

Positive Online Weight Reduction Study (POWeR)

Submission date 02/02/2011	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 02/02/2011	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 03/06/2014	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
9575

Study information

Scientific Title
Positive Online Weight Reduction Study (POWeR): a randomised multicentre interventional prevention/process of care trial

Acronym

POWeR

Study objectives

Obesity is a major and rapidly rising public health threat. Recent NICE guidelines provide recommendations to implement lifestyle changes (diet and exercise) supported by behavioural techniques. Unfortunately, an average practice will have more than 1000 patients with obesity, and most practice staff have neither the training nor the time to implement intensive obesity management programmes based on 1:1 counselling, or even group counselling, to cope with such numbers. The problem will become worse as the obesity epidemic progresses. By providing an intervention which requires fewer resources for training and for intervention this study will allow a much greater group of patients to benefit both locally and nationally.

We have developed written behavioural manuals for both patient and practitioner. This study will create a less resource intensive intervention to support behavioural change by converting these materials into web format, taking advantage of a grant to our group which supports the development of generic web programming for behavioural interventions. After materials have been converted to web format, and are acceptable to patients and practitioners, we will then explore the impact of different levels of nurse support required to achieve effective weight change.

Ethics approval required

Old ethics approval format

Ethics approval(s)

IOW, Portsmouth and SE Hampshire Research Ethics Committee approved on the 13th September 2010 (ref: 10/H0501/46)

Study design

Randomised multicentre interventional prevention/process of care trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Topic: Primary Care Research Network for England; Subtopic: Not Assigned; Disease: All Diseases

Interventions

Phase 1:

Study 1: In-depth interviews will be carried out with up to 30 patients from 3 - 4 GP practices to elicit views of the intervention materials (including reactions to content and usability).

Study 2: Up to 5 focus groups will be held for health professionals with practice nurses and GPs from 10 - 20 practices. Participants will be invited to use the training materials and patient website prior to taking part in the focus group to stimulate discussion of the issues relating to content, format and feasibility.

Phase 2:

Pilot RCT: Patients will be offered:

1. Web access and email support
2. Minimal face to face support
3. Intensive face to face visits
4. Normal care

GP records will be reviewed for cost-effectiveness.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Weight, measured at baseline, one month and end of study (6 months)

Key secondary outcome(s)

1. EQ5D, measured at baseline, one month and 6 months
2. Food Frequency Questionnaire, measured at baseline, one month and 6 months
3. Godin Leisure Time Physical Activity Questionnaire, measured at baseline, one month and 6 months
4. GP record review, measured at the end of the study
5. Physical measurements, measured at baseline, one month and 6 months

Completion date

31/05/2013

Eligibility

Key inclusion criteria

1. Patients aged over 18 years, either gender
2. Body mass index (BMI) greater than or equal to 30 (or 28 with hypertension or hypercholesterolaemia) documented in the GP case records

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Lower age limit

18 years

Sex

Not Specified

Key exclusion criteria

1. Current major mental problems (difficulty completing outcomes)
2. Very ill/unable to change diet (e.g. severe left ventricular failure [LVF])
3. Pregnancy/breast feeding
4. Perceived inability to walk 100 metres (i.e. physical activity difficult)

Date of first enrolment

01/12/2010

Date of final enrolment

31/05/2013

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Primary Medical Care

Southampton

United Kingdom

SO16 5ST

Sponsor information

Organisation

University of Southampton (UK)

ROR

<https://ror.org/01ryk1543>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) programme (ref: PB-PG-0808-17077)

Results and Publications

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	21/05/2014		Yes	No