

My experiences, my rights: Health and wellbeing project



Information about our survey



What you will find in this booklet



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About this booklet



This is an Easy Read information booklet.



It is about some **research** that will be done about health services.



Research is about looking into something to understand it better.



This booklet will tell you about:

- who is doing the research
- what the research is looking into
- how the research will be done
- our survey.

Disabled people can do a **survey** to let us know what they think about health and wellbeing services.



A **survey** is a list of questions that you can answer.

The information in this booklet will tell you more about the survey.



You may find it helpful to get someone to read this booklet with you.

Who is doing the research?



The research will be done by 3 researchers from the Donald Beasley Institute.



They have all worked with disabled people for a long time.

They are:



Dr Brigit Mirfin-Veitch

Dr Robbie Francis Watene



Umi Asaka

Dr Francis Watene and Umi are both disabled people too.

What is the research about?



The **Disability Convention** is an international agreement that sets rules around services for disabled people.



The **Disability Convention** sets out what the New Zealand Government should do to make sure disabled people:



- have the same rights as everybody else
- are able to live a good life.



The research will look at health and wellbeing issues for disabled people in New Zealand.



We would like to hear about the things you think we should look at.



We could look at things like **health** policies and laws.



Health policies and laws are rules made by the Government which health services must follow.



We could also look at things like your **rights**.



Rights are things that you are entitled to.



For example everyone has a right to things like:

- housing
- being kept safe
- having enough food to eat.



Sometimes disabled people do not get their rights.



This research will help make the health system better for disabled people in New Zealand.

Who can do the survey?



Disabled people who are 18 years and older can fill in the survey.



We would like to hear stories about:

- things that you like about your health services
- things that you do not like about your health services



 how health services can be made better.



You can ask someone to help you complete this form.



This can be someone you trust like a:

- family member
- friend



- support person
- carer.



You can get in touch with us if you have any questions.



Our **contact details** are on **page 19** of this booklet.

Giving consent



In the survey we will ask for your consent.



Consent means that you say yes to us using what you write in our survey.



It is also ok if you do not want us to use your information.

Your rights



This research has been agreed by the Northern B Health and Disability Ethics Committee.



This means that a group of people have made sure that the research is safe.



You have the right to:

- 1. ask questions about the research
- 2. only answer questions you want to

What we will do with your information



We will use the information you give us to write reports.

We can give you a copy of the reports.



We will keep your information in a locked cabinet for 10 years.



Only the research team will see your information.

You can see who they are on page X of this booklet.



After 10 years this information will be destroyed.

Keeping your information safe



Confidentiality means keeping the things you let us know about private and safe.

To keep confidentiality:

Your name

 we will not use your name in any of the research reports



 your name will be changed so that no ones who said what.



The research report will be read by:

- disabled people
- people working in government departments
- people working in health services.



How to do the survey



There are 2 ways you can do the survey.

1. You can complete the online survey on our website:

<u>www.donaldbeasley.org.nz/projects/disabled-person-led-monitoring-of-the-uncrpd/</u>



You can download the survey questions from our website and fill out the answers.

After this, you can send us your answers by email or post by **Friday 24th of July 2020**.

More information



If you have questions about keeping safe in this research you can:

Phone: 0800 4 ETHICS

Email: hdecs@moh.govt.nz



For Māori health support:

Phone: 0800 878 839

Email:

admin@donaldbeasley.org.nz



If you want to talk to someone who is not part of this research you can talk the **Advocacy Service**.

The Advocacy Service can:



- listen to anything that you are concerned about
- give you ideas about what you can do



support you to be heard when raising concerns.

To contact the **Advocacy Service**



Phone: 0800 555 050

Fax: 0800 2787 7678



Email:

advocacy@advocacy.org.nz



Website:

https://www.advocacy.org.nz/

How to get in touch with us



If you want to find out more information you can:

Phone:



0800 878 839

This a free call number



Free call number means it will not cost you any money to call this number.



You can **email** us on

uasaka@donaldbeasley.org.nz



See our website:

www.donaldbeasley.org.nz/projects/disabled-person-led-monitoring-of-the-uncrpd/



This information has been translated into Easy Read by the Make It Easy service of People First New Zealand Inc. Ngā Tāngata Tuatahi.



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