

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jul 2026

Efficacy and Safety Study of Injection Iron Sucrose in Pregnant Women with Iron Deficiency Anaemia

Protocol summary

Summary

Objective: To evaluate safety and efficacy of Test Product; Megafer Injection® in Pregnant women with iron deficiency Anemia. Inclusion Criteria: Pregnant Women in second or third trimester of pregnancy, Hb: $\leq$ 8 g/dl and serum ferritin levels are below than normal Exclusion Criteria: Patient allergic to iron preparation or any other ingredient of Injection megafer . Study Population: Pregnant Women Sample Size: 50 subjects Intervention: Megafer Injection® (Iron Sucrose 100mg/5ml), Main Outcomes measures: Increase in Hb level at 2-4 weeks after initiation of treatment, achievement of target haemoglobin.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201611037978N4**
Registration date: **2016-11-18, 1395/08/28**
Registration timing: **retrospective**

Last update:
Update count: **0**

Registration date

2016-11-18, 1395/08/28

Registrant information

Name

Tasneem Ahmad

Name of organization / entity

Pharma Professional Services, Karachi, Pakistan

Country

Pakistan

Phone

(92-21) 34972358, 34820573

Email address

tasneem.ahmed@iccs.edu

Recruitment status

Recruitment complete

Funding source

Surge Laboratories (Private) Ltd.

Expected recruitment start date

2015-07-31, 1394/05/09

Expected recruitment end date

2016-01-18, 1394/10/28

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Efficacy and Safety Study of Injection Iron Sucrose in Pregnant Women with Iron Deficiency Anaemia

Public title

Efficacy and Safety Study of Megafer injection® in Pregnant Women

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion Criteria: Pregnant Women in second or third trimester, Hb: $\leq$ 8 g/dl and serum ferritin levels are below than normal Exclusion Criteria: Patient allergic to iron preparation or any other ingredient of Injection megafer.

Age

From **18 years** old to **40 years** old

Gender

Female

Phase

4

Groups that have been masked

No information

Sample size

Target sample size: **50**

Randomization (investigator's opinion)

N/A

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

Not blinded

Blinding description**Placebo**

Not used

Assignment

Single

Other design features

Single arm, Open label study.

Secondary Ids

empty

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

Ethics Committee of Jinnah Post Graduate Medical Centre

Street address

Rafiqi Shaheed Road, Jinnah Post Graduate Medical Centre

City

karachi

Postal code

75510

Approval date

2015-02-14, 1393/11/25

Ethics committee reference number

NO.F.2-81/2015-GENL/11128/JPMC

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

Iron Deficiency anemia

ICD-10 code

D-50

ICD-10 code description

Iron Deficiency anemia, unspecified

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

Increase in Hb level after initiation of treatment, achievement of target haemoglobin.

Timepoint

2 & 4 weeks after first dose

Method of measurement

Haemoglobin testing

Secondary outcomes**1****Description**

Reported adverse events related to drug only.

Timepoint

At every follow up visit

Method of measurement

Patient monitoring & Follow up calls to patients

Intervention groups**1****Description**

The total calculated dose was divided as 100 to 200 mg dose/day maximum once or twice in week. Firstly, 25mg (25ml) was infused over 30 minutes @12 drops /minute as a test dose and the remaining dose was given at the rate 30-36 drops /minute, aiming the target haemoglobin of 11g/dl.

Category

Treatment - Drugs

Recruitment centers**1****Recruitment center****Name of recruitment center**

Jinnah Postgraduate Medical Centre (JPMC)

Full name of responsible person

Dr. Haleema Yasmeen

Street address

Department of Gynecology, Jinnah Postgraduate Medical Centre (JPMC)

City

karachi

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Surge Laboratories Pvt. Ltd.

Full name of responsible person

Mohammad Younus Batla

Street address

509-510, 5th Floor, Commerce Centre, Hasrat Mohani Road, Karachi, Pakistan

City

Karachi

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

Yes

Title of funding source

Surge Laboratories Pvt. Ltd.

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

City

karachi

Province

Sindh

Postal code

75510

Phone

00922199201300

Fax**Email**

dr.haleemayasmin@yahoo.com

Web page address**Person responsible for general inquiries****Contact****Name of organization / entity**

Nabiqasim Industries (Pvt) Ltd.

Full name of responsible person

Salman Rahim

Position

Assistant Manager Regulatory Affairs.

Other areas of specialty/work**Street address**

510, 5th floor, commerce center, Hasrat Mohani Road

City

Karachi

Province

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salman.rahim@nabiqasim.com

Web page address**Person responsible for updating data****Contact****Name of organization / entity**

Pharma Professional Services

Full name of responsible person

Prof.Dr. Tasneem Ahmad

Position

Chief Investigator

Other areas of specialty/work**Street address**

A-93, Ettawah Society, scheme 33

City

karachi

Province

Sindh

Postal code

75340

Phone

00

Fax**Email****Web page address****Person responsible for scientific inquiries****Contact****Name of organization / entity**

Department of Gynaecology, Jinnah Post Graduate Medical Centre

Full name of responsible person

Dr. Haleema Yasmeen

Position

Assistant Professor

Other areas of specialty/work**Street address**

Rafiqi Shaheed Road, Jinnah Post Graduate Medical Centre

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