

Saturday, October 21 DAY 1 TRAINING DAYS PROGRAM (IN-PERSON ONLY)

KBHN Training Days are two pre-conference days (October 21, 22 2023) of workshops, informal discussion, networking, and Career Pathways.

Time (ET)	Programming / Activity
8:00 am	Registration opens / breakfast
	Location:
	Canadian Room
9:00 am	Opening Remarks
	Location:
	Canadian Room
	Speakers
	Jennifer Zwicker, Samantha Noyek, Samantha Micsinszki
9:30 am	KBHN National Training Program Evaluation: Where We Are and Where We Are Going Join us to explore the KBHN National Training Program and the recent findings from its strategic analysis. The presenters, including current KBHN trainees and persons with lived experience, will draw on the results of an environmental scan, survey, and semi- structured interviews to offer in-depth insight into the training program, including strengths and areas of need. As part of the presentation, attendees will have the opportunity to reflect on their own experiences with the KBHN training program and discuss how to ensure opportunities are aligned with trainees' diverse needs.
	Location:
	Canadian Room
	Speakers
	Alexandra Minuk, Carly Magnacca, Amanda Doherty-Kirby, Jeffrey McCrossin Moderator: Jennifer Zwicker



10:30 am	Break: Canadian Room
10:45 am	Advancing Equity, Diversity, Inclusion, and Justice via Relationships and Research
	Michelle Phoenix, Assistant Profession in the School of Rehabilitation Science (SRS) and CanChild Scientist at McMaster University is
	working to advance equity, diversity, inclusion and justice through her role as Co-Chair of the SRS Anti-Racism, Anti-Bias, Anti-
	Oppression Committee. Michelle will share research, reflection and resources about her experiences integrating EDIJ into education,
	research and service. She will provide opportunities for dialogue, reflection and planning to encourage everyone to advance EDIJ in the
	own work and life.
	Location:
	Canadian Room
	Speakers
	Michelle Phoenix, Moderator: Ayesha Siddiqua
11:45 am	Mitacs: Supporting Equity, Diversity, Inclusion and Decolonization
	Join David Plamondon, Senior Advisor of Indigenous Relations and Initiatives to learn about how Mitacs is actively working to create
	more equitable access to the Innovation and Research Ecosystem for Indigenous Peoples and other equity-deserving groups across
	Canada and abroad. This presentation will include a brief overview of the model of intern co-funding support that Mitacs provides,
	engaging Indigenous Peoples in research and innovation projects, and the newly launched Inclusive Innovation Action Plan.
	Location:
	Canadian Room
	Speakers
	David Plamondon, Moderator: Olivia Tsihlias & Melika Kangarani Farahani
12:45 pm	Lunch: Canadian Room



	This workshop aims to discuss approaches for bridging the gap between research and policy, focusing on improving outcomes for children with neurodevelopmental disabilities and their families. Recognizing the importance of evidence-based approaches, the workshop will discuss policy analysis approaches and strategies for developing inclusive and effective policy documents. Participants will gain insight into crafting policy documents to communicate key priorities and research to decision-makers. Through interactive discussions, the workshop will not only foster a deeper understanding of the complex challenges involved when translating research into policy, but also provide attendees the tools to develop their own policy communications.
	Participants will gain a deeper understanding of the intersection between research and policy impacting children and youth with neurodevelopmental disabilities. They will be equipped with actionable tools and strategies to influence policy decisions and foster collaboration with decision-makers. Ultimately, the workshop will encourage the exchange of ideas and contribute to the development of more informed, inclusive, and effective policy development.
	Location: Canadian Room
	Speakers Jen Zwicker, Sarah Raza; Moderator: Christiane Roth
2:30 pm	Networking BINGO break Take your conference networking to the next level with the exciting Networking Bingo Break! The bingo card contains fun challenges and conversation prompts that will break the ice and spark discussions. Get ready to dive into some fun, engaging conversations with other trainees. From industry insights to personal anecdotes, you'll be amazed at how quickly you connect with fellow attendees. Get ready to Bingo your way to meaningful connections and win some exciting prizes to take home!
	Location: Canadian Room
	Bingo planner and entertainer: Olivia Tsihlias and Ann Yang
3:00 pm	Implementation Science Skills
	Location: Canadian Room



	Speakers Julia Moore; Moderator: Vanessa Thomas
4:30 pm	Evening Out and About in Ottawa Join us for an exciting evening at the KBHN Conference Trainee Games Night on October 21st, from 16:30 to 18:30. The event will take place at the Level One Game Pub, located at 14 Waller Street, Ottawa, Ontario, K1N 9C4 (7-minute walk from conference venue). This private KBHN gathering offers the opportunity to play various games and connect with fellow trainees from across Canada! Dinner and a drink will be provided, including options for vegan and gluten-free diets. Check out https://levelonegamepub.com/ for details about the venue. Don't miss out on this fun event! Event planner and entertainer: Jessica Hanson

Sunday, October 22

DAY 2 TRAINING DAYS PROGRAM (IN-PERSON ONLY)

KBHN Training Days are two pre-conference days (October 21, 22 2023) of workshops, informal discussion, networking, and Career Pathways.

