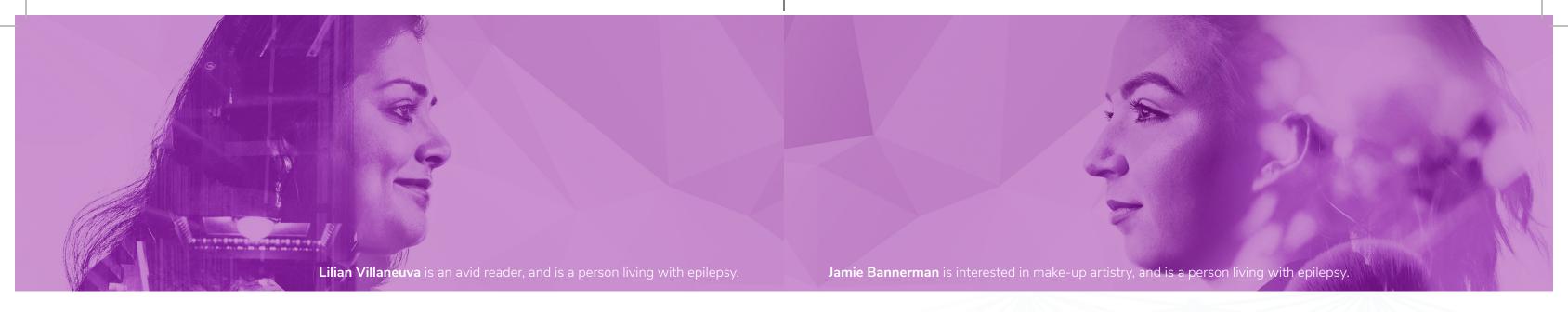


Martin Bell, Director
Cameron Browning, Treasurer
Jessica Chawrun, Director
Brad Cooper, Director
Geoff Hughes, Vice Chair
Marion Jerome, Secretary
Evan Legate, Chair
Mitch McLeod, Past Chair
David Sereda, Director

Wendy Tynan, Director

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#### A MESSAGE FROM EVAN LEGATE



Over the last year, the Board of Directors invested heavily, both financially and timewise, into building for the future of the Epilepsy Association of Calgary.

This financial investment included engaging an executive search firm in order to find the best candidate possible to lead our association forward and implement the necessary changes that we owe both to our membership and to the wider epilepsy community.

After an extensive search, we were thrilled to welcome Laura Dickson as our new Executive Director in December 2019. Laura brought over a decade of experience in leadership roles in the non-profit sector and a stellar track record of building the

capacity of other non-profits. Since joining the association, Laura has already begun to deliver, and she continues to provide strong leadership for our association as we navigate through the COVID-19 crisis.

In February 2020, the Board of Directors held a retreat to define our strategic priorities in order to provide a strong foundation for the association to build upon in the years ahead. At this retreat, the Board of Directors identified and created goals for the following five priorities: redefining a strategic direction, building a strong network of partners, creating meaningful programming, strengthening our finances, and raising awareness.

We are already seeing early results on our efforts to advance these five priorities, and as we head in 2020-21, I am confident that the initial investments made over this last year will pay significant dividends for both our membership and the wider epilepsy community as we look to become a leading epilepsy association within Canada.

Thank you for your support.

Sincerely,

Evan Legate Chair of the Board of Directors

#### **GREETINGS FROM THE EXECUTIVE DIRECTOR**



It is hard to believe that a year has now passed since I first began to explore the role of Executive Director with Epilepsy Association of Calgary. Reflecting on these past months, I am inspired by the kindness, generosity and commitment of the Association's network here in Calgary and across Southern Alberta.

I've met those whose lives are affected by epilepsy and listened to the incredible ways in which people have adapted and to their heart-felt desire to help others by sharing their stories or by simply providing a listening ear tuned in just the right way to another person's fears and worries.

I have connected with medical professionals working in clinics and in community who want to see more opportunities for patients to receive support, information and access to a network of peers in the community as a path to better health outcomes. And, I am looking forward to building upon relationships with Associations just like ourselves across Canada to see how we can work together to better raise awareness, and begin to change the conversation around epilepsy.

