

# drug safety FORUM

Making Safety  
Part of Drug Design

Magdalene College, Cambridge

**October 12th 2023**

9am - 5pm



Time	Title	Speaker
<b>9-945</b>	<b>Arrival and Registration - Coffee and Pastries</b>	
945-10	Introduction	Professor Ruth Roberts, ApconiX
<b>Opening Keynote</b>		
10-1030	Reducing the Risk of Mitochondrially Active Compounds in Medicines Development	Dr Jon Lyon, Director of Preclinical Toxicity and Mitochondrial Network Lead, GSK
<b>Morning Session</b>		
1030-11	Assessing Cardiac Ion Channel Activity in the Drug Discovery Cascade	Dr Mike Morton, ApconiX
11-1130	Making formulation part of drug design: case studies	Dr Linette Ruston, Director, ADME and Modelling Sciences, SEDA Pharmaceuticals
1130-12	The role of Target Safety Assessments (TSAs) in de-risking drug projects	Dr James Sidaway, ApconiX
<b>12-13</b>	<b>Lunch Break</b>	
<b>Afternoon Session</b>		
13-1330	Interfacing DMPK and Safety in Drug Design: Concentration effect relationships	Dr Kevin Beaumont, Senior Director, DMPK, AstraZeneca
1330-14	Developing an ion channel seizure panel	Dr Kim Rockley, ApconiX
14-1430	Designing out secondary pharmacology hits	Dr Mark Anderton, Senior Principal Scientist, MSD
<b>1430-15</b>	<b>Coffee/Tea</b>	
15-1530	State-of-the-art cellular models in de-risking drug design	Dr Colin Brown, Chief Innovation Officer, Newcells Biotech
1530-1600	Making safety part of drug design – a regulatory perspective	Dr David Jones, ApconiX
1600	Panel discussion	Dr David Jones Dr Richard Knight Dr Anthony Holmes Dr Muireann Coen
<b>1630</b>	<b>Close</b>	