

1. What is the research study about?

You are invited to take part in this research study. The research study aims to determine whether family carers of people with traumatic brain injury find an on-line program about how to manage challenging behaviour useful. You have been invited because you have indicated that you have a family member with a TBI and you would be interested in being involved.

2. Who is conducting this research?

The study is being carried out by the following researchers: Professor Skye McDonald, Dr Emily Trimmer, Dr Jill Newby, School of Psychology, UNSW, Professor Graeme Simpson, Ingham Institute of Applied Medical Research, Ms Samantha Grant, Inspire Psychology and Mr Paul Gertler, Gertler Psychological Services.

Research Funder: This research is being funded by iCare, NSW.

3. Inclusion/Exclusion Criteria

Before you decide to participate in this research study, we need to ensure that it is ok for you to take part. The research study is looking recruit people who meet the following criteria:

- Adults aged 18 years or over,
- Identify as a family member of an individual who has experienced a traumatic brain injury
- Identify as caring for that person
- Have access to a computer and printer,
- Prepared to provide your name, phone number, address, and informed consent.
- Fluent English.

4. Do I have to take part in this research study?

Participation in this research study is voluntary. If you do not want to take part, you do not have to. If you decide to take part and later change your mind, you are free to withdraw from the study at any stage.

If you decide you want to take part in the research study, you will be asked to:

- Read the information carefully (ask questions if necessary);
- Sign and return the consent form if you decide to participate in the study;
- Take a copy of this form with you to keep.

5. What does participation in this research require, and are there any risks involved?

Completion of an On-line questionnaire

If you decide to take part in the research study, we will ask you to complete an on-line questionnaire. The questionnaire will ask you questions about your age, your relation to your family member with TBI, whether you are able to read English and ask for contact details. It should take approximately 5 minutes to complete. Information you provide will be linked via a unique participant identification number that is only accessible by the investigators.

We don't expect this questionnaire to cause any harm or discomfort, however if you experience feelings of distress as a result of participation in this study you can let the research team know and they will provide you with assistance.



Participation in the "Carer's Way Ahead" On-line program

If you meet our inclusion criteria on screening we will arrange a telephone interview with you to answer any questions and confirm that you meet inclusion criteria so as to invite you into the study.

Once you have agreed to participate, you will be asked to complete a range of self-report measures before the start of your online program (baseline), post-program (week 9), and again at 3-month follow-up. You will then be provided with access to the online program, which comprises 7 online modules over a 12-week period. You will be able to pick and choose the modules you wish to complete. You will be encouraged to complete one lesson per one to two weeks. You will receive regular phone and/or email contact from us and will be encouraged to carry out practical activities between lessons.

We don't expect the "Carer's Way Ahead" program to cause any harm or discomfort, however if you experience feelings of distress as a result of participation in this study you can let the research team know and they will provide you with assistance. Regular "check ins" will be made throughout the program and on completion of questionnaires to monitor your distress. If you are experiencing overwhelming distress you can discontinue the program at any time and be referred to alternative sources of support.

6. What are the possible benefits to participation?

We hope to use information we get from this research study to benefit others who are also caring for a family member with a TBI and who has challenging behaviour.

7. What will happen to information about me?

By signing the consent form you consent to the research team collecting and using information about you for the research study. Your data will be kept for 7 years after the project's completion. Data will be entered into spreadsheets and data analytic software using password protected files, on a locked computer with password protection. Information about your identification will be stored in a secure excel file with password protected file with the decoding information. Responses recorded on paper from the telephone interview will also be stored in locked cabinets, only accessible by the research team.

All online surveys and participation metrics (e.g., logins, details, time since login) will be collected via the Virtual Clinic website, and linked to your unique individual login code. Each time a questionnaire/survey is available to complete, you will login to the virtual clinic website with your email and a unique password you chose at application stage, and complete the questionnaire of interest. This ensures that all of your responses are linked.

Your information will only be used for the purposes of evaluating the "Carer's Way Ahead Program". Information we receive about you will be collated with that from other people and will only ever be disclosed in research publications in a non-identifiable format.

