

Participant Information

Non-Interventional Study - Adult providing own consent

Title	Map-HD Registry
Short Title	Map-HD Registry
Protocol Number	HDNA-Registry-0001.0
Project Sponsor	Monash University
Coordinating Investigator/ Principal Investigator	Professor Julie Stout email Julie.Stout@Monash.edu phone (03)9905-3987
Associate Investigator	Dr Yifat Glikmann-Johnston email Yifat.Glikmann-Johnston@Monash.edu phone (03)9902-0238
Location	Monash University
Registry Type	Self-Report Registry

1. Introduction

The Map-HD Registry collects information from people in Australia affected by Huntington's disease (HD). This includes people with HD and their family members and carers.

Please read this information to help you decide if you want to take part. Feel free to discuss it with others, such as your family or doctors. Make sure you understand it. You are welcome to contact the registry team to ask questions. Their email addresses and phone numbers are in the table above this section.

Participation in the Map-HD Registry is voluntary. There will be no effect on your care and services if you take part. Participation is free of cost.

To take part in the registry, you will need to consent. The consent form follows this information.

You can download a copy of this information from the Huntington's Disease Network of Australia (HDNA) website at hdna.com.au. You may log in to the registry at any time to view your consent and any information you provide.

2. What is the purpose of the Map-HD Registry?

To enable research that fosters the best quality of life for all people in Australia affected by HD.

The registry is the first-ever collection of data from people across Australia aimed at:

- Finding people that may benefit from new services and treatments
- Increasing the efficiency of HD research in Australia
- Connecting people to research and clinical trial opportunities
- Generating new knowledge about HD to help find treatments and improve care

3. What information do I need to provide if I join the Map-HD Registry?

To participate in the Map-HD Registry, you first need to provide consent. You then answer questions about yourself in the Map-HD Registry portal.

Some questions are **REQUIRED**. They take about 10 minutes to complete:

- Your first and last name
- Your date of birth
- Your post code
- How you prefer us to contact you in future (via text, email, phone call or post)
- Your sex and gender
- Whether you have had a blood test for the HD gene
- Whether you have been diagnosed with HD

Other questions are **OPTIONAL**. They take about 20 minutes to complete:

- Your CAG-expansion (or repeat number) if you have had the HD genetic test
- Which HD specialist clinicians or services you use, if any
- How far you travel to clinical appointments
- If you have accessed the National Disability Insurance Scheme (NDIS) or Aged Care services

4. What steps do I need to take to join the Map-HD Registry?

To join the registry:

Step 1: Create an account by entering a username and password. Your account is stored on a secure and private Map-HD Registry portal.

Use your account whenever you log in. You can register more than one person with your account. For example, using your account, you may register yourself and a person you care for.

Each person registered has their own consent. The registry separates each person's information. Each person's record is their profile.

Step 2: Read this Participant Information or ask someone to read it to you. Make sure you understand the information. If you have any questions, get them answered before continuing.

Step 3: Read the Consent Agreement.

Step 4: Where you see 'Declaration by Participant', fill in your name and the date. By filling in your name you are consenting to participate.

Step 5: The registry gives you questions to complete. You can ask someone you trust to assist you in completing the questions.

You can look at a summary of the responses by going to the summary page.

5. Can I stop before answering all the questions and complete some questions later?

Yes. You may log out and back in at any time.

When answering questions, each page saves when you click the 'Save and Continue' button. If you leave a page before clicking 'Save and Continue' that information will not save.

6. What will happen with the data I provide?

The Map-HD Registry is housed at Monash University, in Victoria, on secured computer systems.

Monash University will use information you provide for the purposes described in this document.

No records will be released with identifying information EXCEPT if you provide clear permission to do so.

Sharing of de-identified data from the Map-HD Registry

Before sharing data, personal information is removed. For example, your name and address are removed. All information that could identify a participant is removed. The data without personal information is called **de-identified data**. De-identification protects the identity of people included in the registry.

We plan to share data with researchers, clinicians, HD advocacy organisations, Government, and industry. To obtain data, a formal request must be made to the Map-HD Steering Committee. The purpose of sharing data is to help generate new understanding of HD and to help find treatments.

