# A primary care randomised controlled trial of probiotics and xylitol for acute sore throat

Submission date	Recruitment status	[X] Prospectively registered
07/12/2004	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
27/01/2005	Completed	Results
Last Edited	Condition category	Individual participant data
19/02/2013	Respiratory	Record updated in last year
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### Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

### Contact name

Dr Zoe Crosby

### Contact details

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# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

RSO2540

# Study information

### Scientific Title

### **Study objectives**

Not provided at time of registration

### 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)

GP practice

### Study type(s)

Treatment

### Participant information sheet

# Health condition(s) or problem(s) studied

Acute sore throat

#### Interventions

Probiotics and xylitol (or placebos)

### Intervention Type

Other

#### Phase

Not Applicable

### Primary outcome measure

Not provided at time of registration

### Secondary outcome measures

Not provided at time of registration

### Overall study start date

### Completion date

31/07/2007

# **Eligibility**

### Key inclusion criteria

Adults attending primary care with acute sore throat

### Participant type(s)

**Patient** 

### Age group

**Not Specified** 

#### Sex

**Not Specified** 

### Target number of participants

Not provided at time of registration

### Key exclusion criteria

Not provided at time of registration

### Date of first enrolment

01/02/2005

### Date of final enrolment

31/07/2007

# Locations

### Countries of recruitment

England

**United Kingdom** 

# Study participating centre Department of Primary Medical Care

Southampton United Kingdom SO16 5ST

# Sponsor information

### Organisation

University of Southampton (UK)

### Sponsor details

Dr Peter Hooper Research Support Office University of Southampton Highfield Southampton England United Kingdom SO17 1BJ +44 (0)2380 598672 info@rso.soton.ac.uk

### Sponsor type

University/education

#### **ROR**

https://ror.org/01ryk1543

# Funder(s)

### Funder type

Government

#### **Funder Name**

Department of Health National Co-ordinating Centre for Research Capacity Development (DoH NCC RCD) Research Fellowship (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