

# Effects of an e-health intervention 'iSupport' for reducing distress of dementia carers

<b>Submission date</b> 27/11/2020	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 09/03/2021	<b>Overall study status</b> Completed	<input checked="" type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 19/05/2025	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

NHS guidelines recommend informal carers of people living with dementia such as family and friends should be offered training to help them develop care skills and manage their own physical and mental health. The World Health Organisation recommend access to affordable, proven, well-designed, online technologies for education, skills training and support for dementia carers. In response to these recommendations, this is the first study in the UK of the benefits of an online training and support programme. This was developed by the World Health Organisation and is called 'iSupport'. It is designed for dementia carers to help them provide good care and take care of themselves. Carers can use iSupport at their own pace. They can access whichever parts of iSupport they feel are most relevant to them from their place of choice using a computer, tablet or smartphone. It is important to do this work because most people living with dementia are cared for at home, supported by a family member or friend who has limited knowledge of the condition. The role is very stressful, and many carers experience more mental and physical illness compared to people who are not dementia carers. COVID-19 has meant that many older people have to self-isolate, placing increasing pressures on carers. The aim of this study is to examine the effectiveness of iSupport in reducing distress.

### Who can participate?

Dementia carers aged 18+ years in England, Wales and Scotland, and 30 young carers

### What does the study involve?

Half of the carers will complete iSupport and half will receive information leaflets. All the carers will answer questionnaires about stress and strain, mental health, resilience and knowledge of dementia. They will do this three times over a 6-month period. By comparing the two groups, the researchers will be able to show whether iSupport is better than simply giving information. To understand what aspects of iSupport carers like the most, and what might put them off using it, the researchers will undertake interviews with a smaller group (50) of the carers and have a longer discussion about their thoughts, feelings and the impact of iSupport on themselves. They will do this three times over a 6-month period. To examine the potential costs and benefits of iSupport carers will answer questions about health and social care service use and their quality of life. They will do this three times over a 6-month period. There are currently no evidence-based support interventions for young carers of people with dementia. The researchers will

work with a group of young carers and adapt iSupport for their use, identify what outcomes are most important to them, and assess this new version of iSupport with 30 young carers.

Where is the study run from?

The study is run from Bangor University in Wales, with partners in the University of Strathclyde, Scotland and UCL in England

When is it starting and how long will it run for?

October 2020 to June 2024

Who is funding the study?

National Institute for Health Research - Public Health Research Programme (UK)

Who is the main contact?

Prof. Gill Windle

g.windle@bangor.ac.uk

## Contact information

**Type(s)**

Scientific

**Contact name**

Prof Gillian Windle

**ORCID ID**

<https://orcid.org/0000-0003-0479-1172>

**Contact details**

Dementia Services Development Centre Wales

Bangor University

Ardudwy

George Site

Holyhead Road

Bangor

United Kingdom

LL57 2EF

+44 (0)1248 38 3968

g.windle@bangor.ac.uk

**Type(s)**

Public

**Contact name**

Mr Gregory Flynn

**Contact details**

Meirion

Normal Site

Holyhead Road

Bangor University

Bangor  
United Kingdom  
LL57 2PZ  
+44 (0)1248 38 2195  
g.flynn@bangor.ac.uk

**Type(s)**  
Scientific

**Contact name**  
Prof Joshua Stott

**ORCID ID**  
<https://orcid.org/0000-0003-1361-053X>

**Contact details**  
1-19 Torrington Place  
Camden  
London  
United Kingdom  
WC1E 7HB  
+44 (0)20 7679 2000  
j.stott@ucl.ac.uk

**Type(s)**  
Scientific

**Contact name**  
Dr Kieren Egan

**ORCID ID**  
<https://orcid.org/0000-0002-1639-4281>

**Contact details**  
Computer & Information Sciences  
Livingstone Tower  
26 Richmond Street  
Glasgow  
United Kingdom  
G1 1XH  
+44 (0)141 552 4400  
kieren.egan@strath.ac.uk

