

Joint Event

4th International Conference on

DRUG DISCOVERY, DESIGNING CHEMISTRY AND PHARMACEUTICAL ANALYSIS

&

International Conference on

BIOBETTERS AND REGULATORY AFFAIRS

June 27 - 28, 2018 | Vancouver, Canada



CONFERENCE HALL: RED CEDAR BALLROOM B

08:30-09:00 Registrations

09:00-09:10 Opening Ceremony

Keynote Forum

09:10-09:20 Introduction

09:20-10:00 Title: Biobetters vs Biosimilars: Opportunities, threats & strategic implications
Gregory K Bell, Charles River Associates, USA

10:00-10:40 Title: The evolving landscape of biobetters: Advantages & challenges of being better
Kooros Motamed-Larijani, NantBioscience Inc, USA

Panel Discussion @ 10:40-10:45

Networking & Refreshment Break 10:45-11:00 @ Red Cedar Foyer

11:00-11:40 Title: Pharmacogenomic aspects in the treatment of type 2 diabetes

Ivan Tkac, Safarik University, Slovakia

11:40-12:20 Title: Biobetters, regulatory frameworks and the 'Right to Health' in Africa

Mandla S Makhanya, University of South Africa, South Africa

12:20-13:00 Title: Treatment of human lysosomal storage disorders with Blood-Brain Barrier penetrating IgG-fusion proteins

Eric Ka Wai Hui, Armagen, USA

Panel Discussion @ 13:00-13:10

Lunch Break 13:10-14:10 @ Red Cedar Foyer

14:10-14:50 Title: Corporate integrity in the medical research and pharmaceuticals industries: The SEC's Whistleblower Program

Steven J Durham, Labaton Sucharow LLP, USA

14:50-15:30 Title: The impact of Pharmacogenomics on the future development of biological therapeutics

Michael D Winther, Genome Institute of Singapore, Singapore

Panel Discussion @ 15:30-15:35

Networking and Refreshment 15:35-15:50 @ Red Cedar Foyer

15:50-16:30 Title: Biosimilars: Challenges in safety and risk management

Mahmood Asif, Pfizer, USA

Video Presentations

16:30-17:10 Title: Quality management and quality audit according to GxP/GMP requirements

Eleonora Babayants, Galaxy Consulting, USA

Panel Discussion @ 17:10-17:20



Keynote Forum

09:20-10:00 Title: Biosimilar by approval – Biogenerics in practice?

Steinar Madsen, Norwegian Medicines Agency, Norway

10:00-10:40 Title: Recent advances in analytical techniques for the implementation of quality by design in biopharmaceutics

Yite Robert Chou, Merck & Co Inc., USA

Panel Discussion @ 10:40-10:45

Networking and Refreshment 10:45-11:00 @ Red Cedar Foyer

11:00-11:40 Title: Challenges in setting acceptance criteria for validation of analytical methods of combination products

Marika Kamberi, Abbott Vascular, USA

11:40-12:20 Title: Pharmacokinetics of Fipronil and Fipronil-sulfone in dogs after oral administration of Fipronil tablets

Yara Peluso Cid, Federal Rural University of Rio de Janeiro, Brazil

12:20-12:50 Title: Pharmacists role in helping patients with biosimilars, education, adherence and appropriate monitoring

Alan Low, BioPro Biologics Pharmacy, Canada

Panel Discussion @ 12:50-13:00

Lunch Break 13:00-14:00 @ Red Cedar Foyer

14:00-14:30 Title: Application of nanotechnology in pharmaceutical formulation design and development

Rahmatullah Hiadery, Kabul University, Afghanistan

Video Presentations

14:30-15:00 Title: Microwave assisted synthesis and evaluation of Polyvinyl alcohol-co-2-acrylamide-2-ethyl-1-propanesulfonic acid hydrogel for oral delivery of Captopril

Furqan Muhammad Iqbal, Bahauddin Zakariya University, Pakistan

15:00- 15:30 Title: GxP/GMP and its consequences for documentation and information technology systems

Eleonora Babayants, Galaxy Consulting, USA

Panel Discussion @ 15:30-15:35

Networking & Refreshment Break 15:35-15:50 @ Red Cedar Foyer

15:50-16:20 Title: Biobetters, they had better be better

William R Strohl, BiStro Biotech Consulting, USA

B2B & Networking @ 16:20-17:00

Panel Discussion & Closing Ceremony @ 17:00-17:30

