ALS SLA ACTION CANADA

ALS SUPER FUND

Photo credit: Vancouver Canucks/Jeff Vinnick

2025-2026 IMPACT REPORT

COMMUNITY, COLLABORATION AND COURAGE continue to be the key drivers of ALS Action Canada and our cause fund, the ALS Super Fund.

From evolving the Courage to Fight ALS campaign to new heights and launching the Ales for ALS initiative with growing partnerships across North America, joining forces with the Canadian Collaboration to Cure ALS to advocate for increased research, to ensuring unique ALS voices are part of health policy considerations, your support has made this impact possible.

We are committed to the focused purpose established by our founders in 2020, the centering and amplification of ALS voices to accelerate treatments, increase advocacy, and mobilize investment in ALS research. It is only because of you we can build on the incredible legacy all those who have contributed to this work. Every ALS voice lost to this devastating disease is irreplaceable, and every ALS voice matters and is woven into the fabric of who we are.

While our valuable charitable status guides our operations, it doesn't wholly define us. We are a movement that is bigger than the sum of its parts.

Thank you to our board of directors, Committee members, Advisory Council, partners, volunteers, and incredible donors – you make our purpose possible.

Don Wright
Chair, Board of Directors,
ALS Action Canada

Leigh Naturkach
Executive Director,
ALS Action Canada

IMPACT HIGHLIGHTS

TOTAL RAISED
$2.2M+

1,650+
DONORS TO DATE

5
COURAGE TO FIGHT ALS NIGHTS WITH THE NHL

LAUNCH OF THE COURAGE TO FIGHT ALS CABINET

14%
INCREASE IN MEMBERSHIP

COLLABORATION WITH
13
ALS & ADVOCACY PARTNER ORGANIZATIONS

3
REPRESENTATIVES ON
2
NATIONAL ALS ADVISORY GROUPS

12+
COMMUNITY FUNDRAISING & AWARENESS INITIATIVES