## **Additional identifiers**

**Clinical Trials Information System (CTIS)**  
Nil known

**Integrated Research Application System (IRAS)**  
311565

**ClinicalTrials.gov (NCT)**

Nil known

**Protocol serial number**

PHR - NIHR130914, IRAS 311565

## Study information

### Scientific Title

A randomized controlled trial and feasibility study of the effects of an e-health intervention 'iSupport' for reducing distress of dementia carers

### Acronym

iSupport

### Study objectives

1. Is carer distress and/or symptoms of depression (primary outcomes) significantly reduced in participants allocated to receive iSupport compared to participants allocated to a comparison group?
2. Are symptoms of anxiety (secondary outcome) significantly reduced, and resilience, relationship quality and dementia knowledge (secondary outcomes) significantly increased in participants allocated to receive iSupport compared to participants allocated to a comparison group receiving standardised information about dementia?
3. What are the participant and contextual barriers and facilitators to the implementation of iSupport?
4. What potential mechanisms might underpin changes in outcomes from using iSupport?
5. What is the cost-effectiveness of iSupport compared to standardised information about dementia?
6. What are the carers' perspectives of iSupport in relation to supporting them in an ongoing or future repeated pandemic such as COVID-19?

The feasibility study will assess if it is feasible, useful and acceptable to refine and digitally deliver a refined iSupport to young carers.

### Ethics approval required

Ethics approval required

### Ethics approval(s)

1. approved 12/04/2021, Bangor University School of Medical and Health Sciences Academic Ethics Committee (AEC) (Bangor University, College Road, Bangor, LL57 2DG, United Kingdom; +44 1248 351 151; hcmsethics@bangor.ac.uk), ref: 2021-16915
2. approved 02/11/2022, London - City & East Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 2071048171; cityandeast.rec@hra.nhs.uk), ref: 22/LO/0688

### Study design

**Trial:** multi-centre pragmatic single-blinded two-arm randomized controlled trial with an internal pilot

**Feasibility study:** non-randomized feasibility study of intervention refinement for younger dementia carers

## **Primary study design**

Interventional

## **Study type(s)**

Quality of life

## **Health condition(s) or problem(s) studied**

Reductions in psychological distress and improvements in skills and resilience in dementia carers

## **Interventions**

WS1 is a multi-centre, pragmatic, single-blinded, two-arm randomized controlled trial (RCT) with an internal pilot. Randomization through a secure online system employing a dynamic randomisation algorithm will allocate patients to the intervention or control-comparison group. Participants assigned to the control-comparison group will receive information about dementia developed by the Alzheimer's Society. A 6-month internal pilot will be nested in this WS at each site. Stop/Review/Go criteria will indicate the success of the participant recruitment targets (e.g. recruitment and set up/ training of sites within time allocated: Go: 3, Review: 2, Stop: 1. Recruitment of participants based on target of n=110 by month 6 of recruitment: Go: >=94 (85%), Review: 56-93 (50 - 84%), Stop: <=55 (<50%)). The study is guided by the 'Context and Implementation of Complex Interventions' checklist.

WS2 PROCESS EVALUATION will run alongside WS1 and apply mixed-methods (semi-structured interviews, quantitative questionnaires and analysis of data from the online platform). It will be conducted in line with established guidance frameworks. It will examine throughout the intervention period how participants engage with and adhere to particular aspects of iSupport (e.g. most/least frequently visited pages, the most 'popular' modules/sessions, sessions with quizzes with the highest rates of wrong responses). A System Usability Scale will be administered at 6-month follow up. This 10-item scale will quantitatively evaluate the overall usability of the iSupport platform. Semi-structured interviews will be undertaken using an internet-based service (e.g. Zoom, Teams, Skype, or GoToMeeting) or telephone, with a sub-sample of the intervention participants. These will be recorded and professionally transcribed. The topic guides will be guided by the process evaluation parameters described in recognised frameworks.

