

# 2022 CADTH Information Session

 Thursday, December 8, 2022

 1:00 p.m. to 4:00 p.m. ET

**Register**

After registering, you will receive a confirmation email with information about how to join the webinar.

1:00 p.m. to 1:10 p.m.

## Welcome

**Suzanne McGurn**, President and CEO, CADTH

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1:10 p.m. to 1:40 p.m.

## Pharmaceutical Reviews Portfolio

With a focus on the strategic pillars and guiding principles from the CADTH 2022–2025 Strategic Plan, the Pharmaceutical Reviews team will conduct a year in review for both manufacturer- and payer-initiated drug technology reviews.

**Brent Fraser**, Vice-President, Pharmaceutical Reviews, CADTH

**Amanda Allard**, Director, Pharmaceutical Reviews, CADTH

**Peter Dyrda**, Director, Pharmaceutical Policy and HTA, CADTH

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1:40 p.m. to 2:10 p.m.

## Medical Devices and Clinical Interventions Portfolio

The work of the Medical Devices and Clinical Interventions portfolio is being reshaped to better address the needs of decision-makers. This update will cover new approaches to address complex health technologies and complex information needs and to evaluate emerging technologies such as digital health.

**Lesley Dunfield**, Vice-President, Medical Devices and Clinical Interventions, CADTH

**Laura Weeks**, Director, HTA, CADTH

**Joanne Kim**, Manager, Clinical Research, CADTH

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2:10 p.m. to 2:30 p.m.

## Status Update on RWE at CADTH

An overview of initiatives to support the optimal integration of real-world evidence (RWE) into decision-making at CADTH by providing guidance on best practices for generating RWE in rare disease registries and reporting of RWE studies in submissions for HTAs and regulatory reviews.

**Nicole Mittmann**, Chief Scientist and Vice-President, Evidence Standards, CADTH

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2:30 p.m. to 2:40 p.m.

## Break

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**2:40 p.m. to 3:10 p.m.**    **Executive Strategy Update**

This update will cover CADTH's approach to respectful engagement with First Nations, Inuit, and Métis peoples; initiatives to incorporate the principles of inclusion, diversity, equity, and access (IDEA) in CADTH's work; and the new Post-Market Drug Evaluation (PMDE) program that extends CADTH's services to cover the full life cycle of health technologies.

**Heather Logan**, Executive Strategy Lead, CADTH

**Tarry Ahuja**, Director, Post-Market Drug Evaluation, CADTH

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**3:10 p.m. to 3:30 p.m.**    **CADTH Myth Busting**

CADTH's Executive Team discusses some of the common misconceptions about CADTH and its work.

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**3:30 p.m. to 3:50 p.m.**    **Open Forum**

This is an opportunity for attendees to ask questions about the material presented during the session and broader questions about anything related to CADTH and the work done at CADTH.

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**3:50 p.m. to 4:00 p.m.**    **Closing Remarks**

**Suzanne McGurn**, President and CEO, CADTH

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