



European
Crohn's and Colitis
Organisation

Inflammatory Bowel Diseases

ClinCom Workshop



Programme 2016-2020

www.ecco-ibd.eu

7th ClinCom Workshop

Date: Thursday, February 13, 2020
Time: 08:00-11:40
Organisation: ClinCom
Target audience: Physicians, Surgeons, Paediatricians, Clinical researchers, Industry
Registration: ECCO Membership 2020 required. No onsite registration.
CME accreditation: 3 CME Credits; prerequisite: completion of the online evaluation form.

Thursday, February 13, 2020

*Schubert 1-3, Level 1, Reed Messe Wien
(as of January 15, 2020)*

08:00 - 08:05	1: Welcome & Introduction K. Gecse, Amsterdam, The Netherlands
08:05 - 09:45	Session 1: Sequencing and combination of medical IBD treatments L. Beaugerie, Paris, France
08:05 - 08:20	2: Sequencing: Does the order of treatment change the biology of IBD? B. Siegmund, Berlin, Germany
08:20 - 08:40	3: Practical approach to treatment sequencing K. Gecse, Amsterdam, The Netherlands
08:40 - 09:00	4: Concomitant immunomodulators – One size fits all? U. Kopylov, Tel Aviv, Israel
09:00 - 09:20	5: Combination of biologics, and small molecules and biologics J. Rahier, Yvoir, Belgium
09:20 - 09:45	6: Round Table B. Siegmund, Berlin, Germany K. Gecse, Amsterdam, The Netherlands U. Kopylov, Tel Aviv, Israel J. Rahier, Yvoir, Belgium L. Beaugerie, Paris, France
09:45 - 10:15	Coffee break

10:15 - 11:30	Session 2: Overcoming barriers in IIS P. Bossuyt, Bonheiden, Belgium
10:15 - 10:30	7: Overcoming challenges with your ethical committee and GDPR T. Ahmad, Exeter, United Kingdom
10:30 - 10:50	8: Feasibility vs practicality S. Sebastian, Hull, United Kingdom
10:50 - 11:10	9: Funding (Tips and tricks) E. Louis, Liege, Belgium
11:10 - 11:30	10: Round Table T. Ahmad, Exeter, United Kingdom S. Sebastian, Hull, United Kingdom E. Louis, Liege, Belgium P. Bossuyt, Bonheiden, Belgium
11:30 - 11:40	11: Summary & Closing remarks U. Kopylov, Tel Aviv, Israel
12:00 - 13:00	ClinCom Workshop Lunch Satellite Symposium

6th ClinCom Workshop

Date:	Thursday, February 15, 2018
Time:	08:00-11:40
Organisation:	ClinCom
Target audience:	Physicians, Surgeons, Paediatricians, Clinical researchers, Industry
Registration:	For registered participants only. No onsite registration
CME accreditation:	3 CME credits, upon return of an evaluation form filled in online

Thursday, February 15, 2018

*Programme overview (as of January 15, 2018) –
Lehar 1, Congress Centre, Messe Wien*

08:00 - 08:05	1: Welcome & Introduction Marc Ferrante, Leuven, Belgium
08:05 - 09:45	Session 1: Evolving endpoints in IBD clinical trials Javier Gisbert, Madrid, Spain
08:05 - 08:25	2: Patient reported outcomes Laurent Peyrin-Biroulet, Vandoeuvre-les-Nancy, France
08:25 - 08:45	3: Defining endoscopic endpoints Krisztina Gecse, Amsterdam, The Netherlands
08:45 - 09:05	4: Cross-sectional imaging Andrea Laghi, Latina, Italy
09:05 - 09:25	5: Histologic remission Roger Feakins, London, United Kingdom
09:25 - 09:45	6: How endpoints can influence trial design? Walter Reinisch, Vienna, Austria
09:45 - 10:15	Coffee break
10:15 - 11:30	Session 2: Comparative effectiveness research (CER) John Mansfield, Newcastle upon Tyne, United Kingdom
10:15 - 10:30	7: General principle Jean-Frédéric Colombel, New York, United States
10:30 - 10:50	8: What is the value of retrospective CER? Pieter Hindryckx, Ghent, Belgium
10:50 - 11:10	9: Head-to-head trials Simon Travis, Oxford, United Kingdom
11:10 - 11:30	10: Setting priorities for IBD (discussion) Jean-Frédéric Colombel, New York, United States Pieter Hindryckx, Ghent, Belgium John Mansfield, Newcastle upon Tyne, United Kingdom Elmer Schabel, Bonn, Germany Simon Travis, Oxford, United Kingdom
11:30 - 11:40	11: Summary & Closing remarks Marc Ferrante, Leuven, Belgium

5th ClinCom Workshop

Date:	March 17, 2016
Time:	08:30-12:10
Organisation:	ClinCom
Target audience:	Physicians, Surgeons, Paediatricians, Clinical researchers, Industry
Registration:	For registered participants only. No onsite registration.
CME accreditation:	3 CME credits, upon return of an evaluation form distributed during the workshop.

Thursday, March 17, 2016

Programme overview (as of February 15, 2016)

Room E102, Level 1, RAI Amsterdam

08:30 - 08:35	1: Welcome & Introduction Alessandro Armuzzi, Rome, Italy
08:35 - 09:55	Session 1: Balance safety – efficacy Vipul Jairath, London, Canada
08:35 - 08:55	2: What has meta-analysis taught us Jean-Frédéric Colombel, New York, United States
08:55 - 09:15	3: How to evaluate safety of biologics Geert D'Haens, Amsterdam, The Netherlands
09:15 - 09:35	4: Cluster randomised trials Vipul Jairath, London, Canada
09:35 - 09:55	5: How to choose your biologics in 2016 Michael Kamm, Melbourne, Australia
09:55 - 10:30	Coffee break
10:30 - 12:00	Session 2: Balance efficacy - costs Marc Ferrante, Leuven, Belgium
10:30 - 10:50	6: Methodology of cost efficacy Keith Bodger, Liverpool, United Kingdom
10:50 - 11:10	7: How to implement results of cost efficacy analysis in clinical practice? Ailsa Hart, London, United Kingdom
11:10 - 11:30	8: Comparing treatment strategies and cost effectiveness Daniel Hommes, Los Angeles, United States
11:30 - 12:00	9: From regulators to payers Elmer Schabel, Bonn, Germany (EMA) Barney Hawthorne, Cardiff, United Kingdom Alessandro Armuzzi, Rome, Italy Ailsa Hart, London, United Kingdom
12:00 - 12:10	10: Summary & closing remarks Fernando Magro, Porto, Portugal