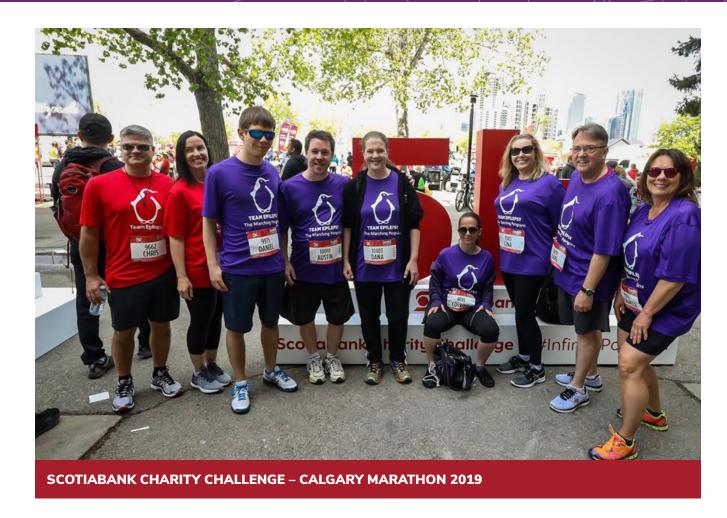
Of course, balancing operations through a global pandemic accompanied by an economic downturn are challenges shared across the non-profit community - and beyond. I am grateful to our donors, grantors and supporters for their belief in, and ongoing commitment to Epilepsy Association of Calgary. As we embark on a journey in 2020-21 of determining how our programs and services can (and should) adapt to better meet the needs of those we serve, I look forward to welcoming new members of the community to the conversation.

Regards,

Laura Dickson
Executive Director

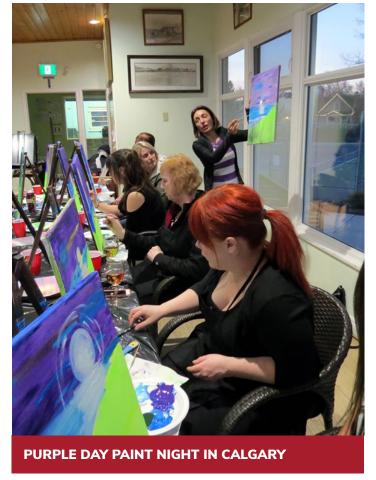
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# **YEAR IN PHOTOS**

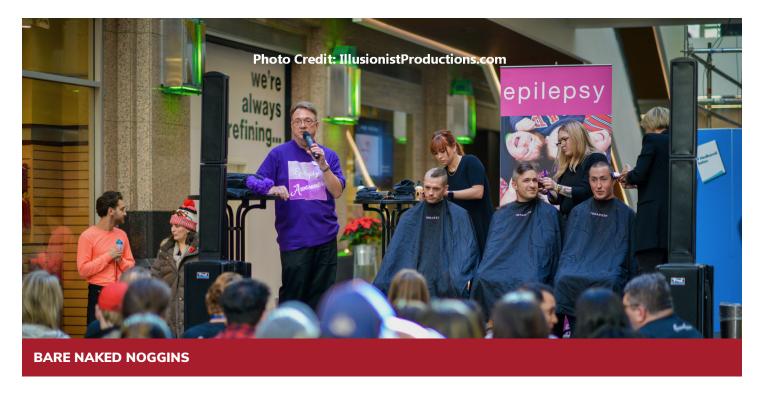




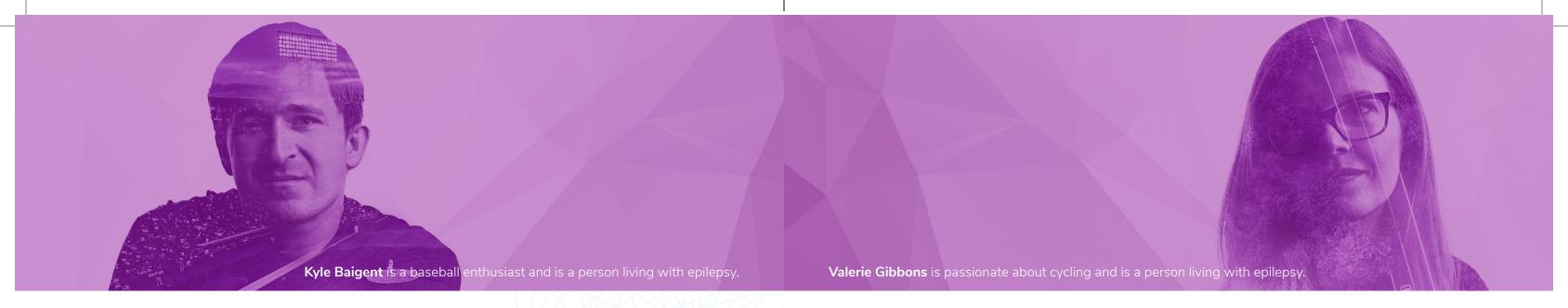








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### **JACK LEARNS OF HIS DIAGNOSIS**

