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Effect of topical honey application along with intralesional injection of glucantime in the treatment of the cutaneous leishmaniasis

Submission date	Recruitment status	[_] Prospectively registered
10/09/2006	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
13/11/2006	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
23/01/2020	Infections and Infestations	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

84210

Study information

Scientific Title
Effect of topical honey application along with intralesional injection of glucantime in the treatment of the cutaneous leishmaniasis

Study objectives
Topical honey along with intralesional glucantime is more effective in the treatment of the cutaneous leishmaniasis than intralesional glucantime alone.

Ethics approval required
Old ethics approval format

Ethics approval(s)
The study was approved by the ethics committee of the Skin Disease and Leishmaniasis Research Center (reference number: SEC. 84210).

Study design
This study was a controlled randomised clinical trial study

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Cutaneous leishmaniasis

Interventions
Topical honey along with intralesional glucantime versus intralesional glucantime alone.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Topical honey and intralesional glucantime

Primary outcome measure
Curing of the leishmaiasis

Secondary outcome measures
Diameter of the lesion and size of the erythema, induration and ulcer

Overall study start date
21/12/2004

Completion date
22/10/2005

Eligibility

Key inclusion criteria
1. Confirmed cutaneous leishmaniasis with direct smear
2. No history of systemic or topical therapy for cutaneous leishmaniasis
3. Absence of the malnutrition or severe predisposing disease such as cardiac, renal or hepatic disease and other contraindication for glucantime

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
100

Key exclusion criteria
N/A

Date of first enrolment
21/12/2004

Date of final enrolment
22/10/2005

Locations

Countries of recruitment
Iran

Study participating centre

Skin Disease and Leishmaniasis Research Center (SDLRC)
Isfahan
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81876-98191

Sponsor information

Organisation
Skin Disease and Leishmaniasis Research Center (Iran)

Sponsor details
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Sponsor type
Research organisation

ROR
https://ror.org/04waqzz56

Funder(s)

Funder type
University/education

Funder Name
Isfahan University of Medical Sciences, Skin Diseases and Leishmaiasis Research Center (Iran)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results:	27/04/2007		Yes	No