Time (ET)	Programming / Activity
8:00 am	Registration opens / breakfast
	Location: Canadian Room
9:00 am	Exploring Family Voices: Youth Engagement Panel This panel explores youth engagement from the different perspectives of family members, including siblings and parents. We highlight examples of youth engagement projects, including: i) the Youth Engagement in Research (@youth_in_research) Instagram account, ii) COVID time capsule, iii) Youth Engagement in Research (YER) project, iv) Sibling Youth Advisory Council (SibYAC) and sibling studies; and v) CHILD-BRIGHT's National Youth Advisory Panel (NYAP); vi) a study about the journeys of fathers of children with cerebral palsy. Youth with lived experience were engaged as collaborators, co-researchers and/or leaders in all projects.



	Location:
	Canadian Room
	Speakers
	Amanda Doherty-Kirby, Kelsey Seguin, Linda Nguyen, Samantha Bellefeuille- Moderator: Samantha Noyek
10:30 am	Break: Canadian Room
10:45 am	Family Engagement in Research (FER) Course in Practice - Innovation and Entrepreneurship in FER
	Location:
	Canadian Room
	Speakers
	Andrea Cross, Connie Putterman, Donna Thomson, Alicia Hilderley, Kristina McGuire-Moderator: Alice Soper
12:00 pm	Lunch: Canadian Room
1:00 pm	Entrepreneurship Skills
	Location:
	Canadian Room
	Speakers
	Speaker: TBD. Venture Labs at SFU? Moderator: Jessica Hanson
2:00 pm	Solutions Rooms: KBHN Training Program-Paving the Path Forward
	This session will follow up on the KBHN National Training Program Evaluation presentation, providing attendees with an opportunity f
	in-depth discussion about key findings. In small groups, attendees will receive a set of findings from the program evaluation to discuss
	with the goal of generating recommendations relative to the future of the training program. Attendees will be able to submit their recommendations along with any other insights they have about the training program for inclusion in the final program evaluation
	report.



	KIGS Brain Health Network
	Location:
	Canadian Room
	Speakers Alexandra Minuk and training program evaluation team- Moderator: Samantha Micsinszki
3:00 pm	Break
3:15 pm	Canadian Neurodevelopmental Research Training Platform (CanNRT) CanNRT is a collaborative training environment that supports sustainable and equitable pathways for Canadian early career researchers and trainees through cross-sector learning, mentorship and connectivity. Come learn about the Platform and interact with current Fellows, faculty and members of our leadership groups. Drinks and canapés will be served during this networking event. Pre-registration is required. Location: Canadian Room
5:00 pm	Speakers Closing Remarks & Prize Winners
	Speakers Jen Zwicker, James Reynolds
5:30 pm	Training Program Ends

Sunday, October 22 RECEPTION AND OUTSTANDING KBHN AWARDS

Time (ET) Programming / Activity



	Conference registration opens / networking
4:30 pm	Location: French Corridor
	Reception and Outstanding KBHN Awards - (Details coming soon)
6:00 pm	Location: Ballroom
8:00 pm	Reception Ends

Monday, October 23 DAY 1 CONFERENCE PROGRAM (HYBRID EXPERIENCE)

Time (ET)	Programming / Activity
7:30 am	Conference registration opens
	Location:
	French Corridor
8:00 am	Breakfast: French Corridor
8:30 am	Conference Opening and Welcome
	Location:
	Ballroom
	Speakers
	Nicky Lewis, James Reynolds



