

Health on the Web study: a workplace study of online feedback for a healthy lifestyle

Submission date 23/07/2012	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 26/07/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 09/07/2015	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

This study aims to determine whether it is feasible to deliver online screening and brief intervention via the workplace, and whether doing so reduces alcohol consumption. The study will determine what proportion of people offered screening accept it, what proportion who accept are drinking above recommended guidelines, what proportion make use of an online intervention to help them drink less and how effective the intervention is amongst those who use it.

Who can participate?

The HOW study aims to recruit 1,840 employees aged 18 and above from a large international organisation, based in the UK.

What does the study involve?

Employees who wish to take part will be asked to complete an online health questionnaire which will take about 10 minutes. The questionnaire includes questions on height, weight, smoking, alcohol intake, diet and physical activity. Participants will receive instantaneous online feedback based on their answers, with information about what they are already doing that's good for their health and what they might want to change. There will also be suggestions on where to get help if needed. Some participants will be contacted again in 3 months' time to complete further questionnaires.

What are the possible benefits and risks of participating?

Participants will receive a personalised health check, together with advice about how to improve their health. This is completely free, and completely confidential. Participants may get a shock if the results of the health check suggest they are at risk of a health problem. If they are worried about their health and feel they need more advice than we can give online, they are advised to go and see their family doctor. The doctor is the person best placed to help with health problems.

Where is the study run from?

The HOW study is based entirely online. It is being conducted by a team of researchers based in the e-Health Unit, at UCL.

When is the study starting and how long is it expected to run for?
Study recruitment is due to begin on 01 August 2012. There is a three week recruitment period, three month trial duration, and three week follow-up period. The study is due to close on the 5th December 2012.

Who is funding the study?
This study has been funded by the NIHR School for Primary Care Research.

Who is the main contact?
Dr Elizabeth Murray
elizabeth.murray@ucl.ac.uk

Contact information

Type(s)
Scientific

Contact name
Dr Elizabeth Murray

Contact details
University College London (UCL)
Research Department of Primary Care & Population Health
Royal Free Hospital
Rowland Hill Street
London
United Kingdom
NW3 2PF
+44 (0)20 7794 0500 ext. 36747
elizabeth.murray@ucl.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title
A randomised controlled trial of work-based online screening and brief intervention for hazardous and harmful drinking

Acronym
HoW

Study objectives

Feedback on alcohol consumption and access to an online intervention will lead to a reduction in drinking, compared with no feedback on drinking in the context of a health check on a range of behaviours.

The null hypothesis is that there will be no difference between groups in their alcohol intake.

Ethics approval required

Old ethics approval format

Ethics approval(s)

UCL Research Ethics Committee, 11/05/2012, ref: 3770/001

Study design

Online randomised controlled trial with wait-list control group

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Quality of life

Participant information sheet

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

Health condition(s) or problem(s) studied

Reducing alcohol misuse

Interventions

Participants randomised to the intervention group will receive individually tailored feedback on body mass index (BMI), smoking status, diet and physical activity. In addition, they will be informed that the information they provided suggests that they are at increased risk of alcohol-related harm, and advised to reduce the amount they drink. The advice will acknowledge that it can be hard to cut down, and state that further support in achieving this is available from the online intervention, Down Your Drink (DYD), with an embedded hyperlink which participants can click on to be taken directly to DYD. Participants can access the DYD website as often as they wish during the 3 month trial duration.

Intervention Type

Behavioural

Primary outcome measure

Past week alcohol consumption, measured by the TOT-AL. This is an online measure of past week alcohol consumption, validated in a UK population.

Secondary outcome measures

1. The number (proportion) of employees completing the online health check
2. The number (proportion) of employees who complete the online health check and score 5 or more on the AUDIT-C
3. The number (proportion) of participants in the intervention group who access DYD at least once
4. Alcohol use disorders, measured by the AUDIT. This will allow comparison of our data with other alcohol trials
5. Health state, measured by the EQ-5D
6. Number of days of sickness absence in past 3 months (self-reported)
7. Number and duration of hospital admissions in past 3 months (self-reported)
8. Costs, staff time and resources required to implement intervention

Overall study start date

01/08/2012

Completion date

05/12/2012

Eligibility

Key inclusion criteria

1. Employees of a large international company aged 18 and over
2. Participants scoring 5 or more on the Alcohol Use Disorders Identification Test (AUDIT)-C questionnaire (three-item screening tool for alcohol misuse)

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

1,840

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/08/2012

Date of final enrolment

22/08/2012

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

University College London

London

United Kingdom

NW3 2PF

Sponsor information**Organisation**

University College London (UK)

Sponsor details

c/o David Wilson

Joint Research Office

149 Tottenham Court Road

London

England

United Kingdom

W1T 7DN

+44 (0)20 3447 5199

david.wilson@ucl.ac.uk

Sponsor type

University/education

Website

<http://www.ucl.ac.uk/>

ROR

<https://ror.org/02jx3x895>

Funder(s)

Funder type
Government

Funder Name
NIHR School for Primary Care Research (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	19/11/2014		Yes	No