



Medical Genetics Seminar Series

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“A Diagnosis for All Rare Genetic Diseases: The Horizon and Next Frontiers”

Friday, January 27, 2023
9:00 am – 10:00 am
ACH Auditorium 4th Floor

Via Zoom: <https://ucalgary.zoom.us/j/98351845596?pwd=a0RZeksrVxHd01HNHh4NGdjLzFhQT09>
Meeting ID: 983 5184 5596 Passcode: 867787

LEARNING OBJECTIVES:

- Define the paradigm shift in the approach to rare diseases since the introduction of genome-wide sequencing analysis;
- Summarize the impact, benefits and challenges of integrating bioinformatics, new technologies, global data sharing strategies and functional studies on the discovery of new causes of rare diseases;
- Propose a clinical and research workflow for how to investigate patients with undiagnosed rare genetic disease