	Kids Brain Health Network
9:00 am	Fraser Mustard Lecture: Childhood Disability in the 21st Century: Can We Create a Canadian Model for the World? The Fraser Mustard Lecture has been a signature event at the Kids Brain Health Network's annual conference since 2012. This lecture is dedicated to the memory of Fraser Mustard, a world-renowned clinician and researcher who pioneered research on the importance of early child development. Each year, KBHN invites an individual who has made significant contributions to advancing the science of early brain development and/or the well-being of children from the perspective of research, clinical practice, or policy and advocacy, to deliver the Fraser Mustard Lecture.
	Location: Ballroom
	Speakers
	Dr. Peter Rosenbaum, M.D., FRCP(C), DSc (HC), FRCPI Hon (Paed) RCPI.
10:00 am	Break: French Corridor
10:30 am	Track A1:
Concurrent	Indigenous Communities Promoting and Supporting Infant and Early Mental Health through Community-Led Models
Sessions (Track A)	Through a collaboration between Indigenous communities and Infant and Early Mental Health Promotion (IEMHP), innovative models for strengthening knowledge and skill specific to infant and early mental health were created. These models of capacity-building engage both community members and professionals to collectively support young children. Unique to these models is the embedding of Indigenous culture which guided content, structure and delivery of all elements. The programs aim to build knowledge and understanding about infant and early mental health and draw attention to the important role both professionals and non-professionals play in supporting the wellbeing of young children. The initiatives focus on the strengths of service providers, families and communities while building an awareness that everyone has something valuable to contribute in supporting future generations. This session examines the collaborative process used to build relationships and support the integration of the science of infant and early mental health through an Indigenous lens. Participants will hear about the importance of engaging both professionals and community members to create a system of change. Two innovative programs will be explored:
	 Nurturing the Seed – A program providing professionals with training on infant and early mental health including the use of developmental support plans that are responsive to a child's needs in a family-friendly and culturally-informed way.



 Natural Helpers – An 8-week hybrid program designed for all community members to enhance knowledge about infant and early mental health and build community capacity through increased interest and commitment to better support very young children.

These initiatives demonstrate the benefits of sharing the science of infant and early mental health with ALL members of a community and doing so through an Indigenous lens. We will highlight preliminary evaluation results and plans for future evaluation strategies based on learnings from the projects to date.

Location:

Speakers

Chaya Kulkarni

Track A2:

Clinician-Caregiver Collaboration in Interventions for Caregivers

There is a dearth of interventions to support the wellbeing of family caregivers of individuals with neurodevelopmental disorders, who often have the responsibility of not only providing direct care for their family member, but also play various roles, including carrying out interventions at home or being an advocate outside the home. Our team has been studying the use of Acceptance and Commitment Training (ACT), a mindfulness and values-based intervention, to support caregiver well-being. One of the unique features of our program is the collaboration of clinicians and caregivers in the co-design and co-delivery of the intervention. We have also developed a comprehensive program to train and support clinician and caregiver facilitators.

In this panel, we will explore: 1) the benefits and challenges of clinician and caregiver collaboration in the design and delivery of the intervention; and 2) strategies, best practices and resources/policies to support, optimize and sustain clinician and caregiver partnerships. Our panelists will involve diverse perspectives, including organizational leadership, caregiver facilitators, clinician facilitators, ACT coaches and researchers, with varying degrees of experience and expertise with the intervention. As our model has been highly valued by clinicians, caregivers, and recipients of the intervention, we believe this will have relevance and implications to other caregiver interventions, training, and research initiatives.

Location:



Speakers

Kenneth Fung, Sacha Bailey, Cindy Hartman, Alyson Wilson

Track A3:

Navigating Childhood Disability Policy in Canada: Discussing perspectives from Families and Case Law

As a signatory to the United Nations Convention on the Rights of Persons with Disabilities (UN CRPD), Canada has committed to protecting the rights of all Canadians with disabilities. Despite this commitment, Canadian children and youth with disabilities often experience barriers to participation in society resulting in higher rates of poor physical and mental health outcomes, lower educational achievement and fewer economic opportunities than youth without disabilities. Parents and caregivers of youth with disabilities also face a high burden of care, resulting in financial and social challenges, and increased parental stress. The care needs of a disabled child often require that family members forgo paid employment outside the home.

This symposium will provide a short overview of disability policy in Canada including laws (such as the Accessible Canada Act, Bill C-22, and Bill C-35), plans of action, and provision of government-funded disability programs by health, education, social services, and economic ministries. The panel will then discuss ways in which disability policy falls short and the practical realities and challenges experienced by children and youth with disabilities, their families, and care teams. Our discussion will center on two main topics:

- Barriers experienced by parents/caregivers when applying for and accessing government-funded disability programs, sourced from interviews with families.
- Implications of current disability policy failing to adequately address the consequences of unpaid carework provided by parents/caregivers. The presumption in most provincial programs that family caregivers ought to be unpaid stands in contrast to personal injury damages awards and public insurance benefits, such as no-fault automobile insurance, which fully reimburse family carework. This means that children born able-bodied who become disabled by an injury are more likely to have access to funding for family caregivers than children born with their disabilities.

We will conclude with a discussion of our recommendations for solutions to these issues that can help to ensure children and youth with disabilities and their families are able to receive the support they require.

