

DRAFT SCHEDULE (11 Feb 2019)

<u>TIME</u>	<u>AGENDA ITEM</u>
9.00am	WELCOME TO SYMPOSIUM – MC - Danny Vadasz
9.10am	GOVERNMENT STATEMENT – Minister or representative
9.15am	KEYNOTE ADDRESS - Pip Brennan (HCCWA) Melissa Fox (HCQ)
9.45am	LESSONS LEARNED – The lived experience of consumers (various)
10.30am	GOVERNMENT’S RESPONSE TO THE SENATE INQUIRY – Senator Rachel Siewert
11.00am	Morning tea
11.20am	REGULATING MEDICAL DEVICES – John Skerritt TGA
11.45am	PANEL DISCUSSION – TGA, ACSQHC, IJC, Politician, Consumers
12.30pm	Lunch
1.30pm	WORKSHOP INTRODUCTIONS
1.35pm	Treatment pathways – Producing a Scorecard for comparing state responses (service quality) and best practice guidelines around consumer led design solutions.
1.50pm	Regulation and policy - Producing a scorecard for evaluating progress on the Government’s response to the Senate Inquiry. Also looking more broadly at questions of device regulation, registration, adverse event reporting, credentialing, complaints processing, product withdrawal and regulation of device manufacturers.
2.05pm	Informed consent – Moving the locus of patient centred care from shared information to decision making tools that enable informed consent based on a full disclosure of risks, benefits and options.
2.20pm	Governance – Affiliation between various mesh bodies, relationship between states and federal jurisdictions. Basic structural requirements of accountability, process etc.
2.35pm	CONCURRENT WORKSHOPS
3.35pm	AFTERNOON TEA
3.50pm	PLENARY REPORT BACKS AND RECOMMENDATIONS
4.50pm	CLOSING REMARKS
5.00pm	CLOSE