21
PARTICIPATING BREWERIES IN ALES FOR ALS™

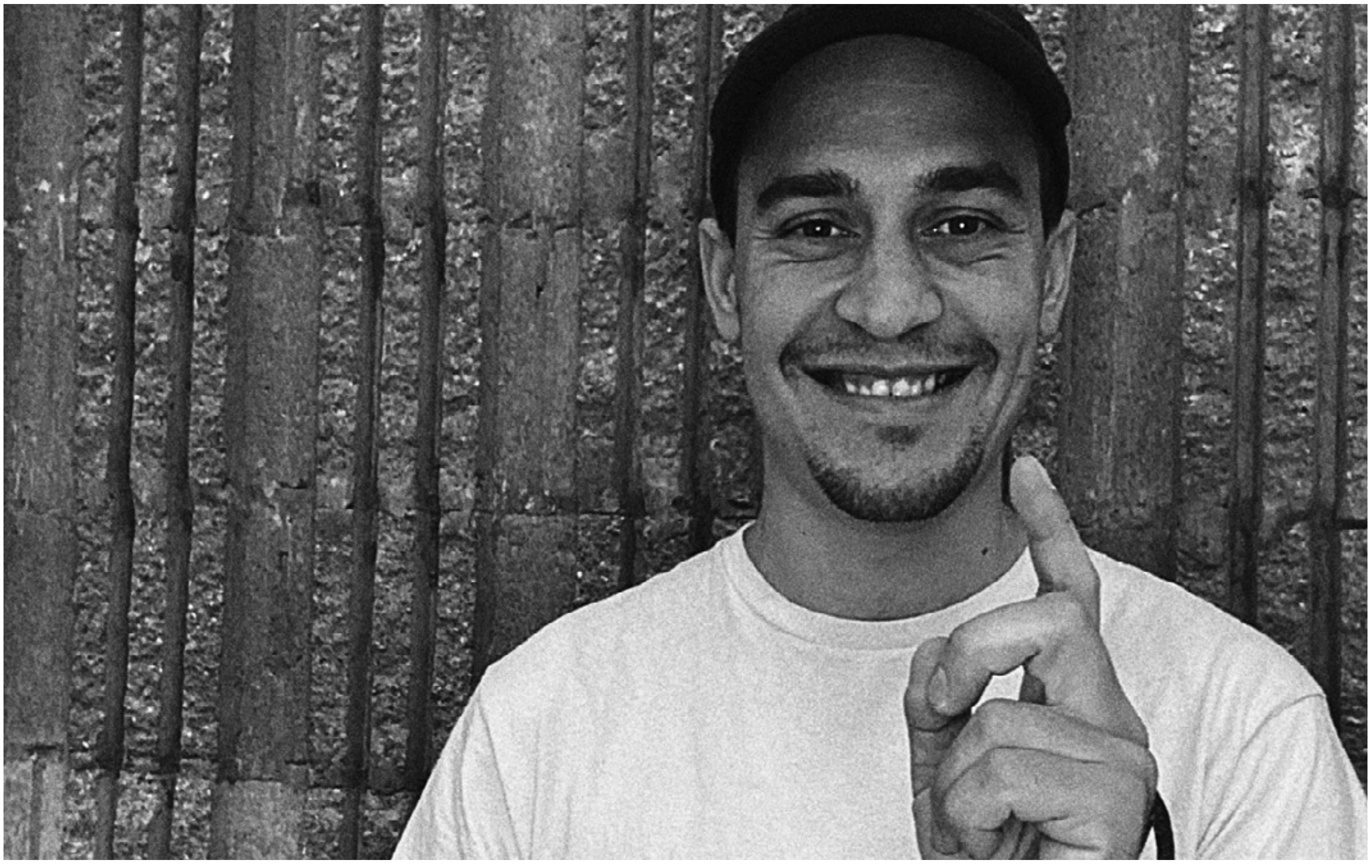
4
GRANTS TO ALS RESEARCH AND ADVOCACY IN NORTH AMERICA

4
FEATURES, WEBINARS & PODCASTS

1
JOURNAL PUBLICATION

3
HEALTH POLICY SUBMISSIONS

15
COMMUNITY STORIES & VOICES FEATURED



ABOUT ALS ACTION CANADA

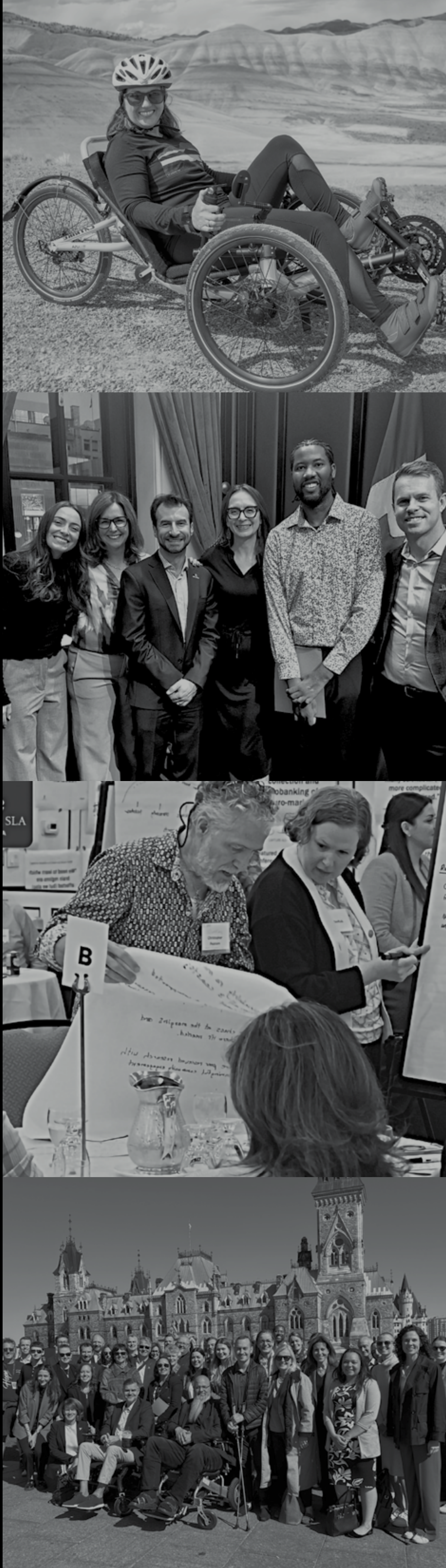
ALS Action Canada is Canada's patient-led ALS organization, centred on the voices and experiences of people living with ALS and caregivers.

Through advocacy, collaboration, storytelling, education, and community-building, ALS Action Canada works to accelerate treatments, mobilize investment in ALS research, and strengthen support for the ALS community across North America.

ALS Action Canada leads the ALS Super Fund, supporting high-impact ALS research, advocacy, and patient and caregiver priorities.

GRANTS

With your support, we fueled important research and patient voice initiatives across North America.



ALES FOR ALS RESEARCH AND PATIENT DRIVEN INITIATIVES

Through the Ales for ALS initiative in partnership with the ALS Therapy Development Institute (ALS TDI), craft brewers, beer drinkers and purchasers helped fuel patient voice work through ALS Action Canada and research through ALS TDI. Every ale sold helped projects such as the ALS Research Collaborative (ARC) Study – one of the largest and longest-running natural history studies in ALS. The study is designed to accelerate research, improve trial readiness, and deepen understanding of disease progression through patient-driven data and biological sample collection.

Through the Courage to Fight ALS campaign in partnership with the National Hockey League, proceeds from auctions, 50/50 draws, concourse activations and more drove support to research projects in BC and Alberta.

