

# Randomised comparison of nicotine replacement therapy and bupropion in the treatment of tobacco dependence

**Submission date**  
06/08/2003

**Recruitment status**  
No longer recruiting

☒ Prospectively registered

☐ Protocol

**Registration date**  
07/08/2003

**Overall study status**  
Completed

☐ Statistical analysis plan

☒ Results

**Last Edited**  
06/05/2014

**Condition category**  
Injury, Occupational Diseases, Poisoning

☐ Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Mr John Stapleton

### Contact details

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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

## Secondary identifying numbers

G0200455

# Study information

## Scientific Title

## Acronym

ZORN

## Study objectives

To compare effectiveness of the three treatments in routine National Health Service (NHS) care.

## 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)

Not specified

## Study type(s)

Treatment

## Participant information sheet

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

Tobacco dependence

## Interventions

Group and individual specialist counselling over six weeks, plus NRT, bupropion or both

## Intervention Type

Drug

## Phase

Not Specified

## Drug/device/biological/vaccine name(s)

Bupropion

**Primary outcome measure**

Smoking status at four weeks and 12 months

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/05/2004

**Completion date**

30/06/2007

## Eligibility

**Key inclusion criteria**

All smokers eligible to use Nicotine Replacement Therapy (NRT) and bupropion who present for help to stop smoking

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Both

**Target number of participants**

1800

**Key exclusion criteria**

Those with contraindications for NRT or bupropion

**Date of first enrolment**

01/05/2004

**Date of final enrolment**

30/06/2007

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**  
**Institute Of Psychiatry**  
London  
United Kingdom  
SE5 8AF

## **Sponsor information**

**Organisation**  
King's College London (UK)

**Sponsor details**  
Institute of Psychiatry  
De Crespigny Park  
London  
England  
United Kingdom  
SE5 8AF

**Sponsor type**  
University/education

**Website**  
<http://www.iop.kcl.ac.uk/>

**ROR**  
<https://ror.org/0220mzb33>

## **Funder(s)**

**Funder type**  
Government

**Funder Name**  
Department of Health (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?
<a href="#">Results article</a>	results	01/12/2013		Yes	No