Jack's epilepsy diagnosis in his late 40s brought a huge adjustment to his life; his driver's license was suspended, and he was unable to work for several months. His neurologists tried various medications to control his tonic-clonic, absence, and focal impaired awareness seizures. The medications had varying effects, all of which made his life even more difficult.

After Jack and his wife found Epilepsy Calgary's website, they began attending an adult support group where he found much-needed support in dealing with his condition and discovered that others had gone through much the same experiences. He learned how to communicate with his doctors so that his questions and concerns were heard and addressed. EAC's Support Coordinator also assisted Jack by writing a letter to CPP in support of his application for disability benefits.

Now, years later, Jack's seizures are under control and Jack still attends the support groups occasionally to offer support to others who are going through what he did when first diagnosed. Jack offers insight in terms of medications and advocating for yourself with your health care team, as well as navigating the confusing systems of medical and disability benefits. Looking back, Jack describes these groups as his lifeline during a difficult period in his life, and now sees them as a way for him to help others.

### **JOAN MAKES A DIFFICULT DECISION**

Joan, a 67-year old woman who had lived with epilepsy all her life, managing with the help of friends and family, had remained fairly independent despite her seizures.

Recently she recognized that her declining memory was becoming an issue. While enroute to her destination she would forget where she was headed and become lost. Other times she would forget there was food cooking on the stove, or fall asleep in the bath, causing safety concerns.

Joan made the difficult decision to move into an assisted living facility, where she found she was still experiencing challenges in dealing with her monthly finances. Forgetting to pay bills on time, sending the wrong cheque to the wrong company, or putting the wrong amounts on her cheques caused issues like her phone being disconnected and her rent to the care centre being bounced more than once.

With the help of a social worker at the care centre, Joan contacted EAC for support and ideas on how to cope with memory loss. EAC's Support Coordinator visited Joan and with Joan's social worker, came up with some strategies that made it easier for Joan to organize and track her financial records. This has alleviated some of Joan's worries and stress.

Joan now has time to connect with other residents and get back to things she enjoys doing, like puzzles and taking walks with others. Joan says, "I appreciate being able to speak to someone who understands epilepsy and the daily struggles that people with epilepsy live through each day."

# **2019 IN THE COMMUNITY**

#### **Engagement & Education**







808
Participants
attended

24
Community
education sessions

107
Requests for educational materials

### **Community Support**







160
Participants
Attended

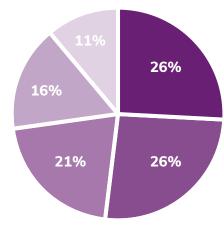
27
Peer support
groups & workshops

255
Total support
hours provided

#### **2019 FINANCIALS**

Epilepsy Association of Calgary is reliant upon the generosity of our donors, one-time funding from grant makers and Alberta Lottery and Gaming to fund our operations. Our goal is to engage with our membership base and establish new connections with those whose lives are affected by epilepsy to continue to strengthen our revenue streams into the future. We are grateful for the support of those taking part in the annual Scotiabank Charity Marathon and the Bare Naked Noggins head-shave. These events not only provided a source of revenue to our operations during 2019, but also offered our network ways to engage with the Association as fundraisers and participants. Thank you!

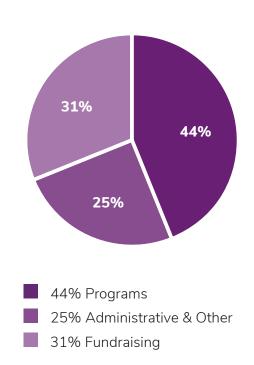
#### **2019 REVENUE**







#### **2019 EXPENSES**



During 2019, Epilepsy Association of Calgary embarked upon a thoughtful re-visioning of strategy, leadership and purpose. We know that investments in these efforts today will pay dividends in the long-run in terms of the quality of programs and services provided to those who rely upon us for information and support. In 2019 we completed our website and re-branding projects and employed a full-time fundraiser. A team of three full time staff was responsible for program delivery, communications, event planning and management, revenue generation and administrative duties while a search for a new Executive Director was completed.