PROJECT HOPE: ADVANCING ALS RESEARCH AND CARE

In partnership with ALS BC, hockey fans helped fuel Project HOPE at UBC's Djavad Mowafaghian Centre for Brain Health, led by neurologist and ALS researcher Dr. Erik Pioro.

Project HOPE reflects a long-term collaborative initiative focused on strengthening ALS research, improving care, and supporting more connected opportunities for people and families impacted by ALS.

Support through the ALS Super Fund contributed to:

- Expanding ALS research initiatives
- Supporting post-doctoral fellows studying disease mechanisms
- Developing advanced imaging tools
- Strengthening clinical programs connecting patients with trials and innovative care
- Building multidisciplinary clinical care teams including occupational and physical therapists, speech therapists, nutritionists, social workers, and nurses

SUNNYBROOK HEALTH SCIENCES CENTRE - QUARTS-ALS TRIAL

The ALS Super Fund continued supporting the QuARTS research program based at Sunnybrook, the largest ALS Clinic in Canada. Led by Drs Zinman, Abrahao, and De Castro, this Trial is exploring whether non-invasive brain stimulation can safely target cortical hyperexcitability – a feature of ALS believed to contribute to disease progression.

During the past year, researchers completed the QuARTS-2 study cohort, evaluating the safety and feasibility of accelerated non-invasive brain stimulation in people living with ALS. The treatment was found to be safe and well tolerated, with most reported side effects mild and temporary.

Early findings suggest the therapy may temporarily strengthen the brain's natural "calming" mechanisms linked to motor neuron activity, supporting continued research into potential therapeutic approaches for ALS.

The findings from QuARTS are helping inform future phases of clinical research, including the planned next stage of the program, QuARTS-3, with the long-term goal of expanding future treatment possibilities for people living with ALS.

ALBERTA ALS RESEARCH NETWORK (AARN)

The hockey community in Alberta helped drive support for The Alberta ALS Research Network (AARN), an Alberta-based collaboration bringing together researchers and clinicians across the University of Calgary and University of Alberta to advance innovation in ALS care, strengthen research capacity, and improve outcomes for people living with ALS.

CANADIAN COLLABORATION TO CURE ALS

ALS Action Canada was proud to participate in the Canadian Collaboration to Cure ALS, a national initiative led by ALS Canada that brought together ALS organizations, researchers, clinicians, advocates, and community leaders from across the country in support of strengthening Canada's ALS research ecosystem.

A key focus of the year was joining forces for Hill Day activities in Ottawa, where dozens of members of the Collaboration engaged directly with Members of Parliament and federal stakeholders to advocate for increased investment and coordination in Canadian ALS research and innovation.

The initiative is calling for $50M in federal support for critical national ALS research and data infrastructure initiatives, including CAPTURE ALS, the Canadian Neuromuscular Disease Registry (CNDR), the Canadian ALS Research Network (CALS), and ACCESS ALS.

The late Matt Brown, former ALS Action Canada Advocacy Committee Member, played an important role in helping inspire this work through his engagement with federal decision-makers prior to his passing.

This work reflected ALS Action Canada's continued commitment to courageous collaboration and building united national movements that strengthen impact, advocacy, and research efforts in service of the ALS community.

NATIONAL ALS RESEARCH SUMMIT

ALS Action Canada participated in the National ALS Research Summit hosted by ALS Canada in June 2025.

Participation reinforced the importance of patient-centred collaboration and long-term coordination across the Canadian ALS research landscape.

Key discussions included:

- Accelerating ALS research collaboration
- Strengthening equity in access to care and clinical trials
- Long-term research priorities in Canada
- The importance of patient-informed approaches to research and policy

AMPLIFYING ALS VOICES

> When we have more and more people like my son wanting to get tested, wanting to find out and get pre-symptomatic care, which doesn't exist equitably across the country right now, I think that's a definite failure on the system. I would also like to see psychology services included in pre-symptomatic care. These are some of the reasons my son and I are Members of ALS Action Canada and something I try to advocate for.
>
> **– Paula Trefiak**

PUBLISHED PATIENT PERSPECTIVE ON ALS DRUG ACCESS

ALS Action Canada contributed an important patient and caregiver perspective to the international conversation surrounding ALS treatment access and regulatory decision-making through the publication of a Letter to the Editor in the peer-reviewed journal Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration.

Led by Advocacy Committee Member Dr. Andrew Darke, a person living with ALS, and Board Director Cali Orsulak, a former caregiver, the publication highlighted concerns surrounding how treatment benefit is assessed in ALS, emphasizing that traditional clinical and statistical measures do not always fully capture what meaningful change looks like for people living with the disease and their families.

