# Training to improve analgesic prescription for chronic pain

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
12/05/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
12/05/2010	Completed	Results
Last Edited	Condition category	Individual participant data
20/04/2017	Signs and Symptoms	<ul><li>Record updated in last year</li></ul>

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Ms Lisa Austin

#### Contact details

University of Bath Norwood house Bath United Kingdom BA2 7AY

## Additional identifiers

Protocol serial number 6840

## Study information

#### Scientific Title

A pilot trial of training in psychological flexibility to improve analgesic prescription for chronic pain in general practice

## **Study objectives**

Can a session of training in psychological flexibility (based on Acceptance and Commitment Therapy [ACT]) have an affect on the way GPs prescribe opioid analgesics to patients with chronic pain and their wellbeing, compared with a control condition?

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

Bath Research Ethics Committee, 01/09/2008, ref: 08/H0101/115

## Study design

Multicentre non-randomised interventional trial

## Primary study design

Interventional

## Study type(s)

Treatment

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

Topic: Generic Health Relevance and Cross Cutting Themes, Primary Care Research Network for England; Subtopic: Generic Health Relevance (all Subtopics); Disease: Other

#### **Interventions**

Intervention group: half a day lecture session with exercises and group work using principles of ACT - psychological flexibility and mindfulness.

Control group: half a day lecture session on guidelines related to pain and exercises and group work based on motivational interviewing techniques.

Follow-up questionnaires two weeks later. Fifty minute lecture on prescribing opioids for persistent pain (all participants).

## Intervention Type

Other

#### Phase

Phase II

## Primary outcome(s)

GP self-reported prescribing, measured at the start of the training day, and two weeks later.

## Key secondary outcome(s))

Psychological acceptance, measured at the start of the training day, at the end of the training day, and two weeks later.

## Completion date

01/10/2011

## **Eligibility**

## Key inclusion criteria

- 1. Must be a general practitioner
- 2. Either sex, any age

## Participant type(s)

**Patient** 

## Healthy volunteers allowed

No

## Age group

Other

#### Sex

All

## Key exclusion criteria

Does not meet inclusion criteria

## Date of first enrolment

01/09/2009

## Date of final enrolment

01/10/2011

## Locations

#### Countries of recruitment

United Kingdom

England

# Study participating centre University of Bath

Bath United Kingdom BA2 7AY

## Sponsor information

## Organisation

University of Bath (UK)

#### **ROR**

https://ror.org/002h8g185

# Funder(s)

## Funder type

Industry

## Funder Name

Reckitt Benckiser (UK)

## **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?

Participant information sheet
Participant information sheet
11/11/2025 No Yes