

11th SwAPP ExEx Meeting: 8 March, 2018 - Fribourg

CONSTRAINT-DRIVEN INNOVATION IN MEDICAL DEVICES

- Operated by the SwAPP Task Force Medical Device -

In the context of an increasing number of medical devices and their combination with medicinal products or biologicals, the complexity of device technology, as well as the implementation of a new EU regulation in Switzerland, the SwAPP has elaborated an Exchange Expertise event for drug and medical device professionals.

Programme

10:00 Visit of the Swiss Integrative Center for Human Health SA (SICHH, www.sichh.ch) - including lunch

13:00 Reception and Coffee break

13:15 Welcome

Mirjam Eglin, President, SwAPP; Jean-Marc Brunner, CEO, SICHH

13:30 Plenary Session

❖ **Latest Technologies in Medical Devices and Drug Delivery Devices**

Bradley Nelson, Professor of Robotics & Intelligent systems, Director of Multi-Scale Robotics Lab (ETH Zurich, CH)

❖ **Development of Innovative and User-centric Drug-Device combination products: a Pharma Perspective**

Ulla Grauschopf, Head of Device Development Europe, F. Hoffmann-La Roche Ltd (Basel, CH)

❖ **The Regulation of Medical Devices: Supporting Creativity while Ensuring Safety**

Bassil Akra, Vice-President, TÜV SÜD Product Service GmbH (Munich, D)

15:30 Coffee break

16:00 Parallel Workshops

Workshop 1: Research	Workshop 2: Development / Registration Process	Workshop 3: Focus on Clinical & Regulatory key points
Biomaterials and Interaction with Body Biological Barriers <i>Priscilla Brunetto</i> <i>R&D Project Leader, University of Fribourg (Fribourg, CH)</i> <i>Vanya Loroach</i> <i>CEO Loroach CTLS (Essertines-sur-Rolle, CH)</i>	Medicinal Products versus Medical Devices: Reaching the Market <i>Christa Spitznagel</i> <i>Sr Regulatory Affairs Manager Alpine, Allergan AG (Zurich, CH)</i> <i>Ivo Schauwecker</i> <i>Manager Clinical Services, AO Foundation (Zurich, CH)</i>	Strategies in Clinical Evaluations <i>Kathrin Abegg</i> <i>Clinical & Regulatory Affairs Manager, ISS AG (Biel, CH)</i> Person Responsible for Regulatory Compliance: Roles & Responsibilities <i>Rainer Voelksen</i> <i>Co-Chair RAPS EU Advisory Committee (Geneva, CH)</i>

17:00 Parallel Workshops (repetition)

17:55 Concluding remarks

18:00 Closing Networking Apéro

End of
early bird
Feb, 12th !

More information and registration at www.swapp.ch/events

Registration

Please register at www.swapp.ch/events/11th-swapp-exex-meeting-8th-march-2018-fribourg/1805/
Registration and payment are required prior to the meeting.

Participation Fees

SwAPP Members	CHF 300,- (Early bird seats available up to Feb, 12th) CHF 400,-
Non-Members	CHF 450,- (Early bird seats available up to Feb, 12th) CHF 550,-
Students	CHF 100,-

Groups, please contact us: swapp@swapp.ch, re: ExEx 2018 groups

Venue

SICHH, Swiss Integrative Center for Human Health SA
Blue FACTORY - Halle Bleue, Passage du Cardinal 13 B, CH- 1700 Fribourg, near to main station

Cancellation Policy

You may cancel 10 business days before the meeting and receive a full refund minus the cancellation fee of CHF 50.-
Cancellations less than 10 business days before the symposium date will not be refunded.

Accreditation

This event is accredited with 4.5 credits by SwAPP/ SGPM.

Learning Objectives

- To familiarize with the latest innovations in the field of medical devices
- To understand why a highly regulated environment can be a motor of innovation
- To learn how Pharma can successfully deal with innovative medical device products
- To understand the phases of their development (versus drugs)
- To become acquainted with some key regulatory requirements of the new EU Medical Device Regulation

Target Audience

Newcomers and experienced professionals in medical devices and combination products development, regulatory affairs, clinical affairs, market access and regulatory compliance (QP); working in the Industry, Contract Research Organisations, Competent authorities, Notified bodies, Ethics Committees or Academia.

Patronages

