

CV QUEST 2009:

Leaders and Legacies in Clinical Trials

Wednesday, October 28, 2009

ACCREDITED SYMPOSIUM

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6:30 - 7:00 a.m. - Breakfast 7:00 - 9:00 a.m. - Satellite Symposium LOCATION: The Fairmont Hotel Macdonald, 10065-100 Street, Edmonton, Alberta, Canada

MEETING ROOM: Empire Ballroom and Foyer

Learn about indispensible findings from the world's leading researchers and clinical trial groups by visiting <u>www.ccrnmd.com</u>

Co-Chairs:



Milan Gupta, MD, FRCPC Associate Clinical Professor of Medicine, McMaster University Assistant Professor of Medicine, University of Toronto Division of Cardiology, William Osler Health Centre, Brampton, Ontario



Subodh Verma, MD, PhD, FRCSC, FAHA Cardiac Surgeon

St-Michael's Hospital Associate Professor University of Toronto Canada Research Chair in Atherosclerosis Director, Traineeship in Atherosclerosis Toronto, Ontario

Faculty:



Deepak L. Bhatt, MD, FACC, FAHA Chief of Cardiology VA Boston Healthcare System Director, Integrated Interventional Cardiovascular Program Brigham and Women's Hospital and the VA Boston Healthcare System Senior Investigator, TIMI Group Boston, Massachusetts, USA

Jean-Claude Tardif, MD

Director, MHI Research Centre

Université de Montréal Research

Professor of Medicine

Chair in Atherosclerosis

Montreal Heart Institute

Université de Montréal

Montreal, Quebec



Robert M. Califf, MD

Vice Chancellor for Clinical Research Duke University Medical Center Director, Duke Translational Medicine Institute, Durham, North Carolina

7.00 0.00 - -

Eva Lonn, MD, MSc, FRCPC, FACC Professor of Medicine, McMaster University Hamilton Health Sciences General Site Hamilton, Ontario

AGENDA:

| AGEINDA: | 7:00 - 9:00 d.m. |
|--|------------------------|
| Opening Remarks | Dr. Milan Gupta |
| The TIMI Experience | Dr. Deepak Bhatt |
| The Population Health Research Institute Experience in Clinical Trials and Epidemiology | Dr. Eva Lonn |
| The Montreal Heart Institute Coordinating Centre (MHICC) and the Canadian Atherosclerosis Imaging Network (CAIN) | Dr. Jean-Claude Tardif |
| The Duke Clinical Research Institute | Dr. Robert Califf |
| Discussion | All |
| Closing Remarks | Dr. Subodh Verma |





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WHO SHOULD ATTEND: CV surgeons, cardiologists, internal medicine specialists, endocrinologists, fellows, residents and nurses interested in these disciplines.

LEARNING OBJECTIVES:

Throughout and following this session, the learner will:

- 1. Review key findings from the Thrombolysis in Myocardial Infarction (TIMI) Experience
- 2. Be familiar with developments in The Montreal Heart Institute Coordinating Centre (MHICC) and the Canadian Atherosclerosis Imaging Network (CAIN)
- 3. Consider the evolution and lessons learned from the Duke Clinical Research Institute
- 4. Examine The Population Health Research Institute Experience in Clinical Trials and Epidemiology

The Canadian Cardiovascular Research Network (CCRN) is a not-for-profit academic research organization that aims to foster basic, translational, clinical and population level research efforts, and generate new knowledge to improve cardiovascular care in Canada. An additional mandate is to develop state-of-the-art knowledge integration platforms that feature multidisciplinary and inter-specialty collaboration focusing on the future of health care delivery, research and academia-industry interface. At the present time, CCRN is actively involved in basic studies evaluating the role of adipokines in atherosclerosis, translational studies in cardiometabolic risk, clinical trials in atherothrombosis, and population based studies in minority groups.

This program will be comprised of internationally recognized principal investigators of major cardiovascular clinical trials. These four distinguished leaders will describe their clinical stream and explain how their investigations and related work have contributed to the current understanding around morbidity and mortality data as well as how their work has contributed significantly to the standard of care in the cardiovascular environment in which we practice today.

The work of these researchers touches on a variety of areas encompassing the investigation of particular therapies including acute coronary reperfusion; the use of aspirin; beta blockers; and ACE inhibitors; and the reduction of blood cholesterol, the intricacies of the RAAS, and the basics on designing and implementing a large, simple trial. Their work has challenged our past and current understanding on the efficacy of therapies such as calcium channel blockers; maintenance heparin use; nitroglycerin; and magnesium, as well as challenges us to examine the studies we design and evaluate. These investigators are well researched in strategies related to their specific area of expertise and plan to bring us 'key learnings' from various realms.

Clearly, the work of these investigators has provided us with key solutions for patient management and their data has been instrumental in influencing the treatment guidelines in Canada and in our own practices. Their knowledge and thoughtful investigation has demonstrated the expertise we should come to expect as we collectively dig deeper towards further understanding in the cardiovascular area and as we plan and execute future research. This program will benefit CV surgeons, cardiologists, internal medicine specialists, endocrinologists, fellows, residents, nurses and pharmacists interested in these disciplines.

THIS EVENT HAS BEEN MADE POSSIBLE THROUGH GRANT SUPPORT TO THE CANADIAN CARDIOVASCULAR RESEARCH NETWORK FROM THE FOLLOWING SPONSORS:

Eli Lilly Canada Inc., Merck Frosst Canada Ltd. / Schering-Plough Canada Inc. (joint venture), Sorin Group Canada, Boehringer Ingelheim (Canada) Ltd., Norvartis Pharmaceuticals Canada Inc., Pfizer Canada, Sanofi-aventis / Bristol Myers Squibb (joint venture), Li-Ka Shing King Saud University St. Michael's Hospital Collaborative Program, Solvay Pharma

> This event is an accredited group learning activity under Section 1 as defined by the Royal College of Physicians & Surgeons of Canada for the Maintenance of Certification program. This program has been reviewed and co-developed by the



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for a maximum of 2 hours of Section 1 credits.