

PROGRAM IN BRIEF

MONDAY 13 SEPTEMBER 2010

9.00 - 9.45	Registration, Welcome Coffee & Tradeshow	
	Opening of Canberra ARCS ASC - Regulatory and Reimbursement Meeting	
9.45 - 12.00	Opening Plenary: Debating dollars and drugs (T1)	
12.00 - 1.00	Lunch & Tradeshow	
1.00 - 3.00	Regulatory	Introduction to the electronic Common Technical Document (Part I) (M2)
	Regulatory	TGAs Business Process Reforms a Year Later (M3)
	Health Economics	How Can You Use Observational Data? (M4)
	Health Economics	Update from the PEB (M5)
	Medical Devices	The New ARGMD and the Key areas of Regulations/Whats New in Medical Devices (M6)
3.00 - 3.30	Lunch & Tradeshow	
3.30 - 5.30	Regulatory	Complementary Medicines. Developments Listing and Pitfall (M7)
	Regulatory	Pre-submission Meetings Around the World (M8)
	Health Economics	Health Economics 101 (M10)
	Health Economics	Cost-Utility, Analysys in PBAC Submissions (M11)
	Medical Devices	New IVD Regulations (M9)
6.00 - 11.30	Gala Dinner	

TUESDAY 14 SEPTEMBER 2010

8.00 - 9.00	Welcome Coffee & Tradeshow	
9.00 - 10.30	Regulatory	Introduction to the electronic common Technical document (Part II) (T5)
	Regulatory	Labelling of Medicines - Best Practice (T2)
	Health Economics	An Overview of Systematic Review and Meta-analysis (T4)
	Health Economics	Pricing Reform Legislation - Practical Understanding (T6)
	Medical Devices	Medical Devices from Innovation to Reimbursement (T3)
10.30 - 11.00	Morning Tea & Tradeshow	

11.00 - 12.00	Plenary: TBC (T7)	
12.00 - 1.00	Lunch & Tradeshow	
1.00 - 3.00	Regulatory	Inside the Mind of an Evaluator (T10)
	Asia Pacific	Recognition of Data by Various Countries in Asia (Mutual Recognition of Data Across Asian Countries (T9)
	Health Economics	Pricing Issues for Drug Listed on the PBS (T8)
	Pharmacovigilance	Risk Management in Australia - Local and Global view. Are you ready for RM in your organisation? (T11)
	Cross-Functional	Consumer Preferences (T12)
3.00 - 3.30	Afternoon Tea & Tradeshow	
3.30 - 5.00	Regulatory	Navigating the Cultural Divides. Working with O/S Regulatory Agencies (T13)
	Regulatory	OTC Medicines (T15)
	Health Economics	Harmonising HTA Approaches (T16)
	Pharmacovigilance	Biosimilars: Beyond Bioequivalence. Development, Registraion and Safety Monitoring of Follow-on Biologics (T14)
	Cross-Functional	e-Health (T17)
	Regulatory	Clinical Evaluation for Medical Devices: Implementing New Regulations 2010 in Europe in Your Global Product Development Strategy (T18)