

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

Preliminary program

- 08.30 *Registration and coffee*
- 09.00 Introduction
- 09.10 **Role of analytical chemistry in drug development – a holistic view
(The T-shaped analytical chemist)**
Anders Karlsson, AstraZeneca
- 09.40 **Early stage therapeutic projects – Characterization and Quality Control**
Patrik Petersson, Ferring
- 10.10 *Coffee*
- 10.35 **Quality Control testing at a CRO**
Lecturer to be confirmed
- 11.05 **Quality Control testing in the commercial manufacturing area**
Jun Yamazaki, AstraZeneca
- 11.35 *Lunch*
- 12.35 **Quality Control testing for continuous manufacturing and Real Time Release Testing
(RTRT)**
Öivind Holte, Medical Products Agency, Oslo, Norway
- 13.05 **Characterization and Quality Control testing of next generation drugs – synthetic
oligonucleotides and mRNA**
- 13.35 **Quality testing of excipients**
Staffan Schantz, AstraZeneca
- 14.05 *Coffee*
- 14.20 **Quality Control testing using monograph procedures**
Lecturer to be confirmed
- 14.50 **Development, validation and review of Quality Control testing procedures – current and
future state**
Lecturer to be confirmed
- 15.20 **Panel discussion (one/two themes to discuss e.g. ICHQ12 and 14, Analytical Risk
assessment (ARA), Critical Quality Attributes (CQA) and Enhanced Control Strategy)**
- 16.00 – 16.10 **Wrap-up**