3. Inclusions

Does your study involve the use of Guthrie cards?

- Yes
- No

Is your study:
- funded by the Health Research Council, and
- not able to be reviewed by an HRCEC-approved university ethics committee?

- Yes
- No

Does your application involve the establishment or maintenance of a tissue bank?

- Yes
- No

2. Exemptions

Does your study involve a medical device that is (or would be) classified as a low risk (class I) medical devices by Australia’s Therapeutic Goods Administration?

- Yes
- No

Is your study a minimal risk observational study?

- Yes
- No

Is your study an audit or related activity?

- Yes
- No

Does your audit or related study involve the use, collection or storage of human tissue without consent?

- Yes
- No

Does a statutory exception to the need to gain informed consent apply to this use, collection or storage?

- Yes
- No

Is your study an observational study that is to be conducted for the purposes of an educational qualification at Masters level or below?

- Yes
- No

1. Main criteria

Will your research use or create a human gamete, a human embryo or a hybrid embryo?

- Yes
- No

Does your study involve human participants recruited in their capacity as:
- consumers of health and disability support services, or
- relatives/caregivers of such consumers, or
- volunteers in clinical trials?

- Yes
- No

Does your study involve the use, collection or storage of human tissue (as defined by the Human Tissue Act 2008)?

- Yes
- No

Does one or both of the exceptions at paras 27.2.1 and 27.2.2 of the SOPs apply to this use, collection or storage?

- Yes
- No

Does your study involve the use or disclosure of health information (as defined by the Health Information Privacy Code 1994)?

- Yes
- No

Does one or both of the exceptions at paras 27.3.1 and 27.3.2 of the SOPs apply to this use or disclosure?

- Yes
- No

Your study must be approved by the Ethics Committee on Assisted Reproductive Technology (www.ecart.health.govt.nz).