

Data Request Form

For access to NSW Health Pathology data for purposes other than direct patient care and management

Please complete and submit ALL pages of this application form

Email this form to: NSWPath-RGO@health.nsw.gov.au

APPLICANT DETAILS

Applicant name:

Applicant email:

APPLICANT'S AFFILIATION / ORGANISATION

Applicant telephone/mobile:

NSW Health Pathology employee:

YES - NSW Health Pathology
Laboratory name:

NO - Local Health District or Hospital or University or
Organisation name:

PROJECT NATURE

Quality Improvement / Health service planning or management / Contractor services

Please provide a copy of the Quality Improvement determination letter email from your local research office

OR

Research

Below, please answer 'Yes' if you have ethics approval or 'No' if ethics is not yet approved

YES

Ethics approval number:

HREC office (that issued ethics approval):

Please provide a copy of:

- *Human Research Ethics Application (HREA)*
- *Ethics approval letter*

NO: *ethics to be confirmed or applied for*

PROJECT OUTLINE

- *Please provide a brief outline of your project. This may include explaining the objective/aim, the type of patients to be studied, and the medical condition of interest.*
- *If you have a protocol or method for your project, please attach this to your application.*

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DATA VARIABLES

To help us better assist your enquiry, please indicate the types of data you require:

PATIENT:

- | | | |
|--|--|------------------------------------|
| <input type="checkbox"/> First name | <input type="checkbox"/> Sex | <input type="checkbox"/> Address |
| <input type="checkbox"/> Surname | <input type="checkbox"/> Medical Record Number (MRN) | <input type="checkbox"/> Post code |
| <input type="checkbox"/> Date of Birth | | |
| <input type="checkbox"/> Other: | | |

TEST:

- | | |
|---------------------------------------|---|
| <input type="checkbox"/> Test type: | |
| <input type="checkbox"/> Test result: | |
| <input type="checkbox"/> Time of test | <input type="checkbox"/> Search test by a period of time: |
| <input type="checkbox"/> Date of test | From: To: |

Test priority code:

Routine
Urgent
Other:

- Do you require the results of any other tests ordered at the same/similar time:

DATA FILTERED BY:

- Ward / Hospital location – please specify:
- Requesting Doctor(s) – please specify:
- Other:

DATA REQUEST IS:

- One-off request
- Ongoing request (specify period): From: To:

COVERAGE OF DATA REQUEST IS:

- | | | |
|---|--|---|
| <input type="checkbox"/> Single hospital | <input type="checkbox"/> Multiple hospitals / LHDs | <input type="checkbox"/> Statewide NSW (All hospitals / all LHDs) |
| <input type="checkbox"/> FASS / Forensic Medicine | | |

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To assist you in completing the form please refer to [Human Research Ethics Committees – Quality Improvement & Ethical Review: A Practice Guide for NSW](#)

Section 1: ISSUES WHICH MAY REQUIRE CONSENT

1. The project involves direct contact with patients, consumers, or members of the public.	Yes	No
2. The project poses additional risks or burdens to the patient beyond their routine care.	Yes	No
3. The data to be collected is of a sensitive nature or application.	Yes	No
4. The purpose of the activity is not 'directly related' to the patient's disease, illness or its management.	Yes	No
5. The data will be used or available in such a way that may identify individuals.	Yes	No

Section 2: PRIVACY AND CONFIDENTIALITY

6. The final dataset will contain information that identifies the participants.	Yes	No
7. Is the proposed activity to be conducted by a person who does not normally have access to the client's health or other records for care or a directly related secondary purpose?	Yes	No
8. The project involves rare conditions or a small community.	Yes	No
9. Data will be selected or identified by: <ul style="list-style-type: none">• Aboriginal or Torres Strait Islander status; or• Ethnic, religious or minority group.	Yes	No
10. Data will be collected beyond that which is normally collected in routine care.	Yes	No

Section 3: OTHER IMPLICATIONS

11. The project uses 'new' interventions, protocols or equipment.	Yes	No
12. The project will involve allocation of patients to groups to enable comparisons.	Yes	No
13. The project will involve genetic tests/testing.	Yes	No
14. The project may potentially infringe the rights, privacy or professional reputation of carers, health professionals or institutions.	Yes	No
15. The project involves use of placebo.	Yes	No
16. The project is likely to generate data that may lead to publication.	Yes	No