

Effect of Fluoride Slow-release Devices on salivary and plaque levels of fluoride, phosphorus and calcium

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 26/02/2020	Condition category Oral Health	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0436133177

Study information

Scientific Title

Effect of Fluoride Slow-release Devices on salivary and plaque levels of fluoride, phosphorus and calcium

Study objectives

The aim of our study is to evaluate the levels of fluoride, calcium and phosphorus in saliva and in dental plaque from the release of fluoride slow-release devices.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised placebo controlled parallel group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Dental materials

Interventions

Randomised controlled trial. Two types of glass devices (one with fluoride, the other without) are randomly attached to the buccal side of maxillary molar.

Intervention Type

Device

Phase

Not Specified

Primary outcome measure

Measurements of levels of fluoride, calcium and phosphorus in saliva and dental plaque

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2003

Completion date

01/07/2004

Eligibility

Key inclusion criteria

Healthy adults aged 18-65 years residing in Leeds

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

65 Years

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/09/2003

Date of final enrolment

01/07/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
The Worsley Building
Leeds
United Kingdom
LS2 9LU

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
Hospital/treatment centre

Funder Name
Leeds Teaching Hospitals 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Thesis results	doctoral thesis			No	No