Drawing from lived experience, the letter underscored the importance of incorporating patient and caregiver perspectives into regulatory, reimbursement, and health policy discussions – particularly in the context of a rapidly progressive disease where preserving function, independence, communication, mobility, or quality of life for even a short period of time may carry significant personal value.

The publication reflected ALS Action Canada's broader advocacy efforts to help ensure that urgency, lived experience, and patient-defined outcomes are considered alongside clinical and economic measures when evaluating emerging therapies and access pathways.

Throughout the year, this work continued to contribute to broader national and international discussions surrounding ALS drug access, patient-centred evidence, and how meaningful benefit is defined within rare and terminal diseases.

Subsequent international research and policy discussions further reinforced ALS Action Canada's position that even modest slowing of disease progression may be meaningful to many people living with ALS and their care partners.

CANADA'S DRUG AGENCY (CDA) SUBMISSIONS

ALS Action Canada provided recommendations and submissions directly to Canada's Drug Agency regarding:

- Treatment access
- Reimbursement review processes
- Patient and clinician engagement
- Access to Tofersen (QALSODY)
- Pre-symptomatic SOD1 treatment access

This work helped ensure patient and caregiver perspectives remained represented within important national conversations surrounding treatment access and reimbursement policy.

Canada's Drug Agency
L'Agence des médicaments du Canada
Drugs. Health Technologies and Systems. Médicaments, technologies de la santé et systèmes.

Submit a Request

ALS Action Canada Executive Director Leigh Naturkach (right) connecting with members of Canada's rare disease community at the CORD Conference. ALS Action Canada is a proud member of the Canadian Organization for Rare Disorders (CORD).

CAMPAIGNS & COMMUNITY ACTIVATION

ALS Action Canada continued building awareness, advocacy, fundraising, and community connection through collaborative campaigns, partnerships, and community-led initiatives across North America.

COURAGE TO FIGHT ALS™ CAMPAIGN EXPANDING AWARENESS THROUGH HOCKEY PARTNERSHIPS

Photo credit: Andy Devlin, Team Photographer Edmonton Oilers

Inspired by the late former NHL player, ALS advocate, ALS Action Canada Board Director, and ALS Super Fund Co-Founder Mark Kirton, the Courage to Fight ALS campaign continued growing into a broader North American movement uniting the hockey and ALS communities through awareness, storytelling, advocacy, and fundraising.

Throughout the 2025–2026 season, Courage to Fight ALS™ Nights helped:

- Raises awareness of ALS across 6 hockey communities
- Generate nearly $130,000 in funds for ALS in Canada. In addition, the Courage to Fight ALS Night in Nashville was acknowledged by Tennessee state representatives in their $10 million allocation in the budget to the Vanderbilt ALS Research Center.
- Elevate patient and caregiver stories
- Strengthen collaboration between ALS organizations, players, alumni, and supporters
- Expand cross-border engagement around ALS advocacy and research

Co-Leads:

Louis Del Re – *Board Director, ALS Action Canada, Augie's Quest and ALS TDI Award Winner, Person living with ALS*

Kevin Moorhead – *Vice-Chair, ALS Action Canada*

Secretariat:

Leigh Naturkach – *Executive Director, ALS Action Canada*

The Cabinet includes ALS advocates, NHL community leaders, philanthropic partners, and media professionals:

ALS Action Canada, pan Canadian

- Jeff Jackson – *Board Member, President & CEO, Edmonton Oilers*
- Jake Thompson – *Member, Person Living with ALS*

Brandon Montour, *NHL Defenceman*

Andrew Jackson, *President, Jackson Events Inc., Ontario*

Augies Quest to Cure ALS, Colorado/California /Washington/Ohio

- Shannon Shryne, *Co-Founder & President*
- Scott Cameron, *Board Member*
- Iain Duncan, *Volunteer Leader, NHL Alumnus*

Bourque Family Foundation, Massachusetts

- Ray Bourque, *NHL Alumnus, Hockey Hall of Fame*

Glenn Healy, *Ontario, President & Executive Director, NHL Alumni Association*

Kelsie Snow, *Former Caregiver, Inaugural Courage to Fight ALS award recipient, Project Manager, ALS Alberta Research Network, Alberta*

Peter Frates ALS Foundation, Massachusetts

- Mike Pitt, *Executive Director*

Tackle ALS, Massachusetts

- Troy Green, *Caregiver, Founder, Strong Companies*
- Tate Green Fox, *Caregiver*

Adam Fox, *NHL Player, New York Rangers*

Team Goose

- Gary Ragusa Jr., *Former Caregiver, Director, New Jersey*
- Luke Campomenosi, *Board Director, New Jersey*

Partners:

ALS Society of Alberta
ALS Society of BC
ALS Quebec

Participating teams included:

Courage to Fight ALS™ Nights included:

- Concourse activations
- Meet and Greets with NHL players
- Executive Suite and special experiences for ALS patients and families
- Auctions and raffles
- Ceremonial puck drops
- Awards for ALS advocates
- Storytelling, in-game interviews and tribute videos

Funds raised through Courage to Fight initiatives supported ALS organizations, advocacy efforts, patient support initiatives, and the ALS Super Fund.

