# Improving dentures for patient benefit

Submission date [X] Prospectively registered Recruitment status 11/11/2009 No longer recruiting [X] Protocol [ ] Statistical analysis plan Registration date Overall study status 04/01/2010 Completed [X] Results [ ] Individual participant data **Last Edited** Condition category 24/08/2016 Other

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

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## Additional identifiers

#### Protocol serial number

N/A

## Study information

#### Scientific Title

Improving dentures for patient benefit: a single centre randomised, controlled, crossover clinical trial comparing materials for complete dentures

#### Acronym

**IMPROVDENT** 

### Study objectives

The aim of this study is to significantly impact on the quality of denture production within NHS dentistry. To achieve this objective, we aim to provide evidence of effectiveness and cost-effectiveness of a development in denture impression materials. The aim is to ask the patients to provide the outcome assessments for the trial; a user-centered approach which will give clinically significant results, empower research participants and facilitate a deeper understanding of patients' denture wearing problems.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Not provided at time of registration

### Study design

Single-centre randomised controlled cross-over clinical trial

### Primary study design

Interventional

## Study type(s)

Treatment

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

Quality of denture production

#### **Interventions**

Patients should be assessed for their suitability to participate in this study before they give their consent and are registered into the trial.

#### Denture construction:

The Prosthodontic Research Team at the Leeds Dental Institute has developed and published a method of producing two or more sets of dentures which are clinically identical apart from the one deliberate alteration which is to be the subject of this trial. This methodology will be used to produce two sets of dentures which are clinically identical apart from the impression material that was used to provide the mould for the fitting surface of each denture.

#### Habituation:

Patients will be given both sets of dentures for a 2-week habituation period. During the 2 weeks the patients will be provided with a diary and asked to record the wearing of each set of dentures. They will then be asked to score each denture for comfort, mobility and chewing efficiency using a 5-point Likert scale. Both sets of dentures will be returned and re-marked for the adjustment stage. The code will be blind to clinicians and patients, and kept confidential until the end of the trial.

#### Adjustment:

Dentures will be given back to patients, one at a time over two adjustment periods (8 weeks each), in random order. If any adjustment or remedial work is required to make the dentures comfortable, retentive and useful this will be carried out by an experienced clinician. Within the limits of the LDI appointment system there will be no restriction on the number of appointments the patients can request during these two 8-week periods. Every appointment

requested by the patient will be noted and a detailed record of the work required will be kept. At the end of each 8-week adjustment period the patient will be asked to attend and report the impact the dentures have had on their quality of life by the completion of a OHIP-EDENT and EQ-5D questionnaires and will rate the denture for comfort, mobility and chewing efficiency using 5-point Likert Scales.

#### Confirmation:

After the two adjustment periods of 8 weeks it is to be expected that each set of dentures will be as good as it is possible to make them. The patient will be asked to take both sets of dentures away for a 2-week confirmation period and return for a final visit to express any preference for the dentures and perform a Likert assessment of comfort, mobility, and chewing efficiency.

#### Intervention Type

Other

#### Phase

Not Applicable

### Primary outcome(s)

Patient preference for unadjusted dentures and overall cost-effectiveness, measured at the end of the habituation period. Patient preference will be established by asking whether one or other denture is preferred, or whether there is no preference. Those expressing no preference will be further asked whether both dentures were satisfactory or unsatisfactory.

## Key secondary outcome(s))

Measured at the end of the secondary impressions, habituation, adjustment and confirmation periods:

- 1. Comfort, Mobility, Chewing Efficiency using 5-point Likert Scales
- 2. Oral Health-Related Quality of Life using OHIP-EDENT
- 3. Patient preference for adjusted dentures
- 4. Patient perception of comfort, taste and ease of use of each impression material using a 5-point Likert scale
- . 5. Qualitative interviews to gain insight into the impact of oral health status on daily lives

Patient preference for adjusted dentures will be established by asking whether one or other denture is preferred, or whether there is no preference. Those expressing no preference will be further asked whether both dentures were satisfactory or unsatisfactory. Patients will be asked to reflect on the whole process of receiving the final denture (including initial comfort, amount of adjustment needed) when expressing preference.

## Completion date

15/01/2012

## **Eligibility**

## Key inclusion criteria

- 1. Patients of either sex who are edentulous
- 2. Subject is available for follow up
- 3. Subject requires replacement complete dentures
- 4. Subject is able and willing to sign the Informed Consent Form
- 5. Aged 18 years or over at the time of signing the Informed Consent Form

## Participant type(s)

Patient

## Healthy volunteers allowed

No

### Age group

Adult

### Lower age limit

18 years

#### Sex

All

#### Key exclusion criteria

- 1. Presence of an oral tumour
- 2. Requirement for an obturator
- 3. Extreme xerostomia (e.g. Sjogren's Syndrome)
- 4. Patients who would benefit from selective pressure impressions
- 5. Known hypersensitivity to silicone or alginate

#### Date of first enrolment

15/01/2010

#### Date of final enrolment

15/01/2012

## Locations

### Countries of recruitment

United Kingdom

England

## Study participating centre Leeds Dental Institute

Leeds United Kingdom LS2 9LU

## **Sponsor information**

### Organisation

University of Leeds (UK)

#### **ROR**

https://ror.org/024mrxd33

## Funder(s)

## Funder type

Government

#### Funder Name

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) programme (ref: PB-PG-0408-16300)

## **Results and Publications**

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	01/08/2014	Yes	No
Results article	results	01/08/2014	Yes	No
Protocol article	protocol	31/08/2012	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes