## **Consent Form**



## Please tick to indicate you consent to the following I have read, or have had read to me in my first language, and I understand the Participant Information Sheet. I have been given sufficient time to consider whether or not to participate in this study. I have had the opportunity to use a legal representative, whanau/ family support or a friend to help me ask questions and understand the study. I am satisfied with the answers I have been given regarding the study and I have a copy of this consent form and information sheet. I understand that taking part in this study is voluntary (my choice) and that I may withdraw from the study at any time without this affecting my medical care. I consent to the research staff collecting and processing my information, including information about my health. If I decide to withdraw from the study, I agree that the information collected about me up to the point when I withdraw may continue to Yes □ No □ be processed. I consent to my GP or current provider being informed about my No □ participation in the study and of any significant abnormal results Yes □ obtained during the study. I agree to an approved auditor appointed by the New Zealand Health and Disability Ethic Committees, or any relevant regulatory authority or their approved representative reviewing my relevant medical records for the sole purpose of checking the accuracy of the information recorded for the study. I understand that my participation in this study is confidential and that no material, which could identify me personally, will be used in any reports on this study. I understand that this form and a secured electronic file with my name and Participant Identification Number will be kept for 6 years, after which time they will be destroyed. I understand that de-identified study data may be kept indefinitely to allow for publication and future re-analysis, and with the permission

the purpose of undertaking related research.

Dated: 13th June 2022

of the lead investigator (A/Prof Fisher) shared with collaborators for

| I understand the compensation provisions in cathe study.  | se of injury during |       |      |
|---|---------------------|-------|------|
| I know who to contact if I have any questions a general.  | bout the study in   |       |      |
| I understand my responsibilities as a study par   | ticipant.           |       |      |
| I wish to receive a summary of the results from   | the study.          | Yes □ | No □ |
| Declaration by participant: I hereby consent to take part in this study.  |                     |       |      |
| Participant's name:   |                     |       |      |
| Signature:  | Date:               |       |      |
| Declaration by member of research team:   |                     |       |      |
| I have given a verbal explanation of the research project to the participant, and have answered the participant's questions about it. |                     |       |      |
| I believe that the participant understands the study and has given informed consent to participate.                                   |                     |       |      |
| Researcher's name:  |                     |       |      |
| Signature:  | Date:               |       |      |

This study has been reviewed and approved by the Health and Disability Ethics Committee [20CEN30] on 05/03/2020 for 5 years.

Lay study title:
PIS/CF version no.:

Brain health in atrial fibrillation 2 (Control participant version)

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