

2021 Accelerator Grant Awardees

	Project Title	Investigator(s)¹	Institution(s)²
01	Surgical genomics: how genetics impact surgical outcomes in Epilepsy	Danielle Andrade (PI) Paula Marques Quratulain Zulfiqar Ali	University Health Network
02	A splicing code resource for the discovery and characterization of neural microexons and associated genetic-based mechanisms underlying neurological disorders	Benjamin Blencowe (PI) Quaid Morris Sabine Cordes	University of Toronto Mount Sinai Hospital Memorial Sloan Kettering Cancer Center
03	Linking genomic data and administrative data to measure health service utilization and costs of genomic sequencing	Yvonne Bombard (PI) Murray Krahn Kevin Thorpe Jordan Lerner-Ellis Raymond Kim	St. Michael's Hospital Mount Sinai Hospital University Health Network
04	Single-nucleus transcriptomics and chromatin accessibility in a mouse model of hypoplastic left heart syndrome	Rajiv Chaturvedi (PI) John Sled Michael Wilson Miriam Reuter	The Hospital for Sick Children
05	Whole Genome Sequencing to understand genetic predisposition for Atypical Femur Fractures (AFFs)	Angela Cheung (PI) Lisa Strug Shinya Ito	University Health Network University of Toronto The Hospital for Sick Children
06	Rapid Genome Sequencing For Diagnosis And Discovery In Severe Infantile Epilepsy	Gregory Costain (PI) Danielle Andrade Vann Chau Robin Hayeems Puneet Jain Christian Marshall Roberto Mendoza-Londono Anna Szuto Ryan Yuen	The Hospital for Sick Children University Health Network
07	Accelerating PRrecision mEDicine for CriTical Care Study (PREDICT)	Claudia Dos Santos (PI) Alexandra Binnie Pedro Castelo Branco Pingzhao Hu Robert Hancock Uriel Trahtemberg Andrew Baker Margaret Herridge Angela Cheung Lisa Strug	St. Michael's Hospital William Osler Health System University of Algarve University of Toronto University of Manitoba University of British Columbia University Health Network
08	Combining bait-capture with whole genome sequencing to improving diagnosis of encephalitis of unknown etiology	Robert Kozak (PI) Amadou Sall Gary Kobinger Samira Mubareka Natalie Knox Robyn Lee Andrew McArthur Jessica Forbes Jalees Nasir	Sunnybrook Research Institute Pasteur Senegal Université Laval McMaster University University of Toronto National Microbiology Laboratory

09	Harnessing multi-omics to deliver innovative diagnostic care for rare genetic diseases in Canada (C4R-SOLVE)	Christian Marshall (PI) Michael Brudno James Dowling Robin Hayeems Roberto Mendoza Kym Boycott	The Hospital for Sick Children University Health Network Children's Hospital of Eastern Ontario
10	Structural variants in childhood heart disease	Seema Mital (PI) Ryan Yuen Rebekah Jobling Raymond Kim Erwin Oechslein	The Hospital for Sick Children University Health Network
11	Defining the chromatin-bound epitranscriptome of human pluripotent stem cells using long-read Nanopore sequencing	Miguel Ramalho-Santos (PI) Jared Simpson	Mount Sinai Hospital Ontario Institute for Cancer Research
12	Identifying candidate molecular markers in epilepsy by integrating DNA methylation and whole genome sequencing data	Rosanna Weksberg (PI) Elizabeth Donner Vann Chau Gregory Costain Evdokia Anagnostou Miriam Reuter Arun Ramani Andrei Turinsky	The Hospital for Sick Children Bloorview Research Institute
13	Profiling genotype and epigenetic status of tandem repeats from nanopore sequencing	Ryan Yuen (PI) Jared Simpson	The Hospital for Sick Children Ontario Institute for Cancer Research

¹ Bold indicates Lead or Co-Lead Principal Investigator (PI)

² Bold indicates Lead Institution