# The FAB Study: Feedback, Awareness and Behaviour in the Fenland Study

Submission date	Recruitment status  No longer recruiting	Prospectively registered	
29/08/2007		[X] Protocol	
Registration date	Overall study status	Statistical analysis plan	
11/09/2007	Completed	[X] Results	
Last Edited	Condition category	[] Individual participant data	
27/09/2013	Nutritional, Metabolic, Endocrine		

#### Plain English summary of protocol

Not provided at time of registration

# Contact information

#### Type(s)

Scientific

#### 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

# Study information

#### Scientific Title

#### Acronym

**FAB** 

#### Study objectives

Aim: To quantify the effect of measurement and feedback on physical activity behaviour

#### Measurement objectives:

- 1. Self awareness of physical activity levels
- 2. Effect of measurement on awareness
- 3. Effect of feedback on awareness and intentions
- 4. Effect of feedback on behaviour
- 5. Level of false reassurance

More details can be found at: http://www.mrc-epid.cam.ac.uk/Research/Studies/FAB/index.html

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Full ethical approval received from Cambridgshire 2 Research Ethics Committee on 03/05/2007 (ref: 07/Q0108/79)

#### Study design

Cohort study with randomisation to one of four different physical activity feedback groups (following participation in the Fenland Study)

#### Primary study design

Interventional

#### Secondary study design

Randomised controlled trial

#### Study setting(s)

Not specified

#### Study type(s)

**Not Specified** 

#### Participant information sheet

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

Physical activity behaviour

#### **Interventions**

The FAB Study has been set up as a nested cohort study within the Fenland Study, but with a very different purpose, and with separate ethical approval.

Volunteers in the Fenland Study have agreed to undergo oral glucose tolerance test and precise measurement of body anthropometry, body composition and physical activity over a one-week period (using individually calibrated Actihearts which measure heart rate and movement). Those who volunteer for FAB are randomly allocated to one of four groups:

Group 1: Control group - no feedback

Group 2: Physical activity feedback type A - Physical Activity Level (PAL)

Group 3: Physical activity feedback type B - PAL + daily heart rate and movement printouts Group 4: Physical activity feedback type C - PAL + daily heart rate and movement printouts + goal setting and modelling information

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome measure

Objectively measured physical activity over 1 week (one month post-feedback), using a combined heart rate and movement sensor.

#### Secondary outcome measures

- 1. Self-reported physical activity RPAQ (baseline, and 1 month after feedback)
- 2. Self-rated physical activity (baseline, immediately after feedback, and 1 month after feedback)
- 3. Worry about own physical activity (baseline, immediately after feedback, and 1 month after feedback)
- 4. Physical activity outcome expectations (baseline, immediately after feedback, and 1 month after feedback)
- 5. Physical activity self-efficacy (baseline, immediately after feedback, and 1 month after feedback)
- 6. Intention to change physical activity (baseline, immediately after feedback, and 1 month after feedback)
- 7. Time orientation (baseline only)

#### Overall study start date

03/09/2007

#### Completion date

03/06/2008

# **Eligibility**

#### Key inclusion criteria

- 1. Aged 30-55 years
- 2. Registered with participating general practices in the Cambridgeshire Primary Care Trust
- 3. Participating in the Fenland Study

#### Participant type(s)

**Patient** 

#### Age group

Adult

#### Sex

Both

#### Target number of participants

400 (100 per group)

#### Key exclusion criteria

- 1. Have diagnosed diabetes
- 2. Have a terminal illness with a prognosis of less than 1 year
- 3. Suffer from psychotic illness
- 4. Are pregnant or lactating
- 5. Are unable to walk unaided

#### Date of first enrolment

03/09/2007

#### Date of final enrolment

03/06/2008

#### Locations

#### Countries of recruitment

England

United Kingdom

# Study participating centre MRC Epidemiology Unit

Cambridge United Kingdom CB2 0QQ

# Sponsor information

#### Organisation

Medical Research Council (UK)

#### Sponsor details

Medical Research Council Head Office 20 Park Crescent London United Kingdom W1B 1AL corporate@headoffice.mrc.ac.uk

#### Sponsor type

Government

#### **ROR**

https://ror.org/03x94j517

# Funder(s)

#### Funder type

Government

#### **Funder Name**

Medical Research Council (UK)

#### Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

#### **Funding Body Type**

Government organisation

#### **Funding Body Subtype**

National government

#### Location

**United Kingdom** 

# **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?
<u>Protocol article</u>	protocol	18/03/2010		Yes	No
Results article	results	16/09/2013		Yes	No