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A study of post-extraction bleeding following warm saline mouth washes compared to no mouth washes during the 24 hours following surgery

Submission date
30/09/2005

Recruitment status
No longer recruiting

Registration date
30/09/2005

Overall study status
Completed

Last Edited
02/06/2017

Condition category
Oral Health

[_] 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
N0205153820

Study information

Scientific Title
A study of post-extraction bleeding following warm saline mouth washes compared to no mouth washes during the 24 hours following surgery

Study objectives
1. Does rinsing of the mouth after dental extractions of tooth induce excessive bleeding?
2. Are traditional post-extraction instructions evidence based?
3. To improve, simplify and validate post-operative instructions after tooth removal

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)
Other

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Dental extractions of tooth

Interventions
A randomised two-group study:
1. Warm saline mouth washes
2. No mouth wash

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
Absence/occurrence of bleeding over 24 hours.

Secondary outcome measures
No secondary outcome measures

Overall study start date
29/11/2004

Completion date
28/02/2005

Eligibility

Key inclusion criteria
60 patients in two groups of 30

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
60 patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
29/11/2004

Date of final enrolment
28/02/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Ground Floor
London
United Kingdom
E1 1BB

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
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United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
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Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Barts and The London NHS Trust (UK)

Funder Name
NHS R&D Support Funding (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