

A clinical comparison of self-etching primer with conventional etch during fixed appliance treatment

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 06/12/2007	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
N0155160660

Study information

Scientific Title

Study objectives

Does self etching primer provide satisfactory clinical performance with similar failure rates compared to conventional etch?

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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Oral Health

Interventions

Randomised controlled trial: self-etching primer compared to conventional etch.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Percentage bond failure rate.

Secondary outcome measures

Not provided at time of registration.

Overall study start date

01/12/2003

Completion date

30/12/2006

Eligibility

Key inclusion criteria

Patients requiring fixed appliance treatment.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

40

Key exclusion criteria

Patients who require bands or have heavily restored teeth.

Date of first enrolment

01/12/2003

Date of final enrolment

30/12/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Dept of Oral and Maxillofacial Surgery

Bury

United Kingdom

BL9 7TD

Sponsor information

Organisation

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

Sponsor details

The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)**Funder type**

Government

Funder Name

Pennine Acute Hospitals 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/12/2007		Yes	No