The information you provide is personal information for the purposes of the Privacy and Personal Information Protection Act 1998 (NSW). You have the right of access to personal information held about you by the University, the right to request correction and amendment of it, and the right to make a compliant about a breach of the Information Protection Principles as contained in the PPIP Act. Further information on how the University protects personal information is available in the <u>UNSW Privacy Management Plan</u>.

8. How and when will I find out what the results of the research study are?

The research team intend to publish and/ report the results of the research study in a variety of ways. All information published will be done in a way that will not identify you.



If you would like to receive a copy of the results you can let the research team know by adding your email address to the consent form. We will only use these details to send you the results of the research.

9. What if I want to withdraw from the research study?

If you do consent to participate, you may withdraw at any time. You can do so by completing the 'Withdrawal of Consent Form' which is provided at the end of this document. Alternatively you can ring the research team and tell them you no longer want to participate. Your decision not to participate or to withdraw from the study will not affect your relationship with UNSW Sydney or iCare lifetime care.

If you decide to leave the research study, the researchers will not collect additional information from you. Any identifiable information about you will be withdrawn from the research project. The research team will destroy any information about you that was collected during your participation in the study.

10. What should I do if I have further questions about my involvement in the research study?

The person you may need to contact will depend on the nature of your query. If you require further information regarding this study or if you have any problems which may be related to your involvement in the study, you can contact the following member/s of the research team:

Research Team Contact Details

Name	Emily Trimmer Project Manager • 02-93853310 e.trimmer@psy.unsw.edu.au	
Position		
Telephone		
Email		

Support Services Contact Details

If at any stage during the study you become distressed or require additional support from someone not involved in the research please call:

Name/Organisation	LifeLine
Telephone	13 11 14

What if I have a complaint or any concerns about the research study?

If you have a complaint regarding any aspect of the study or the way it is being conducted, please contact the UNSW Human Ethics Coordinator:

Complaints Contact

Position	UNSW Human Research Ethics Coordinator	
Telephone	+ 61 2 9385 6222	
Email	humanethics@unsw.edu.au	
HC Reference	HC180364	
Number		



Consent Form – Participant providing own consent

Declaration by the participant

I understand I am being asked to provide consent to participate in this research study;

I have read the Participant Information Sheet or someone has read it to me in a language that I understand;

I understand the purposes, study tasks and risks of the research described in the study;

I provide my consent for the information collected about me to be used for the purpose of this research study only.

I have had an opportunity to ask questions and I am satisfied with the answers I have received;

□ I freely agree to participate in this research study as described and understand that I am free to withdraw at any time during the study and withdrawal will not affect my relationship with any of the named organisations and/or research team members;

 \Box I would like to receive a copy of the study results via email or post, I have provided my details below and ask that they be used for this purpose only;

Name:			

Address: _____

Email Address: ____

I understand that I will be given a signed copy of this document to keep;

Participant Signature

Name of Participant (please print)		
Signature of Research Participant		
Date		

Declaration by Researcher*

□ I have given a verbal explanation of the research study, its study activities and risks and I believe that the participant has understood that explanation.

Researcher Signature*

Name of Researcher (please print)	
Signature of Researcher	
Date	

⁺An appropriately qualified member of the research team must provide the explanation of, and information concerning the research study.

Note: All parties signing the consent section must date their own signature.



Form for Withdrawal of Participation

I wish to **WITHDRAW** my consent to participate in this research study described above and understand that such withdrawal **WILL NOT** affect my relationship with The University of New South Wales, *ICare*. In withdrawing my consent I would like any information which I have provided for the purpose of this research study withdrawn.

Participant Signature

······································	
Name of Participant	
(please print)	
Signature of Research	
Participant	
Date	

The section for Withdrawal of Participation should be forwarded to:

CI Name:	Professor Skye McDonald
Email:	s.mcdonald@unsw.edu.au
Phone:	02-93853029
Postal Address:	School of Psychology, UNSW, Sydney, 2052