Location:

Speakers

Brittany Finlay, Samuel Ragot, Genevieve Currie, Stephanie Chipeur, Elizabeth Benediktsdottir, Brenda Lenahan; Chair: Lucy Lach



11:10 am	Track B1:
Concurrent Sessions (Track B)	Building Systems, Breaking Down Silos: Intersectoral collaboration to support access to care for infants, children, and their families Infancy to early childhood is critical for healthy brain and social-emotional development and an optimal time to identify and address early signs of mental health issues. However, access to care is often complicated and barriers can lead to inequitable care access, delivery and wellness outcomes for infants and young children. Care pathways help families and service providers by taking the guesswork out of what services are available and how to access them to ensure early intervention.
	The Knowledge Institute on Child and Youth Mental Health and Addictions and Infant and Early Mental Health Promotion are working with communities across Ontario to strengthen cross-sector partnerships that support streamlined pathways for infant and early menta health care. We will share outcomes from our pilot project where we partnered with cross-sector infant and early mental health service providers to create, implement and evaluate mental health care pathways for prenatal to age six.
	Through facilitated activities, we will showcase our new pathway resource guide: "From building blocks to care pathways: Working together to support access to infant and early mental health care." This guide leverages the latest evidence on how to plan, develop and implement care pathways with a focus on resources that build on community strengths and support families in accessing the right care in a timely manner. We will outline the processes and considerations that all communities should integrate into planning to sustain care pathways for effective and coordinated infant and child mental health services.
	Location:
	Speakers
	Amanda Davis, Chaya Kulkarni
	Track B2:
	Early Detection and Evidence-Based Early Interventions for Better Outcomes in Young Children with Cerebral Palsy
	International clinical practice guidelines exist for early identification and early evidence-based interventions for young children with cerebral palsy (CP). Adoption of these guidelines requires integrated knowledge mobilization and implementation science strategies to
	support a change in practice. For early detection, different approaches are required for neonatal follow-up medical specialists who see
	children who are at higher risk for developing CP (e.g., born pre-term) compared to community-based primary care physicians where
	children will present during their well-baby care visits with motor impairments. For early intervention, a knowledge gap exists for
	rehabilitation professionals, where there is a need for them to familiarize themselves with different therapy options and the evidence

Kids Brain Health Network #204 – Discovery 2, 8900 Nelson Way Burnaby, BC, V5A 4W9 Phone: 778-782-7283 | <u>www.kidsbrainhealth.ca</u>



	behind these approaches to support them as they move towards 'best' practice. Purpose: To present progress on knowledge mobilization and implementation science strategies to support early identification and intervention for young children with CP across Canada. For Early Detection, an overview of the EDIT-CP Toolkit (Early Detection section), including early attributes that can be detected by primary care providers and caregivers will be provided. Moreover, the progress on current implementation strategies will be summarized. For Early Intervention, an orientation to the Early Intervention section of the EDIT-CP Toolkit, which includes evidence-based information on more than 20 interventions, will be offered. The implementation strategies and the perspectives of a clinical partner will also be presented. These tools will assist physicians and rehabilitation specialists in expediting diagnosis and providing early interventions that are evidence-informed to ensure better outcomes.
	Location:
	Speakers Tatiana Ogourtsova, Annette Majnemer, Jessica Hanson, Filomena Pietrangelo
11:45 am	Lunch: French Corridor
12:30 pm	Poster Session 1 Join our poster event on 2 days to learn about the most recent research to support children with neurodevelopmental disabilities and their families.
	Location: Lauriel Room
2:00 pm	Track C1:
Concurrent Sessions (Track C)	 Individual Orals #1 Moving forward: Mobilizing knowledge to co-develop the Siblings Training, Empowerment, and Advocacy Kit (Siblings TEAKit) Caring for a Child with Prader-Willi Syndrome: A Narrative Inquiry Caring for the Caregiver (C4C): An Integrated Stepped Care Model for Caregivers of Children with Medical Complexity Co-designing solutions to enhance access and engagement in pediatric telerehabilitation KidsAction Coaching in Action with Challenger Baseball-Jays Care Foundation
	Location:



Speakers:

Linda Nguyen & Samantha Bellefeuille, Genevieve Currie, Natasha Bruno, Meaghan Reitzel & Cynthia Lennon, Amanda Field, Moderator: Sam Micsinszki

Track C2:

Supporting diverse families of children with neurodevelopmental disabilities- insights from community-based research

Navigating behavioural and mental health services for children and youth with neurodevelopmental disabilities can be incredibly disorienting, and often lead to high frustrations and stress for parents and families. Importantly, there is a growing awareness of the inequities experienced by families from diverse communities who face more barriers to accessing adequate care. There is an urgent need to act and co-create collaborations with community partners that can utilize research expertise to find solutions to real-world problems faced by diverse families across Canada. This symposium will provide insights from 4 community-based research projects and bring together the perspectives of researchers and community partners who have an interest in supporting families.

