

PREDICT Research Education Plan 2024

MONTH	TOPICS	PRESENTER
7th March 11am-12noon (AEDT)	Different types of research designs <ul style="list-style-type: none"> • Observational- pro vs retro vs cross section • RCT- randomisation schedules, blinding • Adaptive studies • Step wedge 	FRANZ BABL
	Different types of research designs cont. <ul style="list-style-type: none"> • Audit / QI • Qualitative • Mixed methods • Implementation designs 	EMMA TAVENDER (rescheduled)
	Consumer involvement	CATE WILSON (rescheduled)
9th April 11am-12noon	Grant writing <ul style="list-style-type: none"> • Local grants • Different types of grants available • National and international grants • How to get started/elements • How to write a grant 	SIMON CRAIG
	Budgets <ul style="list-style-type: none"> • Invoicing • Staff costs • Strategies on budgeting • How to stretch the research dollar 	CATE WILSON
	Contracts <ul style="list-style-type: none"> • Different type of contracts • MIA vs CTRA vs DTA • How to complete, who to contact • What parts of the contract to pay closer attention to 	CATE WILSON
24TH May 11am–12noon (tbc)	Different types of research designs cont. <ul style="list-style-type: none"> • Audit / QI • Qualitative • Mixed methods • Implementation designs 	EMMA TAVENDER
	Consumer involvement	CATE WILSON

<p>10th Jul 11am-12.30pm</p>	<p>Overview of process from funding to get a study up and running (from design- protocol- consultation (Maori/Atsi)- ethics – local governance – CTRAs- site initiation visit- site activation) monitoring, closure</p>	<p>SHANE GEORGE</p>
	<p>Role of Principal Investigator</p> <ul style="list-style-type: none"> • Different for each study • Safety reporting • Site activation • Delegation • Expectations 	<p>MEREDITH BORLAND</p>
	<p>Setting up a research team and department</p> <ul style="list-style-type: none"> • Rostering staff • Buy in from clinical staff within and outside ED, negotiation skills • How to effectively work with your RAs? 	<p>NATALIE PHILLIPS</p>
<p>30th Aug 11am-12.30pm</p>	<p>HREC</p> <ul style="list-style-type: none"> • Key documents, National Statement, ICH, TGA • Processes and structure • Multisite vs single centre • LNR, governance only • Audits (negligible risk) and QUI • Ethics materials • Types of consent • Difference between states/ between AUS/NZ • Approvals • Annual requirements- project reports, safety reports 	<p>AMANDA WILLIAMS</p>
	<p>Central Filing</p> <ul style="list-style-type: none"> • Trial master file • Investigator site file • What is expected to be in each in detail 	<p>KATE KLEIN</p>
<p>Oct (TBC)</p>	<p>Study governance (MOO)</p> <ul style="list-style-type: none"> • Compliance reporting • Version control • Data dictionary 	<p>AMANDA WILLIAMS</p>

	<ul style="list-style-type: none"> • Source documents • Delegation logs • “how to put together a study binder” • Document storage after a trial • Monitoring 	
	Safety reporting and AE screening <ul style="list-style-type: none"> • Outline • Expectations • Role 	ELLIOT LONG
Dec (TBC)	Enhancing recruitment	SHARON O’BRIEN/NATALIE PHILLIPS
	Pearls and pitfalls/Tricks of the trade/FAQs	SHARON O’BRIEN/NATALIE PHILLIPS