

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

Institute Of Psychiatry

Kings College London

Postal Box P075

4 Windsor Walk

London

United Kingdom

SE5 8AF

+44 (0)20 7848 0450

j.stapleton@iop.kcl.ac.uk

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?
Results article	results	01/12/2013		Yes	No