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A randomised crossover study of citric acid cough challenge in healthy volunteers: Comparison of Mefar Dosimeter and Jaeger Aerosol Provocation System (APS)

Submission date
30/09/2004

Recruitment status
No longer recruiting

Registration date
30/09/2004

Overall study status
Completed

Last Edited
13/04/2018

Condition category
Respiratory

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0084118528

Study information

Scientific Title
A randomised crossover study of citric acid cough challenge in healthy volunteers: Comparison of Mefar Dosimeter and Jaeger Aerosol Provocation System (APS)

Study objectives
Sub-study: To determine whether citric acid inhalation elevates tachykinin and nerve growth factor in sputum of males and females: is this gender related?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled crossover group trial

Primary study design
Interventional

Secondary study design
Randomised cross over trial

Study setting(s)
Hospital

Study type(s)
Other

Participant information sheet
Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Respiratory: Cough challenge

Interventions
Comparison of Mefar Dosimeter and Jaeger Aerosol Provocation System (APS)

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
22/11/2002

Completion date
26/04/2004

Eligibility

Key inclusion criteria
50 healthy volunteers

Participant type(s)
Healthy volunteer

Age group
Not Specified

Sex
Both

Target number of participants
50

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
22/11/2002

Date of final enrolment
26/04/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Castle Hill Hospital
Cottingham

United Kingdom
HU16 5JQ

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
The North and South Bank Research and Development Consortium (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