


# Living Well with HD

we'd love you to join us for a conversation to help find better ways to support the huntington's disease community to live meaningfully and purposefully

*now and into the future.*



We are interested in learning about what living well with Huntington's Disease might mean for you and your family. Your voice is important and we want to hear directly from communities and people living with HD on how you perceive living meaningfully with a progressive disease, and to find ways researchers can work in partnership with the Huntington's Disease community to support wellbeing.





## Who are we looking for?

Age 18+ affected by HD (broadly defined - you do not need to have a diagnosis to participate);  
family members and care partners;  
clinicians;  
community advocates and anyone involved with the HD community



## What is involved?

30-45 min phone call to see if you are eligible and answer your questions.  
Followed by discussions of up to 2 hours on wellbeing and best ways to research this topic.  
In person or online, at a time that works for you  
You will be reimbursed with a \$60 gift voucher.



Please scan the QR code for more info or email [ji.loong@monash.edu](mailto:ji.loong@monash.edu)



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THIS PROJECT HAS BEEN APPROVED BY THE MONASH UNIVERSITY HUMAN RESEARCH ETHICS COMMITTEE MUHREC

**PROJECT ID 41021** LIVING WELL WITH HUNTINGTON'S DISEASE (FOCUS GROUPS/INTERVIEWS).

PLEASE CONTACT JI-SHEN LOONG (JI.LOONG@MONASH.EDU) FOR MORE INFORMATION

DISTINGUISHED PROFESSOR JULIE STOUT IS THE CHIEF INVESTIGATOR OF THIS PROJECT

# We need your help to optimise wellbeing in Huntington's Disease *now and into the future.*

We are interested in learning about what living well with Huntington's Disease means for you and your family.

You are invited to have a conversation with us to support the Huntington's Disease community live meaningfully and purposefully, and to find ways we can work together to support wellbeing.

## WHO ARE WE LOOKING FOR?

- AGE 18+ AFFECTED BY HD (BROADLY DEFINED - YOU DO NOT NEED TO HAVE A DIAGNOSIS TO PARTICIPATE);
- FAMILY MEMBERS AND CARE PARTNERS, CLINICIANS
- COMMUNITY ADVOCATES AND ANYONE INVOLVED WITH THE HD COMMUNITY

## WHAT IS INVOLVED?

- 30-45 MIN PHONE CALL TO SEE IF YOU ARE ELIGIBLE AND ANSWER ANY QUESTIONS YOU HAVE.
- FOLLOWED BY DISCUSSIONS OF UP TO 2 HOURS ON WELLBEING AND BEST WAYS TO RESEARCH THIS TOPIC.
- IN PERSON OR ONLINE, AT A TIME THAT WORKS FOR YOU

YOU WILL BE REIMBURSED WITH A \$60 GIFT VOUCHER.

TO LEARN MORE,  
PLEASE SCAN THE QR CODE  
FOR MORE INFORMATION



OR EMAIL

[JI.LOONG@MONASH.EDU](mailto:JI.LOONG@MONASH.EDU)



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University

This project has been approved by the Monash University Human Research Ethics Committee MUHREC Project ID 41021 Living Well with Huntington's Disease (Focus Groups/Interviews).

Questions can be directed to: Ji-Shen Loong ([Ji.Loong@Monash.edu](mailto:Ji.Loong@Monash.edu)).

Distinguished Professor Julie Stout is the Chief Investigator of this project