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A prospective, placebo controlled, double-blind randomised cross over study evaluating the effects of xylitol-containing chewing gum versus ordinary chewing gum on recurrent purulent upper respiratory tract infections

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 25/11/2019	**Condition category** Respiratory	

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
N0256119254

Study information

Scientific Title
A prospective, placebo controlled, double-blind randomised cross over study evaluating the effects of xylitol-containing chewing gum versus ordinary chewing gum on recurrent purulent upper respiratory tract infections

Study objectives
Does xylitol reduce the number of upper respiratory tract infections?

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 cross over trial

Study setting(s)
Not specified

Study type(s)
Not Specified

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

Health condition(s) or problem(s) studied
Upper respiratory tract infections

Interventions
Clinical trial

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Service outcomes development.

Secondary outcome measures
Not provided at time of registration

Overall study start date
15/11/2002

Completion date
15/10/2003

Eligibility

Key inclusion criteria
30 patients aged between 5-65.

Participant type(s)
Patient

Age group
Mixed

Sex
Both

Target number of participants
30

Key exclusion criteria
Does not match inlcusion criteria

Date of first enrolment
15/11/2002

Date of final enrolment
15/10/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Royal Free Hampstead NHS Trust
London
United Kingdom
WC1X 8DA

Sponsor information

Organisation
Department of Health (UK)

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

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
The Royal Free Hampstead NHS Trust (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