WS3 HEALTH ECONOMICS will evaluate the cost-effectiveness of iSupport from a public sector perspective. The cost of implementing iSupport (e.g., time spent supporting carers to use the tool, and the cost of maintaining technical support for the tool) will be calculated.

WS4 A NON-RANDOMIZED FEASIBILITY STUDY will refine iSupport through co-design workshops with young carers. It will identify what outcomes are most important to young carers in relation to iSupport, and then explore the refined programme. Success will be defined as acceptability of the: a) recruitment process, b) assessments, c) data collection tools, d) intervention content & delivery to participants, as well as compliance. This work will be undertaken in line with WHO's iSupport version 1.0. Adaptation and implementation guide for international uptake.

## **Intervention Type**

Behavioural

### **Primary outcome(s)**

There will be two primary outcome measures for WS1. Sample size has been based on considering both outcomes as primary outcomes where a successful trial would be noted if at least one of the outcomes indicated a statistically significant effect. Both primary outcome measures will be administered at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2):

1. Carers' distress measured by the 12-item Zarit Burden Interview (ZBI-12)
2. Caregiver depression measured by the 10-item Centre for Epidemiological Studies of Depression Scale (CES-D 10)

### **Key secondary outcome(s)**

Current secondary outcome measures as of 04/03/2024:

1. The frequency of common symptoms of anxiety assessed through the Generalised Anxiety Disorder Questionnaire (GAD-7)
2. Improvements in the way carers perceive they can manage the situation assessed through the Resilience Scale-14 (RS-14)
3. The influence of iSupport on the quality of the caregiving relationship assessed through the 14-item Quality of the Carer-Patient Relationship (QCPR)
4. Improvements in how the carer understands their relative assessed through the 25-item Dementia Knowledge Assessment Scale (DKAS)
5. Utility values for Quality-Adjusted Life Years obtained from responses to the EQ-5D-5L
6. Impact on the health-related quality of life of the person being cared for assessed through the DEMQOL-Proxy
7. Out of pocket expenses obtained through a Service Use Questionnaire developed for iSupport
8. Opportunity costs assessed through the iMTA Valuation of Informal Care Questionnaire

The feasibility study (WS4) will adapt the standardised measures above based on focus group responses, and conduct the assessments with younger dementia carers at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2).

One measure, the Short Sense of Competence Questionnaire (SSCQ), was agreed with NIHR to remove from the case report form (CRF) due to overlap with other measures..

Measured at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2):

1. The frequency of common symptoms of anxiety assessed through the Generalised Anxiety Disorder Questionnaire (GAD-7)
2. Improvements in the way carers perceive they can manage the situation assessed through the Resilience Scale-14 (RS-14)
3. The influence of iSupport on the quality of the caregiving relationship assessed through the 14-item Quality of the Carer-Patient Relationship (QCPR)
4. Improvements in how the carer understands their relative assessed through the 25-item Dementia Knowledge Assessment Scale (DKAS)
5. Utility values for Quality-Adjusted Life Years obtained from responses to the EQ-5D-5L
6. Impact on the health-related quality of life of the person being cared for assessed through the DEMQOL-Proxy
7. Out of pocket expenses obtained through a Service Use Questionnaire developed for iSupport
8. Opportunity costs assessed through the iMTA Valuation of Informal Care Questionnaire

The feasibility study (WS4) will adapt the standardised measures above based on focus group responses, and conduct the assessments with younger dementia carers at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2).

One measure, the Short Sense of Competence Questionnaire (SSCQ), was agreed with NIHR to remove from the case report form (CRF) due to overlap with other measures.