Findings will be published in journals. Researchers may describe the findings in talks in Australia and overseas. We will share findings in newsletters, social media and other media. We will describe the de-identified data.

7. What are the possible benefits of taking part in the Map-HD Registry?

- Finding out about opportunities for accessing clinical services, research, treatments, and HD events.
- Creating more knowledge about HD in Australia. More knowledge may speed up the approval of new services and treatments.
- Improved knowledge about HD may increase funding from Government or charities.
- Locations of HD services may be improved to better match where people live.

We cannot guarantee that you will benefit from participating in the Map-HD Registry.

8. What are the possible risks and disadvantages of taking part?

We do not expect risks or disadvantages from taking part. We have detailed policies and procedures to maintain the privacy and security of participant data. You can read the Terms and Conditions on the registry website. You can request additional details by contacting the registry team at info@hdna.com.au.

9. Can I withdraw from the Map-HD Registry?

Yes, you can withdraw from the registry at any time. To withdraw, delete your user profile. You may notify a member of the research team before you withdraw, or to assist you.

If you withdraw, we will not collect additional information from you. We save the information you provided so far. Reports include your saved information as de-identified data. Tell the researchers if you do not want your data included.

10. When will the registry close? Could it stop without warning?

There is no planned end date for the registry. The registry could be stopped if:

- No funding is available to maintain it, or
- it no longer holds value for the Australian HD community.

If the registry closes, we will archive all data. The data will be stored securely at Monash University. We plan to continue sharing de-identified data for approved uses.

You have the right to request access to your information collected and stored by the research team. You have the right to request that any incorrect information is corrected.

11. What should I do if I wish to make a complaint?

If you have complaints about the registry or if you have questions, contact:

Monash Health Human Research Ethics Committee
Executive Officer
Phone: 03 9594 4611

Email: research@monashhealth.org

If taking part causes you to feel upset or distressed, contact a member of the research team for help.

12. Who is organising and funding the Map-HD Registry?

The registry is funded by the Australian Government's National Health and Medical Research Council (NHMRC). The funds come from a grant to Professor Julie Stout at Monash University. Professor Stout's research team led the development of the registry. The team worked with a committee called the Map-HD Working Group of the HDNA. The working group included clinicians, staff from HD advocacy organisations, and people affected by HD.

We plan to support the registry using grant funding. The Australian Government, companies or charities may provide support for the registry.

13. Who has reviewed the ethics of the Map-HD Registry?

A Human Research Ethics Committee (HREC) reviews all registries in Australia. This committee is a group of people independent of the project. The HREC of Monash Health has approved the ethical aspects of the Map-HD Registry.

This registry complies with Australia's *National Statement on Ethical Conduct in Human Research (2018)*. This statement protects the interests of people who participate in human research studies and registries.

14. Further information and who to contact

For more information about the Map-HD Registry, contact Professor Julie Stout (Julie.Stout@monash.edu, 03 9905 3987) or Dr Yifat Glikmann-Johnston (Yifat.Glikmann-Johnston@monash.edu, 03 9902 0238).

Reviewing HREC approving this registry and HREC Executive Officer details

Reviewing HREC Name	Monash Health Human Research Ethics Committee
HREC Executive Officer	Executive Officer
Telephone	03 9594 4611
Email	research@monashhealth.org

Consent Form

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Consent Agreement for participation in the Map-HD Registry

I have read the Map-HD Registry Participant Information or someone has read it to me in a language that I understand. I have had any questions answered.

I consent to the personal information that I provide to the Map-HD Registry being used by Monash University for the purposes set out in the Participant Information.

I consent to the research team contacting me from time to time about updating my profile.

I consent to participate in the Map-HD Registry as described. I understand that I am free to withdraw at any time without it affecting my future health care.

Declaration by Participant – for participants who have read the information

Name of participant (type your name here to indicate your consent)

Date _____

Only complete the section below if the person providing informed consent was unable to read these documents. In such a case, a witness must sign below. Otherwise, no witness is needed.

Declaration for participants unable to read the information and consent form

Witness to the informed consent process

Name (type your name here to indicate you have witnessed the consent)

Date _____

*Witness is not to be the Investigator, a member of the study team or their delegate. Witness must be 18 years or older.