BUILDING THE COURAGE TO FIGHT ALS™ MOVEMENT TOGETHER

The continued growth of Courage to Fight ALS™ is only possible through collaboration across the ALS, hockey, and broader community sectors to strengthen awareness, storytelling, advocacy, fundraising, and community connection.

Additional campaign moments throughout the year included a Toronto Maple Leafs tribute honouring the legacy of Mark Kirton, along with Vancouver Canucks Community Hero recognition for Adrienne Molinski and Greg Gowe.

> “The fight against ALS is the ultimate team sport, where patients, families, and like-minded foundations rely on one another with the same work ethic, fierce loyalty, and protection found in the locker room.
>
> Through our partnership with the NHL and the Courage to Fight ALS initiative, we expand this ‘Team’ into the hockey community — a culture built on the belief that when everyone on the ice and on the bench works as hard as they can for the person beside them, no goal is insurmountable.
>
> Together, we are an unbreakable team driven by a shared mission to provide a lifeline of support and a future free from this disease.”
>
> **– Mike Pitt, Peter Frates ALS Foundation**

> “Collaboration is what turns awareness into action. Through partnerships like Courage to Fight ALS and the incredible support of the NHL community, we’ve been able to give ALS patients and their families something they deserve most — moments to smile, make memories, and forget about the disease, even if just for a night. Team Goose is proud to work alongside so many amazing people who share the same mission of bringing hope, support, and meaningful experiences to the ALS community while continuing the fight for a cure.
>
> **– Gary Ragusa Jr., Director, Team Goose ALS Foundation**

COURAGE TO FIGHT ALS™ AWARD

ALS Action Canada presented the inaugural Courage to Fight ALS™ Award to the Snow family during the Edmonton Oilers' Courage to Fight ALS Night.

The award recognized Kelsie Snow and her children, Cohen and Willa, in memory of Chris Snow, for their courage, advocacy, and leadership within the ALS community. Chris Snow was a respected NHL executive with the Calgary Flames who publicly shared his experience living with ALS, becoming a powerful voice for ALS awareness, advocacy, and research while continuing to work throughout much of his diagnosis.

EDMONTON OILERS
COMMUNITY FOUNDATION

A $10,000 DONATION FROM THE EDMONTON OILERS COMMUNITY FOUNDATION

The donation was directed to the ALS Society of Alberta in honour of the award recipients. The recognition helped elevate awareness of the lived realities of ALS while celebrating leadership, resilience, and courage within the ALS community.

> "Chris' legacy is to always have that positive outlook and to always see the world as beautiful even when it's sad and hard.
>
> **– Kelsie Snow**

ALES FOR ALS™ & BREWERY PARTNERSHIPS GROWING THE ALES FOR ALS™ INITIATIVE IN CANADA

ALS Action Canada launched its collaboration with ALS Therapy Development Institute (ALS TDI) this year through joining ALS TDI's Ales for ALS™ initiative, bringing together breweries, communities, and advocates in support of ALS.

Participating Canadian breweries activated various special beer releases, community gatherings, fundraising efforts, and educational engagement throughout the year. The partnership strengthened connections between the Canadian ALS community and broader North American research and advocacy efforts focused on accelerating progress in ALS treatment development.

In the first year of the partnership, nearly $9,000 was raised through dedicated beer sales.

Participating breweries and partners throughout the year included:

AMSTERDAM Brewery TORONTO'S ORIGINAL
Aquanova BREWING CO.
PASSAGE
BAR VOLO
COLLECTIVE ARTS ART + BREWING
FIRST NATIONS Kahnawake BREWING CO. MICROBREWERY
TOOL SHED BREWING COMPANY
BICKFORD BREWING CO
The GRANITE BREWERY & TIED HOUSE
GREAT LAKES BREWING CO
BOSHKUNG BREWING CO.
5 PADDLES BREWING CO
CLIFFORD BREWING CO.
TRUE HISTORY BREWING
BELLWOODS BREWERY
BASE CAMP BREWING COMPANY
NONSUCH
LONG BAY BREWERY
Blood Brothers Brewing
TOOTH AND NAIL BREWING COMPANY

ALS Action Canada Board Director Louis Del Re was recognized with the inaugural Augie Nieto Legacy Award presented by ALS TDI for his leadership and advocacy related to Ales for ALS™ and broader ALS awareness efforts.

Granite Brewery Owner Ron Keefe – an ALS Action Canada member and the first brewery owner in Canada to participate in Ales for ALS™ – continued advocating for expanded brewery participation and community involvement throughout the initiative.

