

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

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 02/06/2017	<b>Condition category</b> Oral Health	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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### Contact details

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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  
London  
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