

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Double blind, randomized phase 3 efficacy trial of topical nano-liposomal paromomycin in the treatment of cutaneous leishmaniasis caused by *L. major*

Protocol summary

Summary

The aim of this clinical trial is to assess the efficacy of topical nano-liposomal 10 % paromomycin (Lip-PM) in the treatment of cutaneous leishmaniasis (CL) caused by *L. major* compared with placebo. Inclusion criteria are parasitologically proven cases of CL based on positive smear and/or culture, age 12-60 years, duration of lesion less than 3 months, number of lesions at most 5, lesion size not more than 30 mm in one direction, be volunteer to participate and sign informed consent, exclusion criteria are pregnant or lactating women and duration of lesion more than 3 months, participation in any other trial in last 6 months. A total of 144 ZCL patients will be recruited and double blind randomly assigned in 2 arms, 72 patients in each arm. The patients will be double blind randomly receiving either topical Lip-PM or placebo twice daily for 4 weeks (28 days), the study will be completed according to GCP (Good Clinical Practice) guide lines. The patients will be visited at week 4, 8 and at 3 and 6 months after initiation of treatment. At each visit the lesions' characteristics such as size of induration, ulcer and scar will be recorded. The data will be compared between the patients who received the treatment (Lip-PM) and the group who received placebo. The follow up of the patients is for 6 months after beginning of the treatment. The primary clinical efficacy parameter is complete re-epithelization of all lesions with disappearance of induration (with or without scar).

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201108301475N6**

Registration date: **2011-10-05, 1390/07/13**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2011-10-05, 1390/07/13

Registrant information

Name

Ali Khamesipour

Name of organization / entity

Center for Research and Training in Skin Diseases and Leprosy, Tehran University of Medical Sciences

Country

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+98 21 8897 0657

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Recruitment status

Recruitment complete

Funding source

Zoonoses control Center for diseases control Ministry of Health and..; Center for Research & Training in Skin Diseases & Leprosy, Tehran University of Medical Sciences; Nanotechnology Research Center Mashhad University of Medical Sciences

Expected recruitment start date

2011-09-01, 1390/06/10

Expected recruitment end date

2012-03-01, 1390/12/11

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Double blind, randomized phase 3 efficacy trial of topical

nano-liposomal paromomycin in the treatment of cutaneous leishmaniasis caused by *L. major*

Public title

Double blind, randomized phase 3 efficacy trial of topical nano-liposomal paromomycin in the treatment of cutaneous leishmaniasis caused by *L. major*

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Age 12-60 years; Cutaneous leishmaniasis (CL) lesion induced by *L. major*; Parasitological proven cases of CL by direct smear, culture or PCR; Healthy individual otherwise of CL; onset less than 3 months; number of lesion 5 at most; ulcer size not more than 30 mm in one size; willing to participate and sign informed consent. Exclusion criteria: pregnant or lactating women; duration of lesion more than 3 months; serious illness such as heart kidney or liver diseases based on physical exam and laboratory tests; CL lesion resistant to treatment or recidivans form of lesion; number of lesions more than 5; ulcer size more than 30 mm in one direction participation in any other clinical trial in last 6 months.

Age

From **12 years** old to **60 years** old

Gender

Both

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **144**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

University Ethical Committee, Center for Research and Training in Skin Diseases and Leprosy, Tehran

Street address

415 Taleghani Avenue

City

Tehran

Postal code

1416613675

Approval date

2011-05-14, 1390/02/24

Ethics committee reference number

J/423/201

Health conditions studied

1

Description of health condition studied

Cutaneous leishmaniasis otherwise Healthy

ICD-10 code

B55.1

ICD-10 code description

minor skin disease

Primary outcomes

1

Description

Complete cure

Timepoint

at weeks 4 and 8 and at months 3 and 6

Method of measurement

Measurement of reepithelization

Secondary outcomes

1

Description

Partial cure

Timepoint

at weeks 4 and 8 and at months 3 and 6

Method of measurement

Measurement of reepithelization

Intervention groups

1

Description

Treatment with topical Nano liposomal form of 10% paromomycin, twice a day for 28 days

Category

Treatment - Drugs

2

Description

Placebo topically is used twice a day for 28 days

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Isfahan Health Center, Isfahan University of Medical Sciences

Full name of responsible person

Dr Fariba Farid

Street address

Isfahan University of Medical Sciences, Vice Chancellor for Health

City

Isfahan

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Center for Research & Training in Skin Diseases & Leprosy Tehran University of Medical Sciences; Dis

Full name of responsible person

Dr Alireza Firooz, Dr Gooya or Shirzadi, Dr Jaafari

Street address

415 Taleghani Avenue, Hafez Avaneue, BooAli Sq

City

Tehran, Tehran, Mashad

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Center for Research & Training in Skin Diseases & Leprosy Tehran University of Medical Sciences; Dis

Proportion provided by this source

100

Public or private sector

empty

Domestic or foreign origin

empty

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

empty

Person responsible for general inquiries

Contact

Name of organization / entity

Center for Research and Training in Skin Diseases and Leprosy, Tehran University of Medical Sciences

Full name of responsible person

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Position

Prof/PhD

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

empty

Study Protocol

empty

Statistical Analysis Plan

empty

Informed Consent Form

empty

Clinical Study Report

empty

Analytic Code

empty

Data Dictionary

empty