An evaluation of the ADAPT Assistive Technology (AT) Training Programme for health and social care professionals and students

Submission date	Recruitment status No longer recruiting	Prospectively registered		
10/08/2021		[X] Protocol		
Registration date	Overall study status	Statistical analysis plan		
01/09/2021	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
12/12/2023	Other			

Plain English summary of protocol

Background and study aims

Long-term disabilities often result in a loss of autonomy and a breakdown of social interaction, and therefore there is a demand for devices that ensure individuals feel enabled to live independently for as long as possible. Assistive Technology (AT) is therefore an increasingly important aspect of many fields of health and care practice. Several studies highlight challenges and barriers regarding the adoption and usage of AT, and evidence suggests that adequate support, training and education are vital, however, substantial barriers to education exist. The study aims to evaluate the ADAPT AT training programme developed as part of the Interreg funded ADAPT project

Who can participate?

Health and social care professionals and students who have undertaken the ADAPT AT Training programme (i.e. those who use AT, will do so in the future or have an interest in its practical application)

What does the study involve?

Completing a questionnaire following completion of each training unit within the programme and a further questionnaire 6 months later

What are the possible benefits and risks of participating?

Benefits of the training programme include enhanced knowledge and understanding in the area of AT and related issues for studies/careers and evidence of completion of training for CPD (Continuing Professional Development). Benefits of participating in the evaluation study include enhanced knowledge of academic research, contribution to a research study that will benefit the healthcare professional population as a whole, as well as enhanced future AT training where there is currently a gap identified in terms of provision. There are no foreseen risks in this element of the ADAPT Project

Where is the study run from? Canterbury Christ Church University (UK) When is the study starting and how long is it expected to run for? November 2016 to December 2022

Who is funding the study? Interreg VA France (Channel) England (FCE) 2014-202 Programme

Who is the main contact? Prof Eleni Hatzidimitriadou, eleni.hatzidimitriadou@canterbury.ac.uk

Contact information

Type(s)

Scientific

Contact name

Prof Eleni Hatzidimitriadou

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Contact details

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

262502

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

CPMS 49337, IRAS 262502

Study information

Scientific Title

Evaluation Study of the online ADAPT Assistive Technology (AT) Training Programme

Acronym

ADAPT

Study objectives

The evaluation aims to assess learner knowledge, skills, attitudes, confidence and commitment, as well capturing as any behaviour changes or results as a consequence of undertaking the ADAPT AT training

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 01/10/2019, NHS Health Research Authority (Health Research Authority, Skipton House, 80 London Road, London, SE1 6LH, UK; +44 (0)20 7972 2545; hra.approval@nhs.net), ref: 19/HRA/4890

Study design

Interventional non randomized trial

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Internet/virtual

Study type(s)

Other

Participant information sheet

See additional files

Health condition(s) or problem(s) studied

Health Services Research

Interventions

An online questionnaire hosted via Online Surveys has been developed by the CCCU research team.

Participants will be invited to complete the questionnaire at the end of each training unit they undertake. A link will take them to BOS where they can read the Participant Information Sheet, provide their agreement/consent to participate and answer the questions.

The questionnaire is self-completed and includes basic background data (profession, years of practice, level of AT use) and a series of questions around learners' experiences and views of the training materials, as well as feedback on content and use of the online training platform. The questionnaire is anonymous and will take approximately 15 minutes to complete.

Intervention Type

Behavioural

Primary outcome measure

- 1. Learner knowledge is measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months
- 2. Learner skills are measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months
- 3. Learner attitudes are measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months
- 4. Learner confidence is measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months
- 5. Learner commitment is measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months
- 6. Learner behaviour changes are measured using a bespoke questionnaire based on Kirkpatrick's (2016) four levels of training evaluation post training and at 6 months

Secondary outcome measures

There are no secondary outcome measures

Overall study start date

29/11/2016

Completion date

31/12/2022

Eligibility

Key inclusion criteria

Health and social care professionals and students who have undertaken the ADAPT Assistive Technology (AT) Training

Participant type(s)

Health professional

Age group

Adult

Sex

Both

Target number of participants

Planned Sample Size: 277; UK Sample Size: 277

Total final enrolment

544

Key exclusion criteria

Individuals who have not undertaken the ADAPT Assistive Technology (AT) Training

Date of first enrolment

01/10/2019

Date of final enrolment

31/12/2022

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Canterbury Christ Church University

Faculty of Medicine, Health and Social Care North Holmes Road Canterbury United Kingdom CT1 1QU

Study participating centre Kent and Canterbury Hospital

East Kent Hospitals University NHS Foundation Trust Ethelbert Road Canterbury United Kingdom CT1 3NG

Study participating centre Royal Cornwall Hospital

Cornwall Mobility Penventinnie Lane Truro United Kingdom TR1 3LJ

Study participating centre University of Kent

Giles Lane Canterbury United Kingdom CT2 7NZ

Study participating centre University College London

Gower Street London United Kingdom WC1E 6BT

Sponsor information

Organisation

Canterbury Christ Church University

Sponsor details

North Holmes Road Canterbury England United Kingdom CT1 1QU +44 (0)1227922822 debra.towse@canterbury.ac.uk

Sponsor type

University/education

Website

http://www.canterbury.ac.uk/

ROR

https://ror.org/0489ggv38

Funder(s)

Funder type

Government

Funder Name

European Commission

Alternative Name(s)

European Union, Comisión Europea, Europäische Kommission, EU-Kommissionen, Euroopa Komisjoni, Ευρωπαϊκής Επιτροπής, Европейската комисия, Evropské komise, Commission européenne, Choimisiúin Eorpaigh, Europskoj komisiji, Commissione europea, La Commissione

europea, Eiropas Komisiju, Europos Komisijos, Európai Bizottságról, Europese Commissie, Komisja Europejska, Comissão Europeia, Comisia Europeană, Európskej komisii, Evropski komisiji, Euroopan komission, Europeiska kommissionen, EC, EU

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal.

Intention to publish date

31/12/2023

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be stored in a non-publically available repository.

IPD sharing plan summary

Stored in non-publicly available repository

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	version 3	05/11/2019	31/08/2021	No	Yes
<u>Protocol file</u>	version 2	07/11/2019	31/08/2021	No	No
Funder report results		30/11/2023	12/12/2023	No	No