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

Plain English summary of protocolNot provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Peter Andrews

Contact details

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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