

Robarts Translational Research Consortium

presents an inaugural symposium on

JAK-STAT Modulation in IBD - from Bench to Bedside

January 13, 2019

Taos, New Mexico USA

Event Sponsors









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Symposium on JAK-STAT Modulation in IBD – from Bench to Bedside

Agenda for January 13, 2019

at Sagebrush Inn & Suites, Taos, NM

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8:15 AM Welcome & Introductions

Session 1: Clinical Efficacy & Safety Co-chairs: Dr. William Sandborn & Dr. Bram Verstockt 8:45 AM State of the art: mechanism of action of JAK-STAT inhibition Dr. Massimo Gadina - National Institutes of Health, USA 9:10 AM Efficacy in inflammatory bowel diseases Dr. Severine Vermeire – KU Leuven, Belgium 9:35 AM Safety of JAK-STAT inhibition **Dr. Brian Feagan** – Western University / Robarts Clinical Trials, Canada 10:00 AM Panel discussion 10:15 AM Break **Session 2: Clinical Pharmacology & Translational Medicine** Co-chairs: Dr. Stefania Vetrano & Dr. Manuel Braga Neto

10:30 AM Drug interactions, metabolism and excretion

.30 AIVI	Drug interactions, metabolism and excretion
	Dr. Niels Vande Casteele – UC San Diego / Robarts Clinical Trials, USA

- 10:55 AM Peripheral immune system vs. mucosal immunology Dr. Azucena Salas – IDIBAPS, Spain
- 11:20 AM Predictors for response, treatment intensification, and/or discontinuation

Dr. Eoin McKinney – University of Cambridge, UK

11:45 AM Panel discussion

Noon Lunch

Session 3: Drug Design & Development Co-chairs: Dr. Dermot McGovern & Dr. Isabella Dotti 1:00 PM Improving efficacy and safety: Compound engineering vs. drug delivery Dr. Andrew Long - AbbVie, USA 1:25 PM Combination therapy redefined **Dr. Marjolijn Duijvestein** – AMC, the Netherlands 1:50 PM Overview of the pipeline and strategies for efficient drug development Dr. William Sandborn – UC San Diego, USA 2:15 PM Panel discussion 2:40 PM Break **Session 4: Breakout Session** Co-chairs: Dr. Severine Vermeire & Dr. Niels Vande Casteele 3:00 PM Working groups to identify areas of translational research that can guide treatment strategies and accelerate further drug development 4:00 PM Working group presentation & general discussion 5:00 PM Summary of proposed objectives 5:10 PM Adjourn

- Reception to Follow -