

# PROGRAM IN BRIEF

**THURSDAY 27 MAY 2010**

<b>8.00 - 8.30</b>	<b>Registration, Welcome Coffee &amp; Tradeshow</b>	
8.30 - 9.00	Opening of Sydney ARCS ASC - Research and Development Meeting	
9.00 - 11.00	Opening Plenary: Debating dollars and drugs (T1)	
<b>11.00 - 11.30</b>	<b>Morning Tea &amp; Tradeshow</b>	
11.30 - 1.00	Cross-Functional	Therapeutic Area Training for Clinical Trials (T6)
	Clinical Research	Insurance and Indemnity for Clinical Trials (T2)
	Clinical Research	Phase IV/Post-Marketing studies in Devices and Pharma - Are They the Same Beast? (T4)
	Health Economics	Introduction to Health Economics for Non-Economists (T5)
	Asia Pacific	Cultural Diversity and Ethnopharmacology in Global Clinical Trials: Asia Pacific Perspective (T3)
<b>1.00 - 2.00</b>	<b>Lunch &amp; Tradeshow</b>	
2.00 - 3.30	Regulatory/Cross over	Diligence in Early-phase Biotech Development: Preclinical, Regulatory and Clinical Strategies (T10)
	Clinical Research	Theranostics - Moving Closer to Personalised Medicine (T8)
	Workplace Management	Linking Employee Engagement to Effective Deliverables in Clinical Trials - A Perspective (T9)
	Pharmacovigilance	Optimising Post-marketing Surveillance (T7)
	Asia Pacific	Getting it Right! Avoiding Audit Findings in Asia (T11)
<b>3.30 - 4.00</b>	<b>Afternoon Tea &amp; Tradeshow</b>	
4.00 - 5.30	Cross-Functional	Advances in the Design of Dose-Response Studies (T15)
	Clinical Research	Research in Public Hospitals: Advances in Streamlining the Eastern States (T16)
	Workplace Management	Effective Partnering: Pharma, Biotech and CROs (T14)
	Pharmacovigilance	Clinical Aspects of Safety Reporting Including Investigator Initiated Study (T12)
	Asia Pacific	Conducting Clinical Trials in Large Asian Markets (T13)
<b>6.00 - 10.00</b>	<b>Networking Function</b>	

P  
R  
O  
G  
R  
A  
M  
  
I  
N  
  
B  
R  
I  
E  
F

**FRIDAY 28 MAY 2010**

<b>8.00 - 9.00</b>	<b>Registration, Welcome Coffee &amp; Tradeshow</b>	
9.00 - 10.30	Clinical Research	How to Manage Clinical Trial Agreements More Effectively (F1)
	Clinical Research	Lean Six Sigma in Clinical Trials (F3)
	Workplace Management	Performing in Spotlight (F2)
	Workplace Management	Risk Management - Clinical Trial Phase and Beyond (F5)
	Cross-Functional	Practical Workshop on the Implementation of the New MA Code of Conduct (F4)
<b>10.30 - 11.00</b>	<b>Morning Tea &amp; Tradeshow</b>	
11.00 - 12.30	How to Manage Clinical Trial Agreements More Effectively (continued)	
	Lean Six Sigma in Clinical Trials (continued)	
	Performing in Spotlight (continued)	
	Risk Management - Clinical Trial Phase and Beyond (continued)	
	Practical Workshop on the Implementation of the New MA Code of Conduct (continued)	
<b>12.30 - 1.30</b>	<b>Lunch &amp; Tradeshow</b>	
1.30 - 3.00	Clinical Research	How Technology is Empowering Patients, and What this Could Mean for Clinical Research (F6)
	Clinical Research	Managing Trials in the Asia pacific from Australia (F9)
	Workplace Management	Hybrid Models of Conducting Clinical Trials (F7)
	Health Economics	An Overview of Systematic Reviews and Meta-Analysis (F8)
	Cross-Functional	TBC (F10)
<b>3.00 - 3.30</b>	<b>Afternoon Tea &amp; Tradeshow</b>	
3.30 - 5.00	Closing Session: Express Learning & Wine Tasting	