

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jul 2026

Comparison of the effect of intracanal drugs on calcium hydroxide blend with nano silver particles and calcium hydroxide blend and normal saline on pain relief between root canal therapy in patients with symptomatic apical periodontitis

Protocol summary

Study aim

Comparison of the effect of calcium hydroxide with nano-silver particles and calcium hydroxide and normal saline as intracanal medications on pain between sessions in premolar teeth with symptomatic apical periodontitis.

Design

In this single-blind randomized clinical trial study, 30 patients with healthy systemic conditions and premolar teeth with pulp and periapical conditions requiring root canal treatment with symptomatic apical periodontitis were selected. Patients were divided into 3 parallel 10-person groups including study and control groups, and patients in each group received different root canal medications. To choose intracanal medication 10 numbers of each intracanal medications were written on identical pieces of paper then asked the assistant to one sheet of paper for each patient.

Settings and conduct

Study Area: Dental Clinic, Faculty of Dentistry, Army University of Medical Sciences. pain between the root canal treatment sessions was measured using visual analogue scale (VAS) and its relationship with intracanal medications was evaluated.

Participants/Inclusion and exclusion criteria

Inclusion criterion: patient with premolar tooth with closed apex and symptomatic apical periodontitis. Non-inclusion criteria: pregnancy; Not repairable teeth; presence of any resorption; systemic disease; history of chronic use of Analgesics, drugs or alcohol; using antibiotics and pretreatment analgesics.

Intervention groups

Intervention group 1: mixture of calcium hydroxide with silver nano particles. Intervention group 2: Mixture of calcium hydroxide with normal saline. Control group: Sterile dry cotton pellet. Standard root canal treatment was performed for patients in two sessions and at the

end of the first session, the intracanal medications were used for patients in the study groups.

Main outcome variables

Pain

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190323043102N1**

Registration date: **2020-03-14, 1398/12/24**

Registration timing: **retrospective**

Last update: **2020-03-14, 1398/12/24**

Update count: **0**

Registration date

2020-03-14, 1398/12/24

Registrant information

Name

Mohsen Mortezaei

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-07-01, 1398/04/10

Expected recruitment end date

2019-10-03, 1398/07/11
Actual recruitment start date
 2019-07-01, 1398/04/10
Actual recruitment end date
 2019-12-30, 1398/10/09
Trial completion date
 2019-12-30, 1398/10/09

Scientific title
 Comparison of the effect of intracanal drugs on calcium hydroxide blend with nano silver particles and calcium hydroxide blend and normal saline on pain relief between root canal therapy in patients with symptomatic apical periodontitis

Public title
 Effect of nano silver particles on pain control between root canal therapy sessions in premolar teeth

Purpose
 Prevention

Inclusion/Exclusion criteria
Inclusion criteria:
 Patient seeking root canal treatment closed apex
Exclusion criteria:
 Systemic disease tooth with internal or external resorption Use of antibiotics and analgesics before treatment pregnancy Non-repairable teeth A history of chronic use of analgesics, narcotic or alcohol

Age
 No age limit

Gender
 Both

Phase
 1-2

Groups that have been masked

- Participant
- Outcome assessor
- Data analyzer
- Data and Safety Monitoring Board

Sample size
 Target sample size: **30**
 Actual sample size reached: **30**

Randomization (investigator's opinion)
 Randomized

Randomization description
 The patients were categorized by an assistant using random numbers table.

Blinding (investigator's opinion)
 Single blinded

Blinding description
 number 1 was assigned to calcium hydroxide with nano silver particles , number 2 was assigned to calcium hydroxide , number 3 was assigned to sterile dry cotton pellet. Then, on behalf of the intracanal drug type, 10 numbers of each intracanal medication were written on identical so that the numbers could not be identified. The papers were poured into a box and we asked the ward assistant to remove one sheet of paper for each patient and enter it into the file. The patient and data analyzer were unaware of the root canal drug type and after data collection, the number assigned to each intracanal

medication was reported to the data analyzer instead of the type of intracanal medication.

Placebo
 Used

Assignment
 Parallel

Other design features
 Silver nano particles are a new material in dentistry and most of the studies on this material are about its antibacterial properties. Microorganisms are the main cause of pain between root canal treatment sessions with apical periodontitis. So we decided to design a study about the relation of the pain between root canal therapy sessions and nano silver particles.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of AJA University of Medical Sciences

Street address

AJA University of Medical Sciences, Etemadzade avenue., West Fatemi boulevard, Tehran

City

Tehran

Province

Tehran

Postal code

6523545522

Approval date

2019-01-28, 1397/11/08

Ethics committee reference number

IR.AJAUMS.REC.1397.082

Health conditions studied

1

Description of health condition studied

Pain between root canal treatment sessions

ICD-10 code

K04.4

ICD-10 code description

Acute apical periodontitis of pulpal origin

Primary outcomes

1

Description

Post-treatment pain above 3

Timepoint

between two root canal sessions at 6, 12 , 24 and 48 hours after first session (2 session root canal with intracanal medication)

Method of measurement

Visual analog scale (VAS): patients are asked to mark the pain level at 6, 12, 24 and 48 hours after first session of their treatment then the 10 cm line of each period was measured from left (no pain) in millimeter.

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group 1: In patient's premolar teeth with symptomatic apical periodontitis, calcium hydroxide with silver nano particles was used as intracanal medication one time between root canal therapy sessions at the end of the first treatment session after cleaning and shaping of root canal system for 4 days.

Category

Prevention

2

Description

Intervention group 2: In patient's premolar teeth with symptomatic apical periodontitis, calcium hydroxide with normal saline was used as intracanal medication one time between root canal therapy sessions at the end of the first treatment session after cleaning and shaping of root canal system for 4 days.

Category

Prevention

3

Description

Control group: In patient's premolar teeth with symptomatic apical periodontitis, sterile dry cotton pellet was used as placebo medication one time between root canal therapy sessions at the end of the first treatment session after cleaning and shaping of root canal system for 4 days.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Dentistry college of AJA University of Medical Science

Full name of responsible person

Mohsen Mortezaei

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Misagh complex, East 13 alley, Sabari street

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Artesh University of Medical Sciences

Full name of responsible person

Dr. Majid Ajami

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<http://www.ajaums.ac.ir/>

Grant name

Grant code / Reference number

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Artesh University of Medical Sciences

Proportion provided by this source

100

Public or private sector

Public

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Academic

Person responsible for general inquiries

Contact

Name of organization / entity

Artesh University of Medical Sciences

Full name of responsible person

Mohsen Mortezaei

Position

student

Latest degree

A Level or less
Other areas of specialty/work
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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

Patient record, file summary, visual analogue scale (VAS) form, and 5 periapical graphs of patients are available which can be published if necessary after deleting patients' names and telephone numbers.

When the data will become available and for how long

Start of access period 6 months after printing results

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

Those who are researching silver nanoparticles
Comparison of pain levels between sessions in the upper and lower jaw
Comparison of pain between sessions between men and women

From where data/document is obtainable

mmortezaei7777@gmail.com Department of Endodontics, Faculty of Dentistry, Army University of Medical Sciences

What processes are involved for a request to access data/document

The reason for the data request should be verified by the endodontic department.

Comments