BRITISH SOCIETY OF TOXICOLOGICAL PATHOLOGY

CONTINUING EDUCATION SYMPOSIUM 1:
TOXICOLOGY and PATHOLOGY
Cambridge
Tuesday 26th – Thursday 28th March 2019

Approximately 17.25 hours of educational activity will be recorded on Attendance Certificates
Approved by the Royal Society of Biology for the purposes of CPD, this event may be counted as 51 credits

PROVISIONAL PROGRAMME as at 6th February 2019

Tuesday 26th March 2019	
08.45 - 09.15	**Registration**
09.15 - 09.30	**Introduction by the Scientific Organiser** *Ian Taylor, Envigo, UK*
09.30 - 10.15	**Descriptive pathology skills & data recording in toxicological & experimental pathology** *Cheryl Scudamore, Envigo, UK*
10.15 - 10.45	**Break**
10.45 - 12.00	**Immunological techniques in toxicological pathology** *Stewart Jones, AstraZeneca, UK*
12.00 - 13.00	***In situ* techniques using nucleic acids** *Kevin Randall, Medicines Discovery Catapult, UK*
13.00 - 14.00	**Lunch**
14.00 - 14.45	**Approaches to the analysis & interpretation of toxicology clinical pathology data** *Peter Cotton, AstraZeneca, UK*
14.45 - 15.15	**Break**
15.15 - 16.45	**Workshop/Case Studies: Haematology & clinical chemistry** *Peter Cotton, AstraZeneca/Ian Roman, GlaxoSmithKline, UK*

Wednesday 27th March 2019	
08.45 - 09.30	**Safety pharmacology in drug discovery & development/case studies** *Speaker - tbc*
09.30 - 10.30	***In vivo* imaging - overview/case histories** *Juliana Maynard, Medicines Discovery Catapult, UK*
10.30 - 11.00	**Break**
11.00 - 12.00	**Non-animal approaches to safety assessment** *Carl Westmoreland, Unilever, UK*
12.00 - 13.00	**Lunch**
13.00 - 14.00	**Pharmacogenetics & drug toxicity** *Ann Daly, Institute of Cellular Medicine, Newcastle University, UK*
14.00 - 15.00	**Animal models in drug development & discovery** *Peter Clements, GlaxoSmithKline, UK*
15.00 - 15.30	**Break**
15.30 - 16.15	**Animal models in drug development & discovery – case presentations** *Peter Clements, GlaxoSmithKline, UK*
16.15 - 17.15	**Use of nonclinical data for risk assessment & risk management in the clinic: Dose selection, dose escalation & monitoring** *Jim Bush, Covance Clinical Research Unit, UK*

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Thursday 28th March 2019	
08.45 - 09.45	**Reproductive toxicology** *Louise Youngs, Syngenta, UK*
09.45 - 10.45	**Is it adverse? Introduction, case histories & interactive discussion – Part I** *Jan Klapwijk, GlaxoSmithKline, UK*
10.45 - 11.15	**Break**
11.15 - 12.15	**Nonclinical Safety Assessment of Biologics** *Lolke de Haan, Medimmune, UK*
12.15 - 13.15	**Lunch**
13.15 - 14.00	**Non-pharma – adversity – case histories** *John Foster, Consultant Pathologist, UK*
14.00 - 15.00	**A regulatory perspective on pathology data** *David Jones, MHRA, UK*
15.00 - 15.30	**Break**
15.30 - 16.45	**Is it adverse? Case histories & interactive discussion – Part II** *Jan Klapwijk, GlaxoSmithKline, UK*
16.45 - 17.00	**Closing remarks by the Scientific Organiser**