Location:

Speakers:

Vivian Lee, Jessica Remedios, Maria Rogers, Scott Robson

Track C3:

FASD & Me - Strengthening My Community: Leveraging the value of peer support and community advocacy to improve outcomes for youth with FASD

The FASD & amp; Me program aims to decrease stigma, build understanding, and develop actionable resources that are accessible to parents, caregivers, and youth with FASD themselves. FASD & amp; Me resources celebrate children and youth with FASD and their families; equip members of their communities to better support them by strengthening understanding of the unique strengths and challenges of individuals with FASD; and dispel myths and stereotypes.

The FASD & amp; Me: For Teens pilot was added in 2023, providing peer mentoring opportunities to teens with FASD and enabling these youth to gain a better understanding of FASD while developing self-advocacy skills to share with extended family, community members, and potential employers. Peer support, mentoring, and community connections are enhancing lives at all stages through innovative community-inspired approaches.



	Location:
	Speakers: Tracy Moisan, Shelby De Rose, Nicole Williams, Nancy Lockwood
2:40 pm Concurrent Sessions (Track D)	Track D2: From basic science to community, new perspectives from developmental neuroscience There is a movement towards fostering greater researcher-family partnerships within the realm of neurodevelopmental research. These partnerships are important because they have the potential to create better research questions that have stronger and more applicable outcomes for the community. These kinds of partnerships have formed more readily between researchers in applied, health, and clinical settings, yet researchers from the basic sciences (e.g., cognitive and neuroscience) have yet to have as many opportunities to make direct connections with families. In this panel discussion, neuroscientists from Carleton University will describe their work and its applicability to neurodiverse children and youth, and their families. Location:
	Speakers: Amedeo D'Angiulli, Argel Aguilar Valles, Mason Irvine, Vivian Lee
	Track D3: Weathering the Storm: Strengthening outcomes for families experiencing AFCCA (Aggression toward Family/Caregivers in Childhood/Adolescence) Aggression towards Families/Caregivers in Childhood and Adolescence (AFCCA) can be experienced by families impacted by neurodevelopmental disabilities and complex trauma; and this issue has been aggravated by the COVID pandemic. Early indicators support the hypothesis that combining peer supports, supportive communities, and effective evidence-based interventions can significantly change outcomes for both youth and their parents/caregivers, and potentially mitigate the need for more costly, intensive interventions in later developmental stages.
	Research often takes 10+ years to be implemented into clinical practice; these projects show how implementation can be accelerated through effective, meaningful family and community engagement in all stages of research and application.



	Location:
	Speakers: Tracy Moisan, Maude Champagne, Nancy Lockwood, Jennifer Pelissero
3:15 pm	Break: French Corridor
3:45 pm Concurrent Sessions (Track E)	 Track E1: Individual Orals #2 (1) Community Training and Implementation of the Social ABCs, an Intervention for Toddlers with Autism: Spread, Scale, Sustainment (2) Implementation of a Novel Family-Centered Program within an Ontario Autism Service Setting: Baseline Findings from a Mixed Methods RCT (3) The Ultimate Research Partner Guide to the Applicant Profile CV (4) Genetically programmed retinoic acid deficiency during gastrulation phenocopies developmental defects due to acute PAE in FASD (5) The Journey of Becoming and Being a Father to a Child With Cerebral Palsy
	Speakers: Jessica Brian, Terry Bennett, Greta Heathcote, Berardino Petrelli, Kelsey Seguin, Moderator: Jennifer Zwicker Track E2: Innovative programs meeting the needs of students with NDDs in the classroom There are many students, including those neurodevelopmental disorders (NDDs), who require support to be successful in today's classrooms. To be able to provide this support, educators need ongoing access to evidence-based practices. Dr.McGonnell, teacher and psychologist, will open the symposium with an introduction to the current context of these needs and to the importance of evidence-based interventions being accessible to teachers so they can better meet the needs of students with NDDs. Dr.Corkum, child psychologist and researcher, will discuss the development, evaluation, and implementation of the ASSIST (Accessible Strategies



Supporting Inclusion for Students by Teachers) program, an online program to provide classroom teachers with professional development on how to work effectively with children with NDDs, including ADHD, ASD, and LD. This presentation will highlight lessons learned from a national implementation study.

Drs.Rasmussen and Pei will discuss the implementation and evaluation of the Math Interactive Learning Experience (MILE), which was originally developed for children with FASD. MILE supports math learning by building underlying cognitive skills critical to math while as well as supporting the development of problem-solving strategies that facilitate emotional regulation.

