

REGISTRY-BASED CLINICAL TRIALS CONCEPT DEVELOPMENT WORKSHOP

From pragmatic idea to implementation



Program

Friday 25 August, 9.00am - 3.30 pm

Graduate House, Ian Potter Room,

220 Leicester St, Carlton, VIC 3053

Aims:

- To enhance the clinical trial workforce's capacity to conduct a higher number of registry-based clinical trials and elevate the quality and standard of care in registry trial protocols.

Learning objectives: By the end of the session, participants will have had the opportunity to:

- Compare the key differences between conventional trials and Registry-based Clinical Trials
- Explore what data registries are and how they tie into Registry-based Clinical Trials.
- Consider the benefits of Registry-based Clinical Trials and know how they can be easy and cost-effective to run when you are armed with the right knowledge.
- Consider the application of the methodology behind registry-based clinical trials.
- Consider how consumer input can shape a registry-based clinical trial protocol to meet patient needs and standard of care.
- Discuss how potential ideas can be developed into a protocol for registry-based clinical trials.
- Network with multidisciplinary professionals across the registry-based clinical trial industry.

Learning outcomes: By the end of the session, participants will be able to:

- Critically reflect on what knowledge and considerations are needed to develop an idea into a registry-based clinical trial protocol and ensure it is feasible for registry-based clinical trials.
- Evaluate an area of need or develop possible ideas, from within your institution, for where Registry-based Clinical Trials could be beneficial to improve the standard of care.

Program

| Time | Topic | Speaker |
|---------|--|---|
| 9.00am | Arrival and registration (light refreshments) | |
| 9.30am | Session 1 | Chairs: Dr Vanessa Wong & Dr Shehara Mendis |
| 9.30am | Welcome and introduction | Dr Vanessa Wong |
| 9.45am | Barriers and Enablers of Randomised Registry-based Clinical Trials | Bill Karanatsios |
| 10.20am | Radiation Oncology Studies and Registry-based Clinical Trials | A/Prof Gavin Wright |
| 10.35am | Surgical Oncology Studies and Registry-based Clinical Trials | A/Prof Steven David |
| 10:50am | Radiation & Surgical Oncology Registry-based Clinical Trials Discussion | A/Prof Gavin Wright & A/Prof Steven David |
| 11.05am | Morning tea | |
| 11:35am | Session 2 | |
| 11:35am | Biostatistical Considerations | Dr Wei Hong |
| 12:05pm | Ethical Considerations in the Context of Registry-based Clinical Trials | Dr Evelyn Yip |
| 12:30pm | Budgeting for Registry-based Clinical Trials | Maria Edmonds |
| 12:55pm | Lunch | |
| 1:40pm | Session 3 | |
| 1:40pm | Site Perspectives: A Chat | Facilitator: Eleonora Kay |
| 2:10pm | Developing an Idea Into a Registry-based Clinical Trial protocol: Workshop | |
| 3:00pm | Panel Discussion | Facilitator: Dr Shehara Mendis |
| 3.30pm | Close | |



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