A study comparing the construction of stabilization splints using two new methodologies

Submission date	Recruitment status	Prospectively registered
29/09/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Completed	Results
Last Edited	Condition category	Individual participant data
02/02/2015	Oral Health	[] Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof J F McCord

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0453177298

Study information

Scientific Title

A study comparing the construction of stabilization splints using two new methodologies

Study objectives

To compare two new methodologies:

- 1. Clinical Digital gothic arch tracing
- 2. Technical Injection moulded flasking

Against the standard norm:

- 1. Clinical Facebow with bimanual manipulation
- 2. Technical Compression moulded flasking

In the provision of stabilization splints at the University of Manchester Dental Hospital.

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Oral Health

Interventions

Methodology 1 vs methodology 2 vs norm

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

21/02/2006

Completion date

01/07/2008

Eligibility

Key inclusion criteria

Using 25 dental school students

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

25

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

21/02/2006

Date of final enrolment

01/07/2008

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

DEN Central Manchester & Manchester Children's University Hospitals

Manchester United Kingdom M15 6FH

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

Hospital/treatment centre

Funder Name

Central Manchester and Manchester Children's University Hospitals NHS Trust (UK)

Funder Name

NHS R&D Support Funding

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