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

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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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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