## **HREC Proposal Check Sheet**

Project no:											
Project title: Conflicts of interest?							Υ	/	N		
Assessor name and position:					If yes, please specifiy:						
1. Research team						4. Privacy and confidentiality					
Appropriately qualified		/	N	/	NA	Appropriate de-identification	Υ	/	N	/	NA
Appropriate supervision		/	N	/	NA	Safe storage and disposal	Υ	/	N	/	NA
Adequate funding / resources	Υ	/	N	/	NA	Appropriate length of storage	Υ	/	N	/	NA
2. Project						5. Feedback and publication					
Research plan clear	Υ	/	N	/	NA	Appropriate to participants	Υ	/	N	/	NA
Research of significance		/	N	/	NA	Appropriate to communities	Υ	/	N	/	NA
Adequate literature review		/	N	/	NA	Significant conflicts of interest?	Υ	/	N	/	NA
Appropriate methodology		/	N	/	NA	Letters of support?	Υ	/	N	/	NA
3. Risks and benefits to participant	ts					6. Patient information form	Υ	/	N	/	NA
Risk of physical harm	Υ	/	N	/	NA	Letterhead, title	Υ	/	N	/	NA
Risk of psychological harm	Υ	/	N	/	NA	Identifies researchers properly	Υ	/	N	/	NA
Risk of other harm	Υ	/	N	/	NA	Contains relevant information	Υ	/	N	/	NA
Adequate harm minimisation	Υ	/	N	/	NA	Expressed in plain language	Υ	/	N	/	NA
Overly onerous on participants	Υ	/	N	/	NA	Assurance of confidentiality	Υ	/	N	/	NA
Appropriate safeguards for						This is for you to keep	Υ	/	N	/	NA
vulnerable groups *	Υ	/	N	/	NA	Concerns and complaints	Υ	/	N	/	NA
Benefits to participants (direct)	Υ	/	N	/	NA	Risks and benefits	Υ	/	N	/	NA
Benefits to others (indirect)	Υ	/	N	/	NA	7. Consent form	Υ	/	N	/	NA
Benefits proportionate to risks	Υ	/	N	/	NA	Letterhead, title, researchers	Υ	/	N	/	NA
						This means you can say NO	Υ	/	N	/	NA
*Including: children, not mentally or questionably competent, non-English speaking,					Consent for all procedures	Υ	/	N	/	NA	
Aboriginal or Torres Strait Islander, vulnerable or dependent relationship				Space for witness, interpreter	Υ	/	N	/	NA		

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### Nature of study:

Quantitative			/	Ν
	Audit (retrospective)	Υ	/	N
	Cross-sectional	Υ	/	N
	Case control	Υ	/	N
	Cohort	Υ	/	N
	Randomized controlled trial	Υ	/	N
Qualitative			/	N
	Questionnaire or survey	Υ	/	N
	Interviews	Υ	/	N
	Focus groups	Υ	/	N

#### Procedures to be used:

Administration of drugs / therapeutic agents	Υ	/	N
Invasive procedures	Υ	/	N
Collection of blood / human tissue samples	Υ	/	N
Involves human genetic information	Υ	/	N
Involves stem cells / embryos	Υ	/	N
Collection of sensitive information	Υ	/	N
Access to medical or other records	Υ	/	N
Data linkage	Υ	/	N
Audio or video taping	Υ	/	N

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Additional com	iments:				
OVERALL ASSES	SSMENT				
		Minimal	Moderate	High	
Risks to particip	pants				
Benefit to participants					
Overall benefits	S				
		Adequate	Minor concerns	Major concerns	
Informed conse	ent process				
Procedures for confidentiality	maintaining				
ACTION:	Approved				
	Conditionally approved – to chair				
	Conditionally approved – to fast-track				
	Not approved				