Tips for completing an Ethics Application



- Find out the deadlines for CDU-HREC and start your application well in advance. The submission deadlines and meeting dates are available on the <u>CDU website</u>
- Focus on high quality writing and presentation –precise and concise
- Use standard, clear English for intelligent lay person in the application, information to participants, informed consent form and in data collection (e.g. interview guides, questionnaires)
- The ethics application and your Information for Participants Sheet and Consent Form must align and be coherent
- The information for participants needs to be informative including a statement that participation is voluntary, risks and benefits are addressed fully, participant expectations are clearly identified, and language and content is targeted to each cohort.
- Seek out guidance and templates from the <u>CDU-HREC webpages</u>.





Common Pitfalls

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- 1. Poor Research Design
- 2. Lack of information

3. Inconsistency



4. When a researcher proposes to do something that directly goes against the national ethical standards for research

CDU-Human Research Ethics Application Checklist



1.	Information Sheet for Participants (ISP) and Consent Form		
	(Please review the CDU-HREC Guidelines for ISP and Informed Consent.		
	Please use the <u>CDU-HREC Template for ISP and Informed Consent</u>)		
	Current CDU letterhead with full contact details		
	Full official title of the project included on both the ISP and Consent form		
	☐ Includes the statement: "This Is yours to keep", or equivalent		
	If HDR project, the research supervisor is introduced as the PI and is accountable for all details of the study, and the student is introduced as the student (NOT the PI)		
	Include contact details of main researchers		
	Identification of all possible risks		
	Written in clear, succinct, plain English and directed towards the participants		
	Includes statement to direct concerns and complaints to CDU -HREC with correct contact details		
	☐ Includes "This Means You Can Say NO" statement / Consent for ALL procedures		
	Ability for participant to consent to ALL data collection procedures individually (e.g. audio		
	recording videotaping, interviews)		
	NB: It is strongly recommended that you use the Participant Information Sheet Template found here		
2.	Application		
	All relevant answers have beencompleted		
	Research protocol/plan attached, if proposal has not been previously peer reviewed (e.g., through Confirmation of Candidature of research by assessors for a funding body that has awarded a grant for the project)		
3.	HDR candidates		
	Primary research supervisor is the Principal Investigator (PI), HDR student is listed as student		
	Confirmation of Candidature (CoC) approval and date included.		
4	Research team		
	☐ Contact details and qualifications are complete including staff and student numbers. ☐ The PI is appropriately qualified and experienced, and is a CDU staff member or has an		
	honorary appointment		
5.	First Nations Research		
	Section 13 of the <u>Application Form</u> has been completed.		
	☐ A signed ATSIRA is attached to the application.		
	If there is no ATSIRA, ensure that you include evidence of either:		
	☐ A letter or written confirmation from the community and/or relevant local organisation that		
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		details of individuals who will support the study, intellectual property, authorship, timescale, expectations and sustainability?	
		OR	
		Evidence of a reference group that has been consulted that includes First Nations people	
6.	Section 15 –	- Application Declaration	
		Project title included in all applicable areas	
		Risk level checked	
		Signed and dated by PI and (student if HDR project)	
7.	Section 16 – Authorising Officer Declaration		
		Project title to be included on the declaration	
		All relevant boxes checked	
		Application has been reviewed, authorised, signed and dated by the College Dean, orduly appointed agent (Assistant Dean for Research, PVC or University Secretary)	
8.	Final Steps		
		Have any comments made by the Authorising Officer been addressed	
8.	Attachments		
		Information Sheet for Participants	
		Informed ConsentForm	
		Additional attachments (e.g. ATSIRA for First Nations research applications, support letters,	
		research protocol, permits, *Ochre card, example survey,)	
		* valid Ochre card/Working with Children certificate (if project involves children under 18yrs	