

# Comparison of two dental isolation tools used during routine procedures: Rubber Dam and OptiDam

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<b>Registration date</b> 18/07/2025	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 12/09/2025	<b>Condition category</b> Oral Health	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

This study aims to compare the ease of application, time for placement, patient satisfaction, and comfort during common use of the OptiDam versus the conventional rubber dam (CRD) in routine dental practice at Umm Al-Qura University Dental Teaching Hospital.

### Who can participate?

Patients receiving dental treatment at Umm Al-Qura University Dental Teaching Hospital

### What does the study involve?

Participants are randomly allocated to be treated with Rubber Dam or OptiDam during their dental treatment.

### What are the possible benefits and risks of participating?

The possible benefits include improved comfort and efficiency, enhanced quality of care, and contribution to dental research with no extra costs. The risks involved are minimal and similar to those associated with the use of rubber dams in standard dental procedures. If you experience discomfort, the rubber dam can be adjusted or removed.

### Where is the study run from?

Umm Al-Qura University (Saudi Arabia)

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

December 2024 to August 2025

### Who is funding the study?

Investigator initiated and funded

### Who is the main contact?

Dr Rashed Binqali, s441007974@uqu.edu.sa

# Contact information

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# Additional identifiers

## Protocol serial number

HAPO-02-K-012-2024-12-2377

# Study information

## Scientific Title

A clinical comparison of the Rubber Dam and OptiDam in routine dental procedures at Umm Al-Qura University Dental Hospital

## Study objectives

Compare between two isolation systems:

1. Ease of application
2. Time for placement
3. Patient satisfaction
4. Patient comfort

## Ethics approval required

Ethics approval required

## Ethics approval(s)

approved 09/12/2024, Umm Al-Qura University Faculty of Dentistry Biomedical Research Ethics Committee (Taif Road, Al-abdiyyah, Makkah, 24353, Saudi Arabia; +966 (0)125270000; cscenter@uqu.edu.sa), ref: JTIR141124

## Study design

Randomized controlled trial

## Primary study design

Interventional

## Study type(s)

Safety

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

Moisture control during dental procedures

## Interventions

Patients will be assigned to one of two groups (1:1) using computer-generated randomisation:

1. Conventional rubber dam (control group)
2. OptiDam (intervention group)

Treatment duration: each procedure will last 1 to 2 hours.

Follow-up duration: immediate post-procedure evaluation only.

## Intervention Type

Procedure/Surgery

## Primary outcome(s)

1. Time to place isolation system, recorded by using a stopwatch starting at the initiation of isolation placement and ending when the isolation is stable in place and the tooth is ready for treatment
2. Operator preference assessed by questionnaire immediately after placing the isolation system
3. Patient comfort assessed using a Visual Analog Scale (VAS) immediately after the dental

procedure

4. Patient satisfaction assessed by questionnaire after the dental procedure, before patient discharge

### **Key secondary outcome(s)**

There are no secondary outcome measures

### **Completion date**

17/08/2025

## **Eligibility**

### **Key inclusion criteria**

1. Systematically healthy patients: American Society of Anesthesiologists (ASA) I or Controlled ASA II
2. Undergoing routine dental procedures

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

Patient, Learner/student

### **Healthy volunteers allowed**

No

### **Age group**

Mixed

### **Lower age limit**

15 years

### **Upper age limit**

60 years

### **Sex**

All

### **Total final enrolment**

114

### **Key exclusion criteria**

1. Latex allergy
2. Severe gag reflex

### **Date of first enrolment**

10/12/2024

### **Date of final enrolment**

07/08/2025

## **Locations**

## Countries of recruitment

Saudi Arabia

## Study participating centre

**Umm Al-Qura University**

Taif Road

Al-Abdiyyah

Makkah

Saudi Arabia

24243

## Sponsor information

### Organisation

Umm al-Qura University

### ROR

<https://ror.org/01xjqrm90>

## Funder(s)

### Funder type

Other

### Funder Name

Investigator initiated and funded

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated and/or analyzed during the current study are not expected to be made available, as the data can be verified for only validity purposes if required by an authority that has the right to check our result upon request from the author, Rashed Binqali. The data is quantitative data and will be stored in Dr. Omair M. Bukhari's office at Umm Al-Qura University for two years. The data can be accessed by the research team only. All data was taken from the participants after signing an electronic consent to use their data for the purpose of this study only.

### IPD sharing plan summary

Not expected to be made available

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes