# **Cristian Groza**

## **DETAILS**

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*Phone* 438-921-1029

Github https://github.com/cgroza (software projects)

### **EDUCATION**

#### PhD. Quantitative Life Sciences

McGill University

BSc. Joint Computer Science and Biology

McGill University

DEC. Health Science

Dawson College

2019-

2016-2019

2014-2016

# RESEARCH EXPERIENCE

### Japan Society for the Promotion of Science fellowship

Institute for the Advanced Study of Human Biology, Kyoto, Japan

Spring 2023

• Developed novel genome graph methods for comparative epigenomics in primate genomes with Drs. Fumitaka Inoue and Guillaume Bourque at Kyoto University from 3 April to 3 June 2023.

#### Research Assistant

2017 - 2019

Contract at MUGQIC, Dr. Guillaume Bourque's Lab

• Developed pipeline applying personalized genomics to hundreds of epigenomic datasets within the McGill Epigenomics Mapping Centre.

#### Rotating Student

Fall 2019

PhD rotation in Dr Yue Li's lab

Department of Computer Science, McGill University

• Applied topic learning models to single cell transcriptomic data.

#### Rotating Student

Summer 2019

PhD rotation in Dr Celia Greenwood's lab Quantitative Life Sciences, McGill University

• Applied outlier detection algorithms to neuro-imaging data.

## **PUBLICATIONS**

1. Merenciano M, Larue A, **Groza C**, Vieira C, Rebollo R and Goubert C (In preparation) Epigenetics and Genotypic Variation: a transposable elements' perspective. book: Epigenetic and Evolution, Elsevier Press

- 2. **Groza C**, Chen X, Wheeler T, Bourque G and Goubert C (In preparation) GraffiTE: a Unified Framework to analyze Transposable-Element and Repeat-derived Structural Variants using pangenome graphs.
- 3. **Groza C**, Schwendinger-Schreck C, Cheung W, et al (2023) Pangenome graphs improve the analysis of rare genetic diseases. medRxiv. doi: <a href="https://doi.org/10.1101/2023.05.31.23290808">https://doi.org/10.1101/2023.05.31.23290808</a>
- 4. **Groza C**, Chen X, Pacis A, et al (2023) Genome graphs detect human polymorphisms in active epigenomic state during influenza infection. Cell Genomics. https://doi.org/10.1016/j.xgen.2023.100294
- 5. **Groza C**, Bourque G, and Goubert C (2023) A pangenome approach to detect and genotype TE insertion polymorphisms, In: Transposable elements, pp. 85–94 Springer. https://doi.org/10.1007/978-1-0716-2883-6\_5
- 6. Liao W-W, Asri M, Ebler J, ..., **Groza C**, et al (2023) A Draft Human Pangenome Reference. Nature. doi: https://doi.org/10.1038/s41586-023-05896-x
- 7. **Groza C**, Kwan T, Soranzo N, Pastinen, T, Bourque G (2020) Personalized and graph genomes reveal missing signal in epigenomic data. Genome Biol 21, 124. https://doi.org/10.1186/s13059-020-02038-8

## **POSTERS & TALKS**

Pangenome graphs	for the analysis	of rare genetic diseases
15min talk, American	Society of Human	Genetics Annual Meeting,
Los Angeles, CA		

Oct 2022

Human Pangenome Reference Consortium Meeting 2min talk, poster, Bethesda, MD

Oct 2022

Genome graphs detect human polymorphisms in active epigenomic state during influenza infection

Oct 2022

 $15min\ talk,\ poster\ at\ the\ 8th\ Canadian\ Conference\ on\ Epigenetics,\ Esterel,\ QC,\ Canada$ 

Genome graphs detect human polymorphisms in active epigenomic state during influenza infection

Mar 2022

15min talk, Keystone Symposia Transposable Elements, British Columbia, Canada, cancelled due to COVID19.

Leveraging read clouds to locally assemble Alu polymorphisms

Oct 2020

5min talk, poster at T2T—HPRC Towards a Complete Reference of Human Genome Diversity. Virtual.

Personalized and graph genomes reveal missing signal in epigenomic datasets

Nov 2019

Lightning talk, poster at Genome Informatics, CSHL, NY

Personalized and graph genomes reveal missing signal in epigenomic datasets

Oct 2019

Talk at ATCG, McGill University

Personalized genomes for correcting reference bias in epigenetics

Mar 2019

Talk at Montreal Genomics, Université de Montréal

Impact of personalized genomes on histone ChIP-seq peak calls

Sep 2018

Poster at the 5th Canadian Conference on Epigenetics, Esterel, QC, Canada

## **AWARDS AND OFFERS**

Japanese Society for the Promotion of Science

2023

Short-term fellowship, 400 000 JPY

2022 Charles J. Epstein Trainee Awards for Excellence in Human Genetics Research, semifinalist

2022

American Society of Human Genetics, 750 USD

Fonds de recherche du Québec

2022

Nature et technologies, 49 000 CAD, Declined

Postgraduate Graduate Scholarship - Doctoral

2022

NSERC, 63000 CAD

Graduate Excellence Award

2019

Quantitative Life Sciences, McGill University, 4500 CAD

Canada Graduate Scholarship - Master's

2019

Alexander Graham Bell, NSERC, 23000 CAD

PhD. Computer Science

University of Toronto. Declined

Winter 2019

John V Galley Scholarship McGill University, 1000 CAD

Faculty of Science Scolarship

2017

McGill University, 500 CAD

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Dean's Honor List

2017, 2018

First Class Honours

McGill University

June 2016

Dawson College

3e Prix de Chimie

Jan 2014

Concurs Scientifiques Montmorency, 100 CAD

# **OPEN SOURCE SOFTWARE**

GraffiTE 2022

Github - Software

Nextflow pipeline to detect de novo transposable element insertions.

BarcodeAsm 2020

Github - Software

Wrote a C++ program to locally assemble DNA sequences from linked-read libraries.

SeqLib 2020

Github - Accepted pull request

Added multiple features to SeqLib such as GFA output and fully adjustable assembly parameters.

## **TEACHING EXPERIENCE**

#### The 8th Kyoto Course and International Symposium

1 Mar 2023

Kyoto University

Wrote the pangenomics tutorial for the graph genome section of the 8th Kyoto Course.

#### Pharmhackaton 2018

17 Nov 2018

StudentPharma

Helped participants with command line and programming issues in a bio/chemoinformatics themed hackaton.

# **SKILLS**

Programming Python, R, Rust, Java, C/++, Scala, Haskell, LISP, LATEX Statistics Data manipulation (dplyr), presentation (ggplot2) and statistical inference.

*Bioinformatics* Write and run bioinformatic software for genome assembly, pangenomics, structural variation calling and epigenomics. Process, visualize, analyze and interpret next generation sequencing data.