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Australasian Oncofertility Registry: uptake and utilisation of fertility preservation, reproductive health after cancer treatment, and future use and complications of assisted reproductive technologies in cancer survivors

ACTRN12615000221550

Status	RECRUITING
Sponsor	University of New South Wales
Enrollment	2,500 participants

Plain Language Summary

This study is establishing Australia and New Zealand's first fertility data registry for cancer patients. It will track whether cancer patients had fertility preservation (such as freezing eggs or sperm) before treatment, how their fertility was affected by treatment, and whether they were able to start a family afterward. This information will help improve fertility care for future cancer patients.

You may be eligible if:
- You have been diagnosed with any type of cancer
- You are under 45 years old (cancer patients aged 13–45 diagnosed with cancer, or children aged 0–12 referred for fertility preservation)
- You were diagnosed within the last 2 years

You may NOT be eligible if:
- You are a non-cancer patient treated with drugs that affect the reproductive system
- You are a parent or sibling donating fertility preservation for another person
- You do not consent to data collection
- English is not your first language and no interpreter is available
- You have not been diagnosed with cancer
- You are not of childbearing age

Talk to your doctor about whether this trial might be right for you.

Key Eligibility Criteria

Inclusion (6)

- Update
- a. All patients (any cancer diagnosis) diagnosed with cancer
- b. Cancer patients aged 13-45 who diagnosed with cancer
- c. Cancer patients aged 0-12 years of age who are referred for fertility preservation only.
- Those paediatric patients who are not referred for FP will not be eligible.

... and 1 more (see full listing online)

Exclusion (7)

- a. Non-cancer patients who are treated with gonadotoxic drugs will not be registered in phase 1.
- b. Parents/siblings who undergo fertility preservation for a child/sibling.
- c. Patients who do not consent to data being collected and reported.
- d. Patients whose first language is not English when we cannot get an appropriate interpreter to provide consent for the study.

- e. Patients who do not consent or assent to the registry

... and 2 more (see full listing online)

Locations (2 total)

Christchurch, New Zealand
Auckland, New Zealand

https://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?ACTRN=ACTRN12615000221550