

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

Evaluation of using Acellular dermal matrix (ADM) and autogenous free gingival strap on clinical conditions of gingiva

Protocol summary

Study aim

A comparative study of the clinical conditions of gingiva after transplantation by the acellular dermal matrix (ADM) and a narrow free gingiva strap with ADM alone.

Design

This study is a split-mouth randomized controlled clinical trial. 11 patients are enrolled with entry criteria. In cases, each half-jaw was randomized in the test group (Acellular dermal matrix and autogenous free gingival strap) or in the control group (Acellular dermal matrix).

Settings and conduct

This research is done at the Faculty of Dentistry, Islamic Azad University, Tehran. In the control group, ADM was fixed at 7 mm wide, and in the test group, the coronal region of receptor site was covered up to marginal margin by Acellular dermal matrix with a width of 5 mm and free gingival strap was made up to a width of 2 mm from the palate and placed in the apical region of the receptor site.

Participants/Inclusion and exclusion criteria

Entry requirements: A patient has one to two teeth on each mandibular side with insufficient keratinized tissue (1 mm or less). The patient has no systemic disease diagnosed with effect on periodontal tissues. Conditions of non-arrival: Patients lacking entry criteria and non-compliance with essential follow-ups.

Intervention groups

The following measurements will be made on the facial surface of each tooth including the width of attached gingiva (AT), the width of keratinized gingiva (KT) and the pocket depth (PD) on all people entering the study.

Main outcome variables

Keratinized gingiva, Attached gingiva width, probe depth, Shrinkage percentage.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180714040460N4**

Registration date: **2019-06-15, 1398/03/25**

Registration timing: **retrospective**

Last update: **2019-06-15, 1398/03/25**

Update count: **0**

Registration date

2019-06-15, 1398/03/25

Registrant information

Name

Nima Nadafpour

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 2256 4571

Email address

n_nadaf@dentaliau.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-09-23, 1397/07/01

Expected recruitment end date

2019-01-21, 1397/11/01

Actual recruitment start date

2018-11-07, 1397/08/16

Actual recruitment end date

2019-02-13, 1397/11/24

Trial completion date

2019-05-14, 1398/02/24

Scientific title

Evaluation of using Acellular dermal matrix (ADM) and autogenous free gingival strap on clinical conditions of gingiva

Public title

The effect of simultaneous use of Acellular dermal matrix and autogenous free gingival strap on formation of keratinized gingiva

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

The patient should have one to two teeth on each mandibular side with insufficient keratinized tissue (1 mm or less). The patient should not have systemic disease diagnosed with effects on periodontal tissues.

Exclusion criteria:

The patient that does not want to cooperate on the project . The patient with poor oral hygiene .

Age

No age limit

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Outcome assessor
- Data analyser

Sample size

Target sample size: **11**

Actual sample size reached: **11**

Randomization (investigator's opinion)

Randomized

Randomization description

Divide the samples (each jaw) completely randomly into two groups of case and control by dropping the coin and dividing it into two groups of alloderm (control) and alloderm and autogenous free gingival strap (case)

Blinding (investigator's opinion)

Double blinded

Blinding description

All individuals involved in this research, such as clinicians, outcome evaluators, data analyzers, have been unaware of the randomization process of cases, and only researchers and operators in this study are aware of the allocation groups.

Placebo

Not used

Assignment

Parallel

Other design features

The division of samples into two groups test and control was based on the type of graft technique.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Faculty of Dentistry Islamic Azad University, Tehran Branch

Street address

Pasdaran Street, Ninth Neyestan Avenue

City

Tehran

Province

Tehran

Postal code

19585175

Approval date

2018-10-13, 1397/07/21

Ethics committee reference number

IR.IAU.DENTAL.REC.1397.032

Health conditions studied

1

Description of health condition studied

Insufficient keratinized gingiva

ICD-10 code

K06.0

ICD-10 code description

Gingival recession

Primary outcomes

1

Description

Loe and Silness plaque index

Timepoint

At the beginning and 3 months

Method of measurement

Observation

2

Description

Loe and Silness gingival index

Timepoint

At the beginning and 3 months

Method of measurement

Observation and probing

3

Description

Keratinized gingiva width

Timepoint

At the beginning and 3 months

Method of measurement

Periodontal probe

4

Description

Attached gingiva width

Timepoint

At the beginning and 3 months

Method of measurement

Periodontal probe

5**Description**

Probing depth

Timepoint

At the beginning and 3 months

Method of measurement

Periodontal probe

6**Description**

Shrinkage percentage

Timepoint

3 months

Method of measurement

Periodontal probe

Secondary outcomes

empty

Intervention groups**1****Description**

Intervention group: In the cases, each half-jaw was randomly assigned to the test group (the simultaneous use of the acellular dermal matrix and autogenous free gingival strap). In this group, the coronal region of receptor site was covered up to marginal margin by acellular dermal matrix called Ceno Bone from the Ceno Biologics factory with a width of 5 mm and free gingival strap was made up to a width of 2 mm from the palate and placed in the apical region of the receptor site.

Category

Treatment - Surgery

2**Description**

Control group: In the cases, each half-jaw was randomly assigned to the control group (Acellular Dermal Matrix alone). In this group, the acellular dermal Matrix called Ceno Bone by the Ceno Biologics factory will be fixed at 7 mm width in the recipient area with the polyamide thread 5.0.

Category

Treatment - Surgery

Recruitment centers**1****Recruitment center****Name of recruitment center**

Faculty of Dentistry Islamic Azad University of Tehran

Full name of responsible person

Nima Nadaf pour

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Islamic Azad University

Full name of responsible person

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Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Islamic Azad University

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**

Islamic Azad University

Full name of responsible person

Nima Nadaf Pour

Position

Periodontics professor

Latest degree

Specialist

Other areas of specialty/work

Dentistry

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

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Full name of responsible person

Nima nadaf Pour

Position

Periodontics Professor

Latest degree

Specialist

Other areas of specialty/work

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Person responsible for updating data**Contact****Name of organization / entity**

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Dental student

Latest degree

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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

Undecided - It is not yet known if there will be 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

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

Clinical Study Report

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

Analytic Code

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

Data Dictionary

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

Title and more details about the data/document

Evaluation of using Acellular dermal matrix (ADM) and autogenous free gingival strap on clinical conditions of gingiva in patients referring to the dental unit of Islamic Azad University of Tehran in 1397-1398 .

When the data will become available and for how long

6 months after printing the results

To whom data/document is available

All people who need data

Under which criteria data/document could be used

Analyzes that do not damage the outcome of our data analysis are allowed.

From where data/document is obtainable

Faculty of Dentistry Islamic Azad University of Tehran

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

Visit the Faculty of Dentistry, Islamic Azad University of Tehran and request there, and will access the data within 1 week.

Comments