This presentation will 1) highlight adaptations made to the MILE program to allow for classroom-wide implementation and 2) introduce the website-hosted training that supports asynchronous MILE training. Dr.Timmons will discuss the findings a four year-longitudinal mixed-methods study evaluating the effectiveness of an outreach program designed to build capacity of educators to support students with FASD. This presentation will highlight the importance of a collaborative approach to program evaluation while highlighting how practitioners can leverage the results to increase access to equitable learning opportunities for students with FASD globally. The symposium will conclude with discussant Adelee Penner, community partner and educator, reflecting on key points, including EDI considerations, from an educational perspective.

Location:

Speakers:

Kristy Timmons, Melissa McGonnell, Penny Corkum, Carmen Rasmussen, Jacqueline Pei, Adelee Penner

Track E3:

The F-words Implementation Journey: From Publication to Global Impact

Implementation is a central component of CanChild's 'F-words for Child Development' Research and Knowledge Translation Program. Our collaborative implementation work to date has catalyzed our understanding and the momentum of F-words implementation globally and resulted in numerous learnings and subsequent projects/initiatives. This presentation provides an overview of our implementation efforts and highlights the essentiality of Implementation Science to move evidence-based innovations into practice.

Location:

Speakers:



	Tracy Moisan, Maude Champagne, Nancy Lockwood, Jennifer Pelissero
4:25 pm	Track F3:
Concurrent	The Importance of Telling and Sharing Your Story - Using Digital Storytelling to Foster Healing and Resilience
Session	Stories connect us to each other. Stories are a compelling and meaningful way of sharing wisdom, not just data. Join Kristy Wolfe as she
(Track F)	shares what digital storytelling is, what is involved in the process and how it can be used within the context of knowledge translation. This interactive session includes digital story examples, space for conversation and opportunities to reflect on how your research project could benefit from digital storytelling.
	Guests will learn:
	 what a digital story is and why it matters
	 the process of creating a digital story
	 identifying key moments and uncovering lessons
	the impact of sharing digital stories
	Location:
	Speakers:
	Kristy Wolfe

Tuesday, October 24 DAY 2 CONFERENCE PROGRAM (HYBRID EXPERIENCE)

Time (ET)	Programming / Activity
7:00 am	Conference registration opens
	Location: French Corridor



7:30 am	Breakfast: French Corridor
8:30 am	Opening Remarks
	Location:
	French Corridor
	Plenary Session: Mobilizing Knowledge and Changing Practice
	The gap between knowledge creation and knowledge uptake means that research innovations frequently do not reach their intended audience. The Kids Brain Health Network has positioned itself to be an instrument of change in this equation, with a focus on the skills and pathways necessary for implementation of research innovations into practice. In this exciting plenary session, KBHN has invited three change leaders from the areas of research, advocacy and policy who will bring a multisectoral lens to the discussion of what it takes bridge the gap between knowledge creation and uptake to improve outcomes for children with neurodevelopmental disabilities.
8:45 am	Location: French Corridor
	Speakers
	Dr. Susanne Benseler, Director, Alberta Childrens Hospital Research Institute, University of Calgary Sue Robins, Healthcare Activist, Speaker, and Author
	Moderator: James Reynolds
10:00 am	Break: French Corridor
10:30 am	Track G1:
Concurrent	"Nothing About Us Without Us": Equitable and Accessible Youth Engagement in Research
Sessions (Track G)	This session will briefly explain the Youth Engagement in Research Project results. However, more importantly, by utilizing our experience as a research team, we will demonstrate an accessible patient partnership. We will cover our team's journey, from creating
	the research idea to disseminating knowledge. We hope those who attend will leave with a more robust understanding and
	appreciation of how to develop research projects that, from the start, focus on patient engagement in a way that helps not only further research but also continues to push back against the ableist foundations of research.
	Location:



Speakers Amanda St. Dennis, Jessica Geboers

Track G2:

Re-balancing the Wheel in the diagnosis, intervention and prevention of FASD and related NDD

This presentation looks at the accomplishments and the challenges of two community based clinics delivering services for FASD and other NDD's related to adversity. Both clinics opened more than 15 years ago serving youth and families in small but diverse communities--one in the inner city and one in a rural indigenous community. Both view health and healing in a larger social context in terms of strengthening relationships and restoring balanced environments to promote the development of youth gifts. In both communities there were social and economic barriers to accessing diagnostic and intervention services leading to youth health inequities. This in turn perpetuated a cycle of social and economic disadvantage among youth and families living with Neurodevelopmental disabilities that needed accommodations. The Eastern Door opened in 2006 and was the first multi-disciplinary team offering assessment for FASD and related ND conditions in Atlantic Canada. With ED practice there has been an increase in protective factors: earlier diagnosis and implementation of accommodations in school and family supports at home. This has in turn has led to decrease in prevalence and success of individual youth and young adults living with FASD and part of the ED. The ED just won the Korczac prize awarded by social pediatricians at the Pediatric Conference in Halifax.

The Richer initiative (Responsive, Interdisciplinary, Intersectoral Child and Community Health Education and Research) developed as collaborative interdisciplinary partnerships of primary health providers to build evidence based services for socially isolated and materially disenfranchised families in Vancouver's DTES. It has led to a critical difference in the vulnerability of the children in the area. RICHERS interdisciplinary research to practice model has been recognized internationally for its development of an effective health service delivery model linking services across sectors.

Location:

Speakers

Lori Cox

Track G3:

21st Century Philosophies for Developmental Disability: Exploring Approaches and Applications in Practice and Policy



	'Developmental Disability' is undergoing a profound sea change. In this symposium, we will propose a uniquely Canadian approach to 'developmental disability' that is having international impact on the field of child health and development. We will share updated information on CanChild's F-words for Child Development; Family-Centred Services; family engagement in research; and non-categorica approaches to functioning. We will then discuss next steps for working with policy makers and organizations to update and, in some cases, redesign systems of care need to meet these contemporary views. Importantly, we will challenge the traditional approach to services based on diagnosis, and propose and support alternate ways to think, plan and act.
	Location:
	Speakers Peter Rosenbaum, Jennifer Zwicker, Sara Pot, Rachel Teplicky, Patricia Basualto, Angela Senevirathna
11:10 am	Track H1:
Concurrent	Prioritizing children in policy: Lessons learned from a scoping review of national autism strategies around the world
Sessions (Track H)	This session presents select findings from an in-depth international scoping review of national autism strategies with the overarching aim of informing the development of Canada's own federal autism framework. Using the United Nations Convention on the Rights of Persons with Disabilities (UNCRPD) as an organizational framework, panelists will talk about why a national autism strategy is essential and discuss both quantitative findings and qualitative findings of the review. Panelists with lived experience will provide critical context for the panel discussion, offering their perspectives and priorities. Drawing on diverse areas of expertise, panelists will speak to the topics of awareness-raising; health, habilitation, and rehabilitation; political participation; education; employment; intersectionality; an research, statistics, and data collection, which is followed by a question and answer session.
	Location:
	Speakers
	Christiane Roth, Jennifer Zwicker, Yvonne Spicer, Sarah Ewart, Alexandra Minuk, Preeti Kar
-	Track H2:
	Building an Evidence-Informed Knowledge Translation Resource to Promote Healthy Outcomes in Individuals with FASD: Development and Process



	This presentation will characterize the three developmental phases we have undertaken to build COMPASS. In the first section, we will summarize the results of a secondary data analysis of a large national database of individuals with FASD across the lifespan, through which neurodevelopmental periods and their relationship to interventions and recommendations were identified. The second section will describe the developmental process of building a pilot version of COMPASS using the Towards Healthy Outcomes framework, an evidence-based framework developed to help support consistent and strength-based intervention planning across the lifespan. In the third section of the presentation, we will summarize evidence collected through the Tool pilot. We will highlight community-based collaborative processes woven through all COMPASS development and implementation aspects. One of our clinic partners, Dr. Valerie Temple, will provide information on her role in supporting our research team, the challenges and the importance of developing an
	evidence-based resource that effectively communicates knowledge and recommendations for best practice.
	Location:
	Speakers
	Kathleen Kennedy, Kaitlyn McLachlan, Jacqueline Pei, Valerie Temple
11:45 am	Lunch: French Corridor
12:30 pm	Poster Session 2 Join our poster event on 2 days to learn about the most recent research to support children with neurodevelopmental disabilities and their families.
	Location: Lauriel Room
2:00 pm Concurrent	Track I1: Individual Orals #3
Sessions	(1) The forgotten caregivers: A qualitative study exploring the experience of fathers of children with medical complexity
(Track I)	 (1) The folgotter caregivers: A qualitative study exploring the expendice of fathers of emidden with medical complexity (2) The Role of Emotion Regulation and Joint Attention in the Emergence of Autism Spectrum Disorder: An Infant Siblings Study (3) Using Machine learning to identify clinical and socioeconomic predictors of acute care use among medically complex children in Alberta
	 (4) Exploring the relationship between emotion and features of eye-gaze in youth with severe motor and communication impairment (SMCI) and typic



(5) Comparative Strengths and Challenges on Face-to-Face and Computer-Based Attention Tasks in Autistic and Neurotypical Toddlers

Location:

Speakers:

Vanessa Fong, Sarah Raza, Michael Sidra, Mackenzie Collins, Lori-Ann Sacrey, Moderator: James Reynolds

Track I2:

