

# Fluoride glass device for reducing tooth sensitivity

|                                        |                                                   |                                                                                                              |
|----------------------------------------|---------------------------------------------------|--------------------------------------------------------------------------------------------------------------|
| <b>Submission date</b><br>29/09/2006   | <b>Recruitment status</b><br>No longer recruiting | <input type="checkbox"/> Prospectively registered<br><input type="checkbox"/> Protocol                       |
| <b>Registration date</b><br>29/09/2006 | <b>Overall study status</b><br>Completed          | <input type="checkbox"/> Statistical analysis plan<br><input type="checkbox"/> Results                       |
| <b>Last Edited</b><br>11/10/2017       | <b>Condition category</b><br>Oral Health          | <input type="checkbox"/> Individual participant data<br><input type="checkbox"/> 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

**Protocol serial number**  
N0436165569

## Study information

**Scientific Title**  
Fluoride glass device for reducing tooth sensitivity

**Study objectives**

The aim is to compare the slow release fluoride glass device with a desensitising tooth paste to alleviate tooth sensitivity.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Primary study design**

Interventional

**Study design**

Randomised controlled trial

**Study type(s)**

Treatment

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

Oral health

**Interventions**

Randomised controlled trial

**Intervention Type**

Device

**Phase**

Not Specified

**Primary outcome(s)**

Alleviation of tooth sensitivity

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/02/2006

**Eligibility**

**Key inclusion criteria**

Population from Yorkshire

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**Key exclusion criteria**

Does not meet inclusion criteria

**Date of first enrolment**

01/02/2005

**Date of final enrolment**

01/02/2006

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre****Leeds Dental Institute**

Leeds

United Kingdom

LS2 9LU

**Sponsor information****Organisation**

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

**Funder(s)****Funder type**

Government

**Funder Name**

Leeds Teaching Hospitals NHS Trust (UK)

**Funder Name**

NHS R&D Support Funding (UK)

**Results and Publications**

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**

Not provided at time of registration