

A randomised controlled clinical trial comparing aesthetic arch wires and nickel titanium arch wires in initial alignment

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 type="checkbox"/> Results
Last Edited 06/10/2014	Condition category 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
Dr D Bearn

Contact details
Central Manchester & Manchester Children's Uni Hospitals NHS Trust
Dental Hospital
Higher Cambridge Street
Manchester
United Kingdom
M15 6FH

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0453173840

Study information

Scientific Title

Study objectives

To compare two different types of orthodontic braces to see if one works faster than the other.

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

Oral Health: Orthodontics

Interventions

Type A of orthodontic braces vs type B of orthodontic braces

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Time taken to complete levelling and aligning of front teeth

Secondary outcome measures

Not provided at time of registration

Overall study start date

17/10/2005

Completion date

16/10/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Child

Sex

Not Specified

Target number of participants

40 patients in each treatment group. No control group. Total = 80

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

17/10/2005

Date of final enrolment

16/10/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Central Manchester & Manchester Children's Uni Hospitals NHS Trust

Manchester

United Kingdom

M15 6FH

Sponsor information

Organisation

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

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

Central Manchester and Manchester Children's University 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

