

Root coverage with Emdogain versus root coverage with cross-linked hyaluronic acid

Submission date 20/11/2024	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 22/11/2024	Overall study status Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 12/09/2025	Condition category 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

Gingival recession, where the gum tissue recedes and exposes the tooth root, is a common dental condition that can lead to sensitivity, root decay, and aesthetic concerns. This study aims to compare two methods for treating gingival recession when combined with a connective tissue graft (CTG), the gold standard treatment. One method uses cross-linked hyaluronic acid (HA), and the other uses enamel matrix derivative (EMD). By comparing these treatments, the study hopes to identify which approach provides better gum coverage, tissue healing, and patient satisfaction.

Who can participate?

Adults aged 18 to 70 years old with specific types of gum recession (classified as RT1-2 or Miller Class I-III) in good general health can participate.

What does the study involve?

Participants will be randomly assigned to one of two treatment groups:

Group A: Connective tissue graft (CTG) combined with cross-linked hyaluronic acid (HA)

Group B: CTG combined with enamel matrix derivative (EMD)

A minimally invasive gum surgery will be performed in which the CTG will be placed into a tunnel created beneath the gums, and either HA or EMD will be applied to the tooth root. All procedures will be performed under local anesthesia by an experienced periodontist. The study includes follow-up visits to monitor healing and results for six months.

What are the possible benefits and risks of participating?

Participants may benefit from improved gum health, reduced sensitivity, and better aesthetics as part of their treatment. Risks are minimal and include common surgical risks such as swelling, mild discomfort, or temporary bleeding, all of which are manageable and expected to subside quickly.

Where is the study run from?

The Department of Periodontology, Victor Babes University of Medicine and Pharmacy, Timisoara, Romania

When Is the Study Starting and How Long Is It Expected to Run For?

November 2024 to December 2026, with recruitment, treatment, and follow-up taking place over 24 months. Results and analysis will be completed within 27 months.

Who is funding the study?

The University Clinic of Periodontology of Victor Babes University of Medicine and Pharmacy funds,

Who Is the Main Contact?

Dr. Meizi Eliezer, meizi.eliezer@gmail.com

Contact information

Type(s)

Public, Scientific, Principal investigator

Contact name

Dr Meizi Eliezer

ORCID ID

<https://orcid.org/0000-0003-2164-1272>

Contact details

Golomb st 77 apartment 2

Herzelia

Israel

4630577

+9720522577260

meizi.eliezer@gmail.com

Type(s)

Scientific

Contact name

Prof Stefan-loan Stratul

Contact details

Str. Emanoil Gojdu 5

Timisoara

Romania

300176

+40744521470

s.stratul@gmail.com

Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Protocol serial number

Study information

Scientific Title

Root coverage with adjunctive enamel matrix derivative vs. cross-linked hyaluronic acid: 6-months, blinded RCT

Acronym

RC EMD XHYA

Study objectives

It is hypothesized that there is no significant difference in the primary outcomes of the percentage of root coverage and thickness of keratinized tissue between the crosslinked HA and EMD groups.

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 18/11/2024, The Committee on Research Ethics of the Victor Babes University of Medicine and Pharmacy Timisoara (P-ța Eftimie Murgu, nr. 2, Timișoara, 300041, Romania; +40744521470; cecs@umft.ro), ref: 61/18.11.2024

Study design

Single-blind randomized controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Gingival recession coverage treatment

Interventions

The study will include patients presenting with gingival recessions classified as RT1-2 (Cairo, 2011) or Miller Class I-III (Miller, 1985) on maxillary or mandibular teeth. A total of 60 recession defects will be enrolled and randomly allocated into two groups using block randomisation because it will ensure balanced group sizes at all stages of the trial. Group A will undergo recession coverage using a subepithelial connective tissue graft (CTG) combined with cross-linked hyaluronic acid (HA), while Group B will be treated with enamel matrix derivatives (EMD).

To ensure consistency, all surgeries will be performed under local anesthesia by a single experienced periodontist, Dr Meizi Eliezer. The surgical tunnel will be created using a lateral closed tunnel or a modified coronally advanced tunnel technique. The subepithelial CTG will be inserted into the tunnel, and either cross-linked HA or EMD will be applied to the exposed root surface to enhance regeneration and achieve optimal outcomes. Flaps will be repositioned and sutured. Patients will undergo standard post-operative care for regenerative procedures.

Sutures will be removed three weeks after the surgery, and gentle brushing of the operated areas will be resumed after three weeks. Patients will enter supportive periodontal care, depending on the gingival and periodontal status.

Intervention Type

Procedure/Surgery

Primary outcome(s)

Percentage of root coverage measured using a periodontal probe (PCP-UNC15 HuFriedy) with a consistent reference point (CEJ) measured to the gingival margin position at 6 months post-surgery

Key secondary outcome(s)

1. Clinical attachment level gain will be assessed through the following measures:

1.1. Probing depth measured using a periodontal probe (PCP-UNC15 HuFriedy) with a consistent reference point (CEJ) measured to the gingival margin position at 6 months post-surgery

1.2. Complete root coverage (CRC) measured using a clinical assessment of gingival margin position relative to the cemento-enamel junction (CEJ) with periodontal probe (PCP-UNC15 HuFriedy), then CRC is calculated as the percentage of sites where full root coverage (no visible root exposure) has been achieved relative to the total number of sites treated, at 6 months post-surgery

2. Recession depth reduction will be assessed through the following measures:

2.1. Gingival thickness measured using a periodontal probe at 6 months post-surgery

2.2. Patient's perception measured using the Patient-Reported Outcome (PRO) Visual Analog Scale (VAS) immediately post-op (within 2-4 hours), Day 1 (24 hours post-surgery), Day 3 (72 hours post-surgery), Day 7 (1 week post-surgery) and at Day 14 and 1 month for delayed pain assessment

2.3 Esthetic outcomes measured using the Root Coverage Esthetic Score (RES) at the final 6-month visit

Completion date

31/12/2026

Eligibility

Key inclusion criteria

1. Patients with RT1-2 or Miller 1-3, gingival recessions on maxillary or mandibular teeth

2. Aged between 18-70 years old

3. In good general health

4. Non-smokers or smoking up to 10 cigarettes/day

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

70 years

Sex

All

Total final enrolment

90

Key exclusion criteria

1. Systemic diseases affecting periodontal health
2. Without periodontal active disease
3. Smoking above 10 cigarettes/day
4. Pregnancy

Date of first enrolment

21/11/2024

Date of final enrolment

01/08/2025

Locations**Countries of recruitment**

Romania

Study participating centre

University Clinic of Periodontology, Victor Babes University of Medicine and Pharmacy Timisoara

Bv. Revolutiei nr.9

Timișoara

Romania

300176

Sponsor information**Organisation**

Victor Babeș University of Medicine and Pharmacy Timișoara

ROR

<https://ror.org/00afdp487>

Funder(s)

Funder type

University/education

Funder Name

Victor Babeş University of Medicine and Pharmacy

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analyzed during the current study will be available upon request from Dr. Meizi Eliezer, meizi.eliezer@gmail.com, and from Prof.Dr. Stefan-Ioan Stratul, [s.stratul@gmail.com](mailto:stratul@gmail.com)

IPD sharing plan summary

Available on request