Granite Brewery also launched “Keep on Buzzin’,” a brew created in honour of Louis Del Re and the ALS community.

The initiative also reflected the strong family and community roots of Granite Brewery, including the involvement |of Ron Keefe’s daughter, Mary Keefe, Head Brewer at Granite Brewery, in supporting Ales for ALS™ initiatives.

ALS Action Canada also participated in the Ontario Craft Brewers Conference during the year, helping strengthen relationships with brewery leaders and expand awareness of the Ales for ALS™ initiative across Canada.

Together, these collaborations helped:

- Expand ALS awareness into new community spaces and audiences
- Strengthen community-led fundraising and engagement
- Build new partnership opportunities across Canada and the United States
- Create accessible opportunities for storytelling, advocacy, and education
- Reinforce the importance of collaboration in advancing ALS awareness and research support

> Being the first brewery in Canada to participate in the Ales for ALS program has been one of our greatest honours in our 35 years. It has brought our staff and customers together for such an important cause. It also reminds us, as we encourage others to join us, what a truly great industry and community we are part of.
>
> **– Ron Keefe, Granite Brewery**

STRATEGIC PARTNERSHIP: AMSTERDAM BREWING COMPANY

In 2026, ALS Action Canada launched a strategic partnership with Amsterdam Brewing Company focused on awareness, fundraising, storytelling, and community engagement in support of people and families impacted by ALS.

The collaboration helped expand public awareness of ALS while creating new opportunities to engage broader audiences through community-driven events and partnerships.

The Amsterdam partnership also reflected ALS Action Canada's growing focus on values-aligned collaborations that combine awareness, fundraising, advocacy, and storytelling in creative and accessible ways.

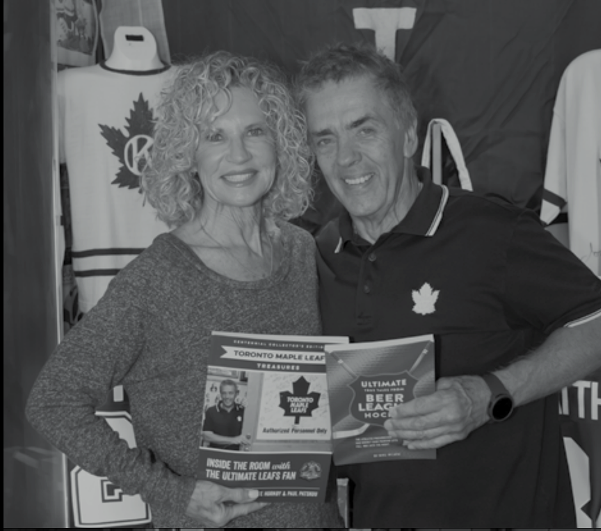
The partnership officially launched during Amsterdam Brewing's St. Patrick's Day Festival and included:

- **"A Legendary Stout" brewed for Ales for ALS™**
- **Festival activations and**
- **Featured menu items supporting ALS research**
- **Community storytelling opportunities**
- **On-site TipTap donation initiatives**
- **Cause-driven merchandise and engagement activities**

> Founded by uncompromising individuals and fearless innovation, we are proud to partner with ALS Action Canada. As the only Canadian ALS organization driven by ALS patients and caregivers, they share our approach and values in driving change....we stand alongside all those in the fight against ALS.
>
> **– Amsterdam Brewing Company**

COMMUNITY & FUNDRAISING HIGHLIGHTS

From galas and golf tournaments to brewery events and grassroots fundraisers, these initiatives reflected the growing momentum of a community united in support of people and families impacted by ALS.



KEEP ON BUZZIN' 2ND ANNUAL GALA

The Keep on Buzzin' 2nd Annual Gala brought together more than 650 supporters in support of Louis Del Re and Emily Robinson, while also raising incredible awareness for and $10,000 for the ALS Super Fund to help fuel advocacy, awareness, and research initiatives.

Led by the Del Re family and a dedicated volunteer committee, the event featured NHL alumni including Tom Fergus, silent auctions, Mike Wilson's OT Raffle in support of the ALS Super Fund, live music, dancing, and an evening focused on ALS education, storytelling, community care, and bringing people together in support of families affected by ALS. The gala continued building momentum for communty-led fundraising and awareness efforts connected to the ALS community, creating space for connection, advocacy, and meaningful conversations surrounding the lived realities of the disease.

The initiative's continued momentum was further reflected in the announcement of a third annual gala or May 2026.

FOREFEST CHARITY SCRAMBLE

Held in September 2025, the Forefest Charity Scramble raised more than $20,000 in support of the ALS Super Fund.

Led by Ryan Ferguson, the tournament selected the ALS Super Fund as its charitable beneficiary for the first time in recognition of Ryan's friendship with Louis Del Re and a shared commitment to supporting families affected by ALS.

