

Role of Analytical Chemistry in Drug Product Development – Characterization and Quality Control,

March 28, AstraZeneca, Mölndal

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Dro	lim	inarv	program	
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08.30	Registration with coffee and sandwich
09.00	Introduction Chair: Mareike Lutz and Anders Karlsson
09.10	Role of Analytical Chemistry in Drug Development – a Holistic View (The T-shaped analytical chemist) Anders Karlsson, AstraZeneca, Mölndal
09.40	Early Stage Therapeutic Projects – Characterization and Quality Control Patrik Petersson, Ferring, Copenhagen
10.10	Coffee
10.35	Build Measurement Procedures to Support Continuous Manufacturing and Real Time Release Testing (STST) Peter Hamilton, AstraZeneca Macclesfield
11.05	Characterization and Quality Control Testing of Next Generation Drugs – an Introduction to Antisense Oligonucleotides Daniela Balas, AstraZeneca, Mölndal
11.35	Lunch
12.35	Analytical TT, Getting Value out of your CDMO Per-Ivar Corin, Recipharm, Stockholm
13.05	Analytical Chemistry at a Commercial Site – Release Testing and Support Manufacturing Magnus Liljenberg, AstraZenca, Södertälje
13.35	Quality Testing of Excipients Staffan Schantz, AstraZeneca, Gothenburg
14.05	Coffee
14.20	Use of Analytical Chemistry in Development and Commercial Space <i>Anders Johansson and Petros Tesfai</i> , J&J McNeil AB, Helsingborg
14.50	Review of Development and Validation of Quality Control Testing Procedures from an Agency Perspective Annika Ridell, Swedish Medical Products Agency, Uppsala
15.20	Online Analysis of Cell Culture Media Hans Christian Høiberg, Agilent, Gothenburg
15.50	Open discussion focus on highlights from the day
16.30	End of the day

Exhibitors







