

Clinical Trial Protocol

Iranian Registry of Clinical Trials

18 Sep 2026

Evaluation of the efficacy of topical Gabapentin for the treatment of pruritus in patients with Epidermolysis bullosa, Faghihi hospital, 2021

Protocol summary

Study aim

Evaluation of the efficacy of 10% gabapentin cream compared to base cream without Gabapentin in the treatment of pruritic lesions in patients with Epidermolysis bullosa.

Design

A non-randomized, double-blind; placebo-controlled clinical trial with blinded outcome assessment, 19 patients; convenient sampling.

Settings and conduct

A total of 19 patients with various forms of epidermolysis bullosa referred to the Faghihi Hospital will be included (Available/ convenient Sampling). The dermatologist first identifies two pruritic areas in each patient (A and B), each less than 3% of BSA, and then measures the size and erythema of each area. The lesions will be photographed and the Leuven-Itch-scale questionnaire is completed by the patients. Patients will receive 10% gabapentin cream on one side and base cream on the other (patients and dermatologists are unaware of the type of treatment on each side). Two similar containers are designed with labels A or B, and patients apply them 3 times a day for 6 weeks. Patients then return to the dermatology clinic at the end of week 6, and the rate of erythema, area size, and leuven-Itch-scale and imaging on each side are performed by a dermatologist.

Participants/Inclusion and exclusion criteria

Patients older than 6 years; different forms of epidermolysis bullosa (histologically confirmed); unresponsive to conventional antipruritic treatment, no pregnancy or lactating; no underlying systemic disease or disorders that can make them prone to itching, regardless of skin disease

Intervention groups

Two pruritic lesions are identified in each patient, then one of the lesions is treated with Gabapentin 10% cream (Prepared and formulated in the Dr. Rastegar pharmacy in Shiraz) and another lesion with a gabapentin-free base cream three times a day for 6 weeks.

Main outcome variables

Erythema, lesion size and leuven-itch-scale

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210822052258N1**

Registration date: **2021-09-15, 1400/06/24**

Registration timing: **prospective**

Last update: **2021-09-15, 1400/06/24**

Update count: **0**

Registration date

2021-09-15, 1400/06/24

Registrant information

Name

Samira Vahedi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

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Email address

svhd7988@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-09-23, 1400/07/01

Expected recruitment end date

2021-11-22, 1400/09/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the efficacy of topical Gabapentin for the treatment of pruritus in patients with Epidermolysis bullosa, Faghihi hospital, 2021

Public title

Evaluation of the efficacy of topical Gabapentin for the treatment of pruritus in patients with Epidermolysis bullosa

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

Age equal or more than 6 years; Histologically confirmed any three major forms of epidermolysis bullosa (EB simplex (EBS), junctional EB (JEB), and dystrophic EB (DEB)) complain of persistent pruritic lesions unresponsive to the standard treatment

Exclusion criteria:

Use of oral gabapentin Pregnancy or lactation Concomitant kidney, liver, blood, thyroid, and psychiatric diseases or any other systemic disease that may cause pruritus

AgeFrom **6 years** old**Gender**

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser

Sample sizeTarget sample size: **19**

More than 1 sample in each individual

Number of samples in each individual: **2**

Two pruritic areas of skin are selected in each patient, one for medication and the other for placebo

Randomization (investigator's opinion)

Not randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description

In each patient, two pruritic areas are defined for the use of medication or a placebo. The type of intervention in each lesion remains unclear to both, the patients and the researcher until the end of the study. Products are delivered within completely similar 50 gr containers. Containers are identified with adhesive labels (A or B) so the subjects and investigators are not aware of the kind of treatment in each lesion (double-blind design).

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics committee of Shiraz University of Medical Sciences

Street address

Local Ethics Committee, Shiraz University of Medical Sciences, Setad Squ., Zand Blvd., Shiraz.

City

Shiraz

Province

Fars

Postal code

7176913861

Approval date

2021-07-14, 1400/04/23

Ethics committee reference number

IR.SUMS.MED.REC.1400.210

Health conditions studied**1****Description of health condition studied**

Epidermolysis bullosa

ICD-10 code

Q81

ICD-10 code description

Epidermolysis bullosa

Primary outcomes**1****Description**

itching

Timepoint

Baseline and at the end of treatment (end of 6th week)

Method of measurement

Leuven_itch_scale

2**Description**

Pruritic area size

Timepoint

Baseline and at the end of treatment (end of 6th week)

Method of measurement $(\text{Length} \times \text{width}) \div 2$

3

Description

Erythema

Timepoint

Baseline and at the end of treatment (end of 6th week)

Method of measurement

Examination: absent (0), mild (pink-1), moderate (pink to red-2), severe (red-3)

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Intervention group receives topical Gabapentine 10%(plus 10%Glycerine in base of cold cream)formulized and prepared in Dr.Rastegar pharmacy on the assigned side 3 times daily for 6 weeks .

Category

Treatment - Drugs

2

Description

Control group: Control group uses base cream(1. % glycerin in cold cream) on the assigned side 3times daily for 6 weeks.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Faghihi hospital

Full name of responsible person

Nasrin Saki

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Dermatology clinic, Faghihi hospital, Setad Sque., Zand Blvd., Shiraz.

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shiraz University of Medical Sciences

Full name of responsible person

Dr Younes Ghasemi

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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

Shiraz 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

Shiraz University of Medical Sciences

Full name of responsible person

Samira Vahedi

Position

Dermatology resident

Latest degree

Medical doctor

Other areas of specialty/work

Dermatology

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Shiraz University of Medical Sciences

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Position

Dermatology resident

Latest degree

Medical doctor

Other areas of specialty/work

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be 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

Not applicable