

Nicotine nasal spray for smoking cessation in primary care

Submission date 12/03/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 15/04/2010	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 08/08/2011	Condition category Mental and Behavioural Disorders	<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
9820384; G7804283

Study information

Scientific Title
Randomised placebo controlled trial of nicotine nasal spray for smoking cessation in primary care

Acronym

GPNNNS

Study objectives

To compare the effectiveness of nicotine nasal spray and placebo when given with brief GP and nurse support in primary care.

Ethics approval required

Old ethics approval format

Ethics approval(s)

King's Lynn Medical Research Ethics Committee approved on the 21st November 1995 (ref: 95-NNNS-014 (34/95))

Study design

Multicentre randomised placebo controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Tobacco dependence

Interventions

Active or placebo nicotine nasal spray for 12 weeks plus brief primary care counseling support. Total duration of follow-up is 12 weeks.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Co-verified complete abstinence from smoking for weeks 3 to 12.

Key secondary outcome(s)

1. Co-verified complete abstinence from smoking during 12 weeks
2. Adverse events

Completion date

01/02/1999

Eligibility**Key inclusion criteria**

1. Smokers: smoking for at least 3 years
2. Aged 20 - 60 years, either sex
3. Smokes 15 or more cigarettes a day
4. Motivated to stop

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Myocardial infarction within 3 years
2. Chronic nasal disorder
3. Pregnancy, intending pregnancy
4. Breast feeding
5. Current psychiatric care
6. Current use of psychotropic drugs

Date of first enrolment

01/02/1996

Date of final enrolment

01/02/1999

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Institute Of Psychiatry

London

United Kingdom

SE5 8AF

Sponsor information

Organisation

McNeil AB (Sweden) - formally Pharmacia and Upjohn

ROR

<https://ror.org/020jwmq86>

Funder(s)

Funder type

Research council

Funder Name

Kings College London (UK) - Institute of Psychiatry, staff supported by MRC Programme Grant (ref: G7804283)

Funder Name

McNeil AB (Sweden) - formally Pharmacia and Upjohn, supported trial costs

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	01/04/2011		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes