

Clinical Trial Protocol

Iranian Registry of Clinical Trials

06 Jun 2026

Investigation of the efficacy of a topical dosage form consisting lawsonia inermis extracts in reducing Fluoropyrimidines induced Hand-Foot Syndrome in cancerous patients

Protocol summary

Summary

The purpose of this study is to introduce a suitable topical product of lawsonia inermis to improve Hand-foot syndrome symptoms in cancer patients who are taking Fluoropyrimidine drugs. The study population is cancer patients who are taking IV injection of 5-fluorouracil or oral Capecitabine and showing hand-foot syndrome. These patients are entered in study with consent. Patients in the event of the death or absence of these conditions, as well as allergy to formulation are excluded. The clinical trial of this formulation will be conducted on 20 cancer patients with Hands and foot syndrome of one medical center in Tehran , in a double-blind, placebo-controlled and parallel study. Trial group is combination of phase 1 and 2. The sample is consisted of 40 pairs of hands and feet. Patients are randomly divided into two groups, pink and green; pink group receive the placebo gel to their hands and the drug gel to their feet; the green contrary receive the placebo gel to their feet and the drug gel to their hands for two weeks. The beginning and the fourteenth day of study, the severity of lesions are checked, photographed and questionnaire is completed; syndrome is graded according to data by the National Cancer Institute protocols and changes are investigated.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014051017643N1**
Registration date: **2015-08-27, 1394/06/05**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2015-08-27, 1394/06/05

Registrant information

Name

Farhad Shahi

Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Pharmaceutical Sciences branch, Islamic Azad University

Expected recruitment start date

2014-06-11, 1393/03/21

Expected recruitment end date

2016-01-05, 1394/10/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigation of the efficacy of a topical dosage form consisting lawsonia inermis extracts in reducing Fluoropyrimidines induced Hand-Foot Syndrome in cancerous patients

Public title

Investigation of the efficacy of a herbal topical gel in reducing the skin reactions in cancer patients taking chemotherapy drugs Fluoropyrimidines induced Hand-

Foot Syndrome in cancerous patients

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria: Taking 5-fluorouracil (5-fu) by injection or orally Capecitabine as an anticancer drug; showing hand-foot syndrome; patient satisfaction; Having cancer. Exclusion criteria: Treatment change and discontinuing 5-fluorouracil (5-fu) and oral Capecitabine administration; Showing no hand-foot syndrome; Patient's death; Lack of cancer; Sensitivity to compounds of formula.

Age

No age limit

Gender

Both

Phase

1-2

Groups that have been masked

No information

Sample size

Target sample size: 40

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

Islamic Azad University Of Pharmaceutical Sciences
Branch

Street address

Iakhchal Avenue, Qolhak, Dotor Shariati Avenue,
Tehran

City

Tehran

Postal code

19395-6466

Approval date

2013-12-03, 1392/09/12

Ethics committee reference number

11994

Health conditions studied

1

Description of health condition studied

hand-foot syndrome

ICD-10 code

L27.1

ICD-10 code description

localized skin eruption due to drugs and medicaments

Primary outcomes

1

Description

severity of HFS symptoms

Timepoint

Beginning of the study , the fourteenth day

Method of measurement

NCI Hand-Foot Syndrome staging version 4.0

Secondary outcomes

1

Description

Skin Sensitivity

Timepoint

From the beginning of the study

Method of measurement

Observational

Intervention groups

1

Description

The topical gel will be administrated to the foot or hand of the patients randomly four times a day for two weeks

Category

Treatment - Drugs

2

Description

The topical placebo will be administrated to the foot or hand of the patients taking the topical drug randomly four times a day for two weeks

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Imam Khomeini Hospital of Tehran

Full name of responsible person

Dr.Farhad Shahi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Pharmaceutical Sciences branch, Islamic Azad University(a part of razieh mohajerani's PharmD thesis

Full name of responsible person

Dr.Sepide Arbabi

Street address

Islamic Azad University Of Pharmaceutical Sciences Branch, Iakhchal avenue, Dotor Shariati avenue, Tehran

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Tehran

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Pharmaceutical Sciences branch, Islamic Azad University(a part of razieh mohajerani's PharmD thesis

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

Islamic Azad University Of Pharmaceutical Sciences Branch

Full name of responsible person

Razieh Mohajerani

Position

student of pharmacy

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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