Better Nights, Better Days for Children with Neurodevelopmental Disorders Program: So What Happens After the RCT? Sleep problems are highly prevalent and impactful on children with NDDs and their families. Dr. Corkum, child psychologist and researcher, will introduce the Better Nights, Better Days for Children with NDDs (BNBD-NDD) program, an online, transdiagnostic, selfguided behavioural sleep intervention. She will review the steps taken to develop this program and the preliminary results of a Canadawide randomized controlled trial (RCT). Alzena Ilie, Clinical Psychology PhD student, will discuss the results of exit interviews with 20 parents who completed the RCT of BNBD-NDD. This presentation will highlight the barriers and facilitators that affect the Reach, Effectiveness, Adoption, Implementation, and Maintenance of the program. Katie Vaughan, BNBD-NDD Project Manager, will discuss how feedback from parents who participated in the RCT and exit interviews was used to modify the BNBD-NDD program. This presentation will highlight the modification process, as well as preliminary findings of the ongoing implementation study. Mya Dockrill, incoming Clinical Psychology PhD student, will discuss how an EDI lens can be applied to the BNBD suite of programs. This is particularly relevant given that there is a higher prevalence of sleep problems among children from racial and ethnic minorities, resulting in the need for equitable and inclusive access to sleep health care. Jocelyn Paul, a Mi'kmaw Clinical Psychology PhD student, will share the results of a scoping review examining the current literature about sleep and Indigenous peoples and discuss the next steps including learning circles and usability studies. The results of these studies will be used to determine if BNBD is able to be modified to be culturally appropriate and usable in this population. The symposium will conclude with lived experience discussant, Connie Putterman, reflecting

Location:

Speakers:

Katie Vaughan, Penny Corkum, Alzena Ilie, Mya Dockrill, Jocelyn Paul, Connie Putterman

on key points from the perspective of a parent of a child with NDD and insomnia.



	Rius Brain Health Network
	Track 13: The Family Engagement in Research Training Program: An Opportunity to Connect, Collaborate, and Co-Create Solutions to Common Challenges in Patient-Oriented Research In 2018, family partners and health service researchers at CanChild Centre for Childhood Disability Research at McMaster University partnered with Kids Brain Health Network to launch the Family Engagement in Research (FER) Course. Over the past five years, this 10- week online course for researchers and family members/caregivers with lived/living experience has evolved into an internationally recognized multi-faceted training program. Our program is unique in that it is designed to nurture connections, collaboration, and co- creation, and to build trusting relationships between people with lived/living experience and researchers. We achieve this by enabling co-learning in a safe and welcoming environment. In this workshop, we will facilitate roundtable interactive solutions-focused discussions about strategies to address two common barriers to engagement identified from graduates and instructors of the FER Course: i) readiness to partner in research for family partners and researchers; and ii) creating psychologically safe spaces and welcoming environments for partnership. Workshop presenters and small group facilitators include FER Course developers, instructors, coordinators, knowledge brokers, and graduates who are both researchers and family partners. Workshop participants will be invited to join an ongoing FER Community of Practice to support continued connection, collaboration, and co- creation the identification of strategies and development of collaborative working groups that will support the advancement of best practices in family engagement in research across Canada. Location:
	Speakers: Andrea Cross, Genevieve Currie, Krista McGuire, Linda Nguyen, Connie Putterman, Donna Thomson
2:40 pm Concurrent Session (Track J)	Track J1: Pediatric Cognitive Rehabilitation: Implementation lessons learned across diverse contexts. Attention and executive functions (A/EF), critical for academic success and quality of life, are almost universally disrupted in children with neurodevelopmental disorders (NDDs). Untreated A/EF problems lead to long-term mental health/behavioural, academic, and occupational/social difficulties, in addition to lower quality of life. While a number of cognitive rehabilitation programs have sought to address the needs of children with NDDs, those currently available are typically ineffective, expensive, or inaccessible to those in remote communities or without access to specialized professionals. Dino Island is an innovative cognitive intervention for children with NDDs that is designed to bridge this significant treatment 'gap'. It combines the appeal of tablet-based gaming with the effectiveness of an in- person, interventionist-child structure. We will discuss the unique implementation considerations of Dino Island in three distinct contexts: parent/home delivery, therapist/hospital-based delivery, and international remote delivery (Malawi, Africa). Our symposium



will focus on unique considerations for delivering cognitive rehabilitation across these distinct contexts in addition to community impacts, successes, and lessons learned.
Location:
Speakers:
Sarah Macoun, John Sheehan, Yu Shi, Ryan D'Arcy, Dacey Doyle, Martin Doyle Award Presentation & Closing Remarks
Location:
Speakers:
End of 2023 KBHN Conference