---

Previous secondary outcome measures:

Measured at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2):

1. The frequency of common symptoms of anxiety assessed through the Generalised Anxiety Disorder Questionnaire (GAD-7)
2. Improvements in the way carers perceive they can manage the situation assessed through the Resilience Scale-14 (RS-14)
3. Sense of competence developed for dementia caregivers assessed through the 7-item Short Sense of Competence Questionnaire (SSCQ)
4. The influence of iSupport on the quality of the caregiving relationship assessed through the 14-item Quality of the Carer-Patient Relationship (QCPR)
5. Improvements in how the carer understands their relative assessed through the 25-item Dementia Knowledge Assessment Scale (DKAS)
6. Utility values for Quality-Adjusted Life Years obtained from responses to the EQ-5D-5L
7. Impact on the health-related quality of life of the person being cared for assessed through the DEMQOL-Proxy
8. Out of pocket expenses obtained through a Service Use Questionnaire developed for iSupport
9. Opportunity costs assessed through the iMTA Valuation of Informal Care Questionnaire

The feasibility study (WS4) will adapt the standardised measures above based on focus group responses, and conduct the assessments with younger dementia carers at baseline (T0), 3 months after baseline (T1) and 6 months after baseline (T2).

## **Completion date**

30/06/2024

# **Eligibility**

## **Key inclusion criteria**

WS1 & WS3

Dementia carers (age 18+ years):

1. Self-identify as an unpaid carer (partners, children, friends, etc.) of a person with dementia who is not living in a full-time care facility, caring at least weekly for at least 6 months
2. Self-identify as experiencing at least some stress, depression or anxiety
3. The care recipient has to have a confirmed diagnosis of dementia (through self-report by the carer, to reflect the 'real world' application of iSupport)

WS2 PROCESS EVALUATION will run alongside WS1 to undertake a mixed-methods exploration in line with Medical Research Council guidance. It will assess the implementation of iSupport, the participant's perspectives of the intervention, including the barriers, facilitators and perceived benefits, and theorize mechanisms of change

WS4

Young dementia carers (age 11-17 years):

1. Young people between the age of 11-17 years (secondary school age) who self-identify as a carer of a person with dementia who is not living in a full-time care facility, caring at least weekly for at least 6 months
2. The care recipient has to have a confirmed diagnosis of dementia (through self-report by the carer, to reflect the 'real world' application of iSupport)

**Participant type(s)**

Carer

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

11 years

**Sex**

All

**Total final enrolment**

352

**Key exclusion criteria**

Trial exclusion criteria:

1. Receiving psychological treatment from a mental health specialist at the time of recruitment
2. Unable to comprehend written English
3. No access to the internet
4. Unable to give informed consent to the trial
5. Have previously used iSupport materials (in the last 12 months)

Feasibility study (WS4) exclusion criteria:

1. Receiving treatment from Child and Adolescent Mental Health Services (CAMHS) at the time of recruitment
2. Unable to comprehend written English
3. No access to the internet
4. Have previously used iSupport materials (in the last 12 months)

**Date of first enrolment**

01/12/2021

**Date of final enrolment**

31/03/2023

**Locations**

**Countries of recruitment**



United Kingdom

England

Scotland

Wales

**Study participating centre**

**North Wales Organisation for Randomised Trials in Health (NWORTH)**

Y Wern

Normal Site

Holyhead Road

Bangor University

Bangor

United Kingdom

LL57 2PZ

## **Sponsor information**

**Organisation**

Bangor University

**ROR**

<https://ror.org/006jb1a24>

## **Funder(s)**

**Funder type**

Government

**Funder Name**

National Institute for Health Research

**Alternative Name(s)**

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

Location  
United Kingdom

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Prof. Gill Windle (g.windle@bangor.ac.uk).

### IPD sharing plan summary

Available on request

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>		20/11/2024	28/11/2024	Yes	No
<a href="#">Protocol article</a>		21/09/2022	22/09/2022	Yes	No
<a href="#">Basic results</a>	version 1	27/09/2024	27/09/2024	No	No
<a href="#">Other publications</a>	Cost-consequence analysis	16/05/2025	19/05/2025	Yes	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes
<a href="#">Statistical Analysis Plan</a>	version 1	29/09/2023	04/03/2024	No	No
<a href="#">Study website</a>	Study website	11/11/2025	11/11/2025	No	Yes