A randomised controlled trial to measure the impact of behavioural SMS reminders messages on missed appointments among IAPT patients

Submission date	Recruitment status	[X] Prospectively registered
06/02/2018	No longer recruiting	∐ Protocol
Registration date	Overall study status	Statistical analysis plan
15/02/2018	Completed	Results
Last Edited	Condition category	Individual participant data
04/04/2023	Mental and Behavioural Disorders	Record updated in last year

Plain English summary of protocol

Background and study aims

The NHS Improving Access to Psychological Therapies (IAPT) services currently experience a relatively high rate of missed appointments (roughly 11%). This imposes a cost on the services and increases waiting times for others. The aim of this study is to test whether changes to the content of SMS (text message) reminders sent to IAPT patients before their appointments can reduce the rate of missed appointments. More specifically two different messages are tested against the existing business as usual message; one based on implementation intentions and the other on reciprocity. In addition to this, the study tests whether the behavioural messages can increase completion of treatment and ultimately improve mental health outcomes.

Who can participate?

Patients of one of the three participating IAPT services who have an appointment during the study period

What does the study involve?

Participants are randomly allocated to receive either the business as usual SMS reminder message or one of the two new messages. One of the new reminders asks the patient to take a moment to ensure that they are able to make their appointment. These messages are based on the psychological insight that asking people to consider the when, where and how of their behaviour can help translate intentions into action. The other new reminder lets the patient know that an appointment has been booked for them by their therapist. These messages are based on the concept of reciprocity whereby when someone does something on our behalf we reciprocate by going along with their request. All SMS messages are sent to patients 1-2 days before their appointment. The message also contains a prompt to rearrange the appointment if necessary. The messages purposely contain no message content that is clinical in nature. Appointment attendance and completion of treatment are recorded by the IAPT services at the end of the study period.

What are the possible benefits and risks of participating?

The new messages aim to increase the likelihood that patients attend their appointments.

Attending an appointment should increase the likelihood that the patient recovers and also help them recover sooner. Given that this study is being conducted with participants referred to or undergoing mental health treatment the main risk is that the new messages backfire and increase missed appointments or dropout. The messages have been carefully worded to guard against this happening.

Where is the study run from?

The study will be run at three participating NHS IAPT providers:

- 1. West London Mental Health Trust (UK)
- 2. Camden and Islington NHS Foundation Trust (UK)
- 3. Norfolk and Suffolk Foundation Trust (UK)

When is study starting and how long is it expected to run for? January 2017 to September 2018

Who is the main contact? Edward Flahavan Edward.Flahavan@bi.team

Contact information

Type(s)

Public

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 2017175

Study information

Scientific Title

A randomised controlled trial to measure the impact of behavioural SMS reminders messages on missed appointments among IAPT patients

Study objectives

The hypothesis is that changes to the wording of SMS appointment reminders informed by behavioural science can reduce the missed appointment rate among IAPT patients.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Other

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied

Mild to moderate anxiety or depression

Interventions

The intervention in this trial is a change to the wording of the SMS appointment reminders sent to patients 1-2 days prior to their IAPT appointment. Method of randomisation is a random allocation mechanism with SQL. This will be carried out by our partner on the trial Mayden (the company who send the SMS messages). Patients of participating IAPT services will be allocated 1: 1:1 to receive either the business as usual SMS reminder messages or one of the two behavioural SMS treatment arms. Once allocated to a trial arm, the patient will receive messages from that arm for the duration of the trial period.

- 1. Control this is the business as usual SMS reminder message.
- 2. Implementation plans these reminders ask the patient to take a moment to ensure that they will be able to make their appointment. These messages are based on the psychological insight that asking people to consider the when, where and how of their behaviour can help translate intentions into action. The concept was developed by psychologist Paul Gollwitzer.
- 3. Reciprocity these reminders let the patient know that an appointment has been booked for them by their therapist. These messages are based on the concept of reciprocity whereby when

someone does something on our behalf we reciprocate by going along with their request. The Behavioural Insights Team found this to be effective in encouraging job seekers to attend careers fairs.

These SMS messages will be sent to patients 1-2 days before their appointment. The message will also contain a prompt to rearrange the appointment if necessary. The messages purposely contain no message content that is clinical in nature.

The trial will run for 4 months, data collected during this period will be collected for analysis. After that point no new data will be collected for the trial.

Intervention Type

Behavioural

Primary outcome measure

Recorded by the IAPT services at the end of the trial period:

- 1. Appointment attendance whether or not a patient misses their appointment
- 2. Completion of treatment whether or not a patient completes their treatment

Secondary outcome measures

Reliable recovery, assessed using the PHQ-9 and/or the GAD-7 at initial assessment and at the end of treatment (or last available appointment where an assessment was taken)

Overall study start date

01/01/2017

Completion date

01/09/2018

Eligibility

Key inclusion criteria

- 1. Patients of all conditions, age and gender will be eligible
- 2. Patients has an appointment with one of the participating IAPT services during the trial period
- 3. Patient has consented to receive SMS communication from the service
- 4. Patient has supplied a valid mobile phone number to the service

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

It is expected that roughly 10,000 participants will be recruited into the trial during the period

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment 01/03/2018

Date of final enrolment 01/07/2018

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Camden and Islington NHS Foundation Trust

Room 208 3rd Floor South Wing St Pancras Hospital 4 St Pancras Way London United Kingdom NW1 0PE

Study participating centre Norfolk and Suffolk NHS Foundation Trust

Drayton High Road Norwich United Kingdom NR6 5BE

Study participating centre West London Mental Health Trust (Ealing)

3rd Floor 84 Uxbridge Road West Ealing London United Kingdom W13 8RA

Study participating centre West London Mental Health Trust (Hammersmith and Fulham)

1 Hammersmith Broadway

Sponsor information

Organisation

The Behavioural Insights Team

Sponsor details

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Sponsor type

Other

Website

https://www.bi.team/

ROR

https://ror.org/03mk5b468

Organisation

Mayden House Ltd

Sponsor details

1 Widcombe Crescent Bath England United Kingdom BA2 6AH

Sponsor type

Industry

Website

mayden.co.uk

Funder(s)

Funder type

Industry

Funder Name

Behavioural Insights Team

Funder Name

Mayden House Limited

Results and Publications

Publication and dissemination plan

The protocol is not currently available online and there are no immediate plans to publish this. Results will be published on the company website and in policy publications. The trialists also plan to publish the results in a high-impact peer-reviewed journal.

Intention to publish date

01/09/2019

Individual participant data (IPD) sharing plan

The participant level data is being shared with BIT for the trial in accordance with a data sharing agreement agreed with the participating IAPT providers. In this it is agreed that the raw data will not be made public. Though the data is anonymous the services are understandably very conscious of data protection issues.

IPD sharing plan summary

Not expected to be made available