#### **THANK YOU FOR CARING!**

We would like to extend our sincere thanks to everyone who has supported the Epilepsy Association with a donation. Every gift helps make a difference in the life of someone affected by epilepsy. We also acknowledge and thank those donors who prefer that we not share their names.

### **\$1000 AND ABOVE**

#### **INDIVIDUALS**

Cheryl Cohen Kathy & Derek Fyfe Kelly Hewson & Gordon Salahor Cheryl Menegoz Dale Poloway Jessica Weatherhead

#### **ORGANIZATIONS**

Estate of Eleanor Ganes







Harry & Martha Cohen Foundation







\$500 - \$999 LEVEL

#### **INDIVIDUALS**

Maria & Brad Cooper Kim & Stefan Hrach Terry Poole David Tod John Zutter

#### **ORGANIZATIONS**







#### \$100 - \$499 LEVEL

#### **INDIVIDUALS**

Thomas Albert William Appleby Sherry Austin Deborah Baigent Kevin Beavis Gary Ballantyne Kevin Bamford Gord Bandy Adela Bannerman Paula Bannerman

Jamie Bannerman-Flaterud

Janice Bateson Ross Beasley Garry B. Beres Gerry Bowland Scott Budau

Hanh Dung (Lee) Bui Wayne & Shannon Burwash

David Butler Brian Callow Rob Campbell Lorna Carpenter Ross Collett Ron Conley

Mansil & Marilyn Davis Brian Dowse Kevin Dradon Linda Eirikson Richard Fleurant Simon Forman Jodi Forzani Allan Fowler Adelein Frey Louise Gallagher Richard Galloway David Gammie Michael Gaschnitz Dale Getty Theo Goethals

Tom Guebert Don Hadley

Ken & Christine Hale

Carol Haley Patti Harrison Marlene Hein Blair Hicken **Bob Holmes** Greg Horton Donna Mae Howes Shawn Irwin Hasneen Jalil Margaret Johnson Cori Johnston **Edward Johnston** Paul Kelly

Craig Kemery

Amanda & David Kemery Dana Kemery Valguardson

Patty Kilgallon Zora Ljubojevic Ron Mackenzie Howard MacKichan Lana Macneil

Janelle Marietta-Vasquez

Allison Martin Linda McClure John McDonald Susan McLean Mitch McLeod Retta McLeod Cheryl McNab Francesco Mele Mary Merriman Rebecca Mesa Laurie Middlehurst Marc Murdoch Brendan Murray Elena Nechushkina

Gary Nissen Shelly Norris Karen O'Connor Mary & Ross Phillips

Art Piercy David Pyke Fred Pynn Lori Ramcharan Charles Ramcharan Rick Riffel

Linda & Rowland Rochow Stephen Rodrigues

Mary Rooney Hans & Mary Lou Saeby

Adam Sanvido Shelby Schultz Zach Schultz

Kristin Shandro Kerry Smith Roger Smith

John Spronk Christopher Stanley Paula Steele John Thomson Helen Tod William G. Turnbull Karla Turner Tim Tycholis Adam Walsh Frank Walsh

#### **ORGANIZATIONS**

Chris Wolfenberg

Deborah Warren

Helen Warren





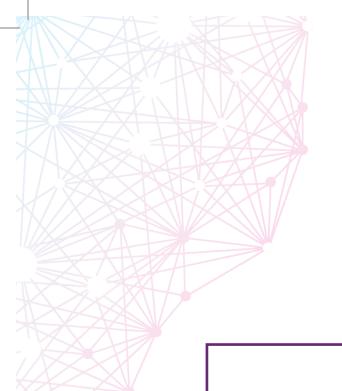






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# LIST OF MONTHLY SUPPORT GROUPS & WORKSHOPS

bit.ly/eac-support

# GET INVOLVED AS A VOLUNTEER WITH EAC

bit.ly/eac-volunteer

# **BOOK A COMMUNITY EDUCATION SESSION**

bit.ly/eac-education





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