

# The effect of viewing digital images of the severe consequences of poor oral hygiene on patients' adherence to good oral hygiene practice

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<b>Registration date</b> 09/01/2024	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 09/01/2024	<b>Condition category</b> Oral Health	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

Background and study aims.  
Adhering to good oral hygiene is vital for people's oral health and dental treatment success. However, it is difficult to achieve good oral hygiene adherence in dental patients. Several studies have explored different methods to increase patients' oral hygiene adherence. For instance, studies have used verbal oral hygiene instructions, while others employed written leaflets or digital oral hygiene instructions. This pilot trial aims to study the effect of oral hygiene instructions given via video while supplemented with images showing the severe consequences of poor oral hygiene on patients' oral hygiene adherence.

Who can participate?  
Healthy female adults aged 18-30 years

What does the study involve?  
The participants will be divided into three oral hygiene instruction groups:  
Group 1: They will receive verbal oral hygiene instructions  
Group 2: They will receive video-based oral hygiene instructions  
Group 3: They will receive video-based oral hygiene instructions similar to group 2 but will also be shown images of the severe consequences of poor oral hygiene  
The oral hygiene instructions (video or verbal) will be given to each participant at the initial visit. The researchers will also measure participants' oral hygiene status at the initial visit using three methods:  
1. Plaque accumulation on teeth surfaces after rinsing with a discoloring solution.  
2. The percentage of teeth that exhibit gum bleeding  
3. Gum bleeding index based on the total percentages calculated from method number 2.

What are the possible benefits and risks of participating?

This study involves no expected risks to participants. This study's obtained results will be of great importance for dental practitioners and especially for orthodontic specialists in the control of optimum oral hygiene in pre-orthodontic patients.

Where is the study run from?

Qassim University (Saudi Arabia)

When is the study starting and how long is it expected to run for?

May 2022 to November 2023

Who is funding the study?

Investigator initiated and funded

Who is the main contact?

Dr Nabeel Almotairy, n.s.almotairy@qu.edu.sa

## Contact information

### Type(s)

Public, Scientific, Principal Investigator

### Contact name

Dr Nabeel Almotairy

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## Additional identifiers

### EudraCT/CTIS number

Nil known

### IRAS number

### ClinicalTrials.gov number

Nil known

### Secondary identifying numbers

Nil known

## Study information

**Scientific Title**

Clinical efficacy of verbal versus video oral hygiene instructions in reducing plaque accumulation and gingival bleeding in healthy female adults

**Study objectives**

The null hypothesis is that showing patients' digital images displaying the consequences of poor oral hygiene (OH) will not influence their oral hygiene adherence.

**Ethics approval required**

Ethics approval required

**Ethics approval(s)**

Approved 01/06/2022, Committee of Research Ethics at Qassim University (AlQassim, Buraidah, 51452, Saudi Arabia; +966 (0)163010355; bioethics@qu.edu.sa), ref: 21-19-05

**Study design**

Prospective single-blinded randomized clinical three-arm parallel trial

**Primary study design**

Interventional

**Secondary study design**

Randomised parallel trial

**Study setting(s)**

University/medical school/dental school

**Study type(s)**

Prevention

**Participant information sheet**

Not available in web format, please use contact details to request a participant information sheet.

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

Oral hygiene adherence in healthy female adults

**Interventions**

The experiment will be conducted by a dentist with a minimum of 2 years' experience.

Using the randomization.com website, a computer-generated random selection list with a 1:1:1 allocation ratio for the three intervention groups will be prepared. Sealed opaque envelopes will be prepared following the generated list to reduce the risk of concealing the group allocation to the examining dentists before recruiting the study participants. Trained and calibrated dentists who are not involved in generating the random list or creating the sealed envelope will be responsible for recruiting the participants and collecting the oral health parameters. After ensuring the participation eligibility, each participant will be allowed to draw an envelope to be assigned to one of the three groups.

The participants will be randomly divided into three groups of patients:

1. The group will be given verbal oral hygiene instructions
2. The group will be shown a one-to-one 5-minute video-based presentation containing recommended brushing and flossing techniques WITH images displaying the consequences of poor oral hygiene will be presented
3. The group will be shown a one-to-one 5-minute video-based presentation containing recommended brushing and flossing techniques WITHOUT images displaying the consequences of poor oral hygiene will be presented.

The verbal or video-based oral hygiene instructions will be given at the baseline visit only. Each participant will be asked if she understood the verbal or video-based instructions, and she will be given the chance for any clarifications. Then, their oral hygiene status will be assessed using the below outcome measures.

The participants will come back after 4 weeks to assess their oral hygiene status.

### **Intervention Type**

Behavioural

### **Primary outcome measure**

Plaque score: a modified O'Leary Index is used to define the number of surfaces free of plaque as a percentage of surfaces in the mouth following a rinse with a discoloring solution — six surfaces will be assessed per tooth at baseline and 4 weeks post-intervention.

### **Secondary outcome measures**

1. Gingival bleeding index as follows: 0: no bleeding on probing, 1: a single bleeding point, 2: several bleeding points or thin line, 3: interdental space filled with blood
  2. Bleeding on probing: the percentage of teeth surface numbers that are free from gingival bleeding — four surfaces per tooth.
- Each will be measured at baseline and 4 weeks post-intervention.

### **Overall study start date**

15/05/2022

### **Completion date**

20/11/2023

## **Eligibility**

### **Key inclusion criteria**

1. Have at least 20 natural teeth
2. Be willing to participate in this study and sign the consent form

### **Participant type(s)**

Healthy volunteer

### **Age group**

Adult

### **Lower age limit**

18 Years

**Upper age limit**

30 Years

**Sex**

Female

**Target number of participants**

21

**Total final enrolment**

21

**Key exclusion criteria**

1. Students in medical or dental fields
2. Under current dental or orthodontic treatment
3. Active smoker
4. Has acute or chronic immune disturbances
5. Currently pregnant
6. Have any systemic diseases that could directly or indirectly inhibit optimum oral health maintenance, such as diabetes, known antibiotic treatments, and oral-related diseases such as periodontitis
7. Has insufficient Arabic language skills than can inhibit understanding of the study goals and oral health instructions

**Date of first enrolment**

15/08/2023

**Date of final enrolment**

20/10/2023

## **Locations**

**Countries of recruitment**

Saudi Arabia

**Study participating centre**

**Qassim University**

College of Dentistry

AlQassim

Buraidah

Saudi Arabia

51452

## **Sponsor information**

**Organisation**

Qassim University

**Sponsor details**

AlQassim, Buraidah  
Buraidah  
Saudi Arabia  
51452  
+966 (0)163010355  
bioethics@qu.edu.sa

**Sponsor type**

University/education

**Website**

<http://www.qu.edu.sa/Pages/Home.aspx>

**ROR**

<https://ror.org/01wsfe280>

**Funder(s)****Funder type**

Other

**Funder Name**

Investigator initiated and funded

**Results and Publications****Publication and dissemination plan**

Planned publication in a high-impact peer-reviewed journal.

**Intention to publish date**

10/04/2024

**Individual participant data (IPD) sharing plan**

The dataset generated during and/or analysed during the current study will be available upon request from Dr Nabeel Almotairy (n.s.almotairy@qu.edu.sa). The type of data that will be shared is currently not known. The timing for availability is upon a reasonable request. Informed consent was obtained.

**IPD sharing plan summary**

Available on request