The event brought together more than 100 golfers, volunteers, sponsors, and community supporters for a day focused on fundraising, awareness, and community connection in support of ALS Action Canada and the ALS Super Fund. ALS Action Canada extends its sincere thanks to Ryan Ferguson, the organizing committee, volunteers, golfers, and supporters whose efforts helped make the tournament a meaningful success.

The continued momentum of the event further reflected the growing strength of community-led fundraising initiatives connected to the ALS community, with plans already underway for the 2026 tournament.

KATHY CLASSIC GOLF TOURNAMENT

The Kathy Classic Golf Tournament, in memory of Kathy Boyle, directed $7,500 in proceeds toward ALS research initiatives supported through the ALS Super Fund.

Now entering its fifth year, the tournament continued building momentum as a growing community-led initiative supporting ALS awareness, research, and collaboration.

ALS Action Canada extends its sincere thanks to Amanda Diraddo The Diraddo real estate Team & BroLaws Team the tournament organizers, volunteers, golfers, sponsors, and supporters whose continued dedication has helped make the Kathy Classic an impactful community initiative in support of families affected by ALS.

ALS Action Canada is also honoured that the Kathy Classic Golf Tournament has selected the ALS Super Fund as its chosen charity once again for 2026.

PORTLAND CRAFT ALS FUNDRAISER

The Portland Craft ALS Fundraiser brought together supporters and community partners in support of ALS awareness and fundraising initiatives.

Together, these community-led initiatives reflected the strength, generosity, and commitment of the ALS community.

GREYSTONE CHARITY GOLF CLASSIC

The Greystone Charity Golf Classic raised more than $5,000 while honouring the legacy of Mark Kirton and supporting the ALS Super Fund.

Held shortly before Mark's passing, the event became a meaningful tribute to his leadership, courage, and lasting impact on the ALS community.

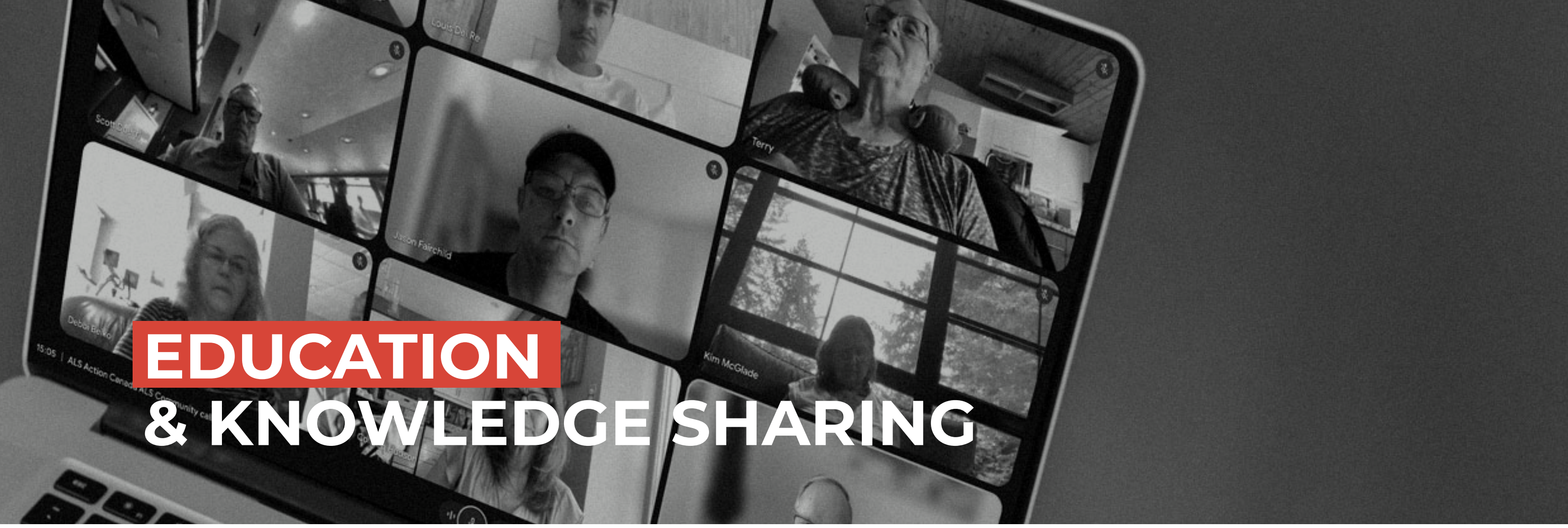
WILDEYE BREWING ALS PUB NIGHT

Wildeye Brewing's ALS Pub Night brought together the Vancouver community over two evenings in support of ALS advocacy, awareness, and research initiatives. Nearly $10,000 has been raised to date to the ALS Super Fund by Greg's Legacy Crew Giving Group. Greg Gowe was a co-founder of ALS Action Canada, advocate, entrepreneur, and a powerful voice within the ALS community.

STRENGTHENING CARE AND CONNECTION

The ALS Community Group is a patient-driven, biweekly virtual support and connection group that brings together people living with ALS, caregivers, family members, and advocates from across Canada to share experiences, discuss challenges, and support one another.

Facilitated by ALS Action Canada Board Director Rick Zwiep, this Group helped foster connection, reduce isolation, and create space for meaningful conversations surrounding the lived realities of ALS.

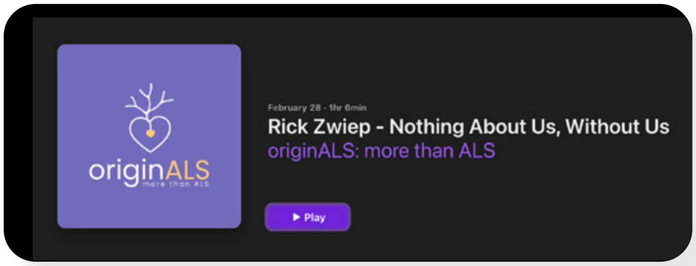


EDUCATION & KNOWLEDGE SHARING

Throughout the year, we connected the community to educational, research, and engagement opportunities across the ALS sector.

This included sharing and amplifying webinars, research updates, and educational initiatives related to:

- Approved ALS treatments
- Platform trials
- Clinical trial education
- Research developments
- Brain-computer interface studies
- International ALS research initiatives



February 28 · 1hr 6min

Rick Zwiep - Nothing About Us, Without Us

originALS: more than ALS

We were pleased to participate in the Canadian ALS Learning Institute (CALI) alongside ALS organizations and research leaders from across Canada, helping support greater awareness, education, and engagement within the ALS community.

For more about the CALI,
visit https://als.ca/advocacy/canadian-als-learning-institute/.

We've also aggregated some of the community-sourced resources and information on our new website, amplifying the efforts of many who lead this work in various ways.

Ideas? Recommendations? Share them with us at community@alsactioncanada.org.

POLLOCK FAMILY FOUNDATION

> We are proud to support ALS Action Canada because of the organization's uniquely ALS patient-centered approach. Supporting ALS Action Canada reflects our hope in a future where families facing ALS are met with compassion, dignity, and ultimately, ztreatment to end to this disease.
>
> **– Pollock Family Foundation**

THANK YOU

to Mike Wilson and Deb Thuet for their contribution through the OT Raffle fundraising initiative.

THANK YOU TO ALL OUR DONORS FOR YOUR SUPPORT

Akira Tamaki Charitable Account

Allen

Jennifer Amaral

Amsterdam Brewery Giving Group

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Braemar Parents Advisory

Beverley Brown

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Canucks for Kids Fund

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Dan

David

David Diamanti

Gino Del Re

Louis Del Re
and Emily Robinson

Dennis

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Fore Fest Golf Tournament

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Christine Grant

Great Lakes Brewing Company Inc.

Greg's Legacy Crew

Tim Greer

Greystone Golf Tournament

Ajeet Grover

Wade G Hall

David C Hann

Susan Audrey Hollinshead

Martin Horak

Colleen Hudson

Jacob Hudson

Martin Isaif

In Memory of Matt Brown
Giving Group

Katie Jamieson

Carolyn Jones

Kate

Kathy Classic Golf Tournament

Keep on Buzzin' Fundraiser

Keep on Buzzin' Giving Group

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Dale H Kendel
and Barbara H Kendel

Michael King

Jo-anne Kirton
and Kevin Marshman

Chris Krecisz

Yayu Li

Lisa Kirton Team

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Maple Leaf Sports and Entertainment

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Vanesse

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Michael A Wekerle

William

Michael Wilson and Debra Thuet (OT Raffle)

Don Wright

Victoria Young

Rick Zwiep

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Ken Karakashian

Veronique Saguin

Diane Shymkiw

Outram Tirbeni

If you see any errors or omissions, please contact: **info@alsactioncanada.org**

THANK YOU
TO OUR LEADERSHIP, VOLUNTEERS & CONTRIBUTORS

Our work is only made possible because of the caring, dedicated, and tenacious volunteers, advocates, contributors, and community leaders who continue to lead the way in support of people and families affected by ALS.

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Leigh Naturkach
Paul Thandi
Rick Zwiep

We were pleased to welcome Kevin Dawson and Lisa Kirton as new Board Directors in 2025.

We extend our sincere thanks to key organizational and operational partners whose support helped make our day-to-day work possible, including Power Play Production Group, Charitable Impact, Jordan Durham, Doane Grant Thornton, and Cadence Charity Services.

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