

Clinical Trial Protocol

Iranian Registry of Clinical Trials

25 Jul 2026

In vivo bioequivalence study if Amantadine 100 mg capsule manufactured by Fatak shimi Co. in comparison to innovator product Symmetetrel

Protocol summary

Study aim

Investigating In vivo bioequivalence study of Amantadine 100 mg capsule manufactured by Fatak shimi Co. in comparison to innovator product

Design

Twenty-four healthy male volunteer will enter the study based on random numbers table as two groups of twelve people. Each volunteer will receive a single dose of drug in two periods. In one period the test formulation and in another period the reference formulation (without knowing the formulation identity in each period). Therefore, each volunteer will be his own "Control".

Settings and conduct

After administration of one 100 mg capsule to volunteer, the blood samples will be taken in predetermined time intervals up to 72 hours. The samples will be stored in freezer -4 degrees centigrade until analysis and sample quantitation. In this study the volunteer would not be aware of the formulation identity in each period

Participants/Inclusion and exclusion criteria

Inclusion criteria: general health (Liver, Heart and Kidney); age: 20-60 years old. Exclusion criteria: smoking; history of cardiovascular disease, liver and kidney disease; pregnancy; alcohol and drug addiction; history of drug allergy.

Intervention groups

Test group: administrating the test product. Control group: administrating the innovator product.

Main outcome variables

Drug plasma concentration

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210519051345N7**

Registration date: **2021-11-27, 1400/09/06**

Registration timing: **registered_while_recruiting**

Last update: **2021-11-27, 1400/09/06**

Update count: **0**

Registration date

2021-11-27, 1400/09/06

Registrant information

Name

Parvin Zakeri-Milani

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 41 3334 8801

Email address

pzakeri@tbzmed.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-11-21, 1400/08/30

Expected recruitment end date

2021-12-26, 1400/10/05

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

In vivo bioequivalence study if Amantadine 100 mg capsule manufactured by Fatak shimi Co. in comparison to innovator product Symmetetrel

Public title

Bioequivalence study of Amantadine 100 mg capsule

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

General Health (Liver, Heart and Kidney), Age (20-60)

Exclusion criteria:
Smoking, History of cardiovascular disease, liver and kidney disease, Pregnancy, Alcohol and drug addiction, History of drug allergy

Age
From **20 years** old to **60 years** old

Gender
Both

Phase
Bioequivalence

Groups that have been masked

- Participant

Sample size
Target sample size: **24**

Randomization (investigator's opinion)
Randomized

Randomization description
"Random number table" will be used. Each volunteer will be assigned a two-digit number. After making the table, we need randomly point to a spot on the table and select the numbers.

Blinding (investigator's opinion)
Single blinded

Blinding description
In this study, Volunteers participating will be blinded to the type of product they are taking in each period (test or reference product). This means that the product will be given to the volunteers for administration outside the original packaging

Placebo
Not used

Assignment
Crossover

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee
Biomedical Research Committee, Tabriz University of Medical Sciences

Street address
Golgashat st Attar Neishaboori st, Faculty of Pharmacy, Tabriz University of Medical Sciences

City
Tabriz

Province
East Azarbaijan

Postal code
51664-14766

Approval date
2021-11-15, 1400/08/24

Ethics committee reference number

IR.TBZMED.REC.1400.742

Health conditions studied

1

Description of health condition studied

In the present study, no diseases will be examined and products will be administered by healthy volunteers

ICD-10 code

ICD-10 code description

Primary outcomes

1

Description

Plasma Drug Concentration

Timepoint

0.5-72 hours in predetermined time intervals after drug administration

Method of measurement

HPLC

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group will receive one test drug product. Blood samples will be taken from the volunteers for 72 hours at the mentioned times after drug administration and the concentration of drug in blood samples will be measured by liquid chromatography with mass spectroscopy detector.

Category

Treatment - Drugs

2

Description

Control group will receive one reference drug product. Blood samples will be taken from the volunteers for 72 hours at the mentioned times after drug administration and the concentration of drug in blood samples will be measured by liquid chromatography with mass spectroscopy detector.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Faculty of Pharmacy, Tabriz University of Medical Sciences

Full name of responsible person

Parvin Zakeri-Milani

Street address

Golgasht st Attar Neishaboori st, Faculty of Pharmacy,
Tabriz University of Medical Sciences

City

Tabriz

Province

East Azarbaijan

Postal code

51664-14766

Phone

+98 41 3334 8801

Email

pzakeri@tbzmed.ac.ir

Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Tabriz University of Medical Sciences

Full name of responsible person

Mohammad Sameie

Street address

No.2 Central Building 3rd Floor, Daneshgah st. Tabriz
University of Medical Sciences

City

Tabriz

Province

East Azarbaijan

Postal code

51664-14766

Phone

+98 41 3334 8801

Email

Samiei.moh@gmail.com

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

No

Title of funding source

Fatak Shimi Pars Co.

Proportion provided by this source

100

Public or private sector

Private

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin**Type of organization providing the funding**

Industry

Person responsible for general inquiries

Contact**Name of organization / entity**

Tabriz University of Medical Sciences

Full name of responsible person

Parvin Zakeri-Milani

Position

Professor

Latest degree

Ph.D.

Other areas of specialty/work

Medical Pharmacy

Street address

Golgasht st Attar Neishaboori st, Faculty of Pharmacy,
Tabriz University of Medical Sciences

City

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Province

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Person responsible for scientific inquiries

Contact**Name of organization / entity**

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Person responsible for updating data

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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

Undecided - It is not yet known if there will be a plan to
make this available