

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

A Randomized, Multi-center, Single Dose, Cross over Study Comparing the Pharmacokinetic of Bio-similar EPTACOG Alfa with NOVOSEVEN®, in Patients with Congenital Factor VII Deficiency

Protocol summary

Study aim

Evaluation of pharmacokinetic(PK) of biosimilar eptacog alfa(AryoSeven)and NovoSeven in phase A(PK)and monitoring inhibiting antibody formation,lack of efficacy,collection of safety data in the following 12 months(Phase B)

Design

A phase III multicenter double-blind cross-over study with 24 sample size who based on 1:1 randomization receive either single dose of AryoSeven and NovoSeven 30 mcg per kg separated by 3days washout period.Patients hospitalized at time of medication administration and plasma sampling.Plasma sampling atleast 5ml at 10min before,10,20min,1,3,6,8,12,24,30h post drug administration.Phase B:At end of PK phase all patients receive AryoSeven on demand for every bleeding episode that occurs during 12 months or prophylaxis.The treatment modality(on demand at study center or home,or prophylaxis)decided by Investigator.Sampling for determination of antibody taken every 3 months.

Settings and conduct

Multicenter Iran and Turkey.First phase 1 week with infusion of single dose aryoseven and novoseven randomly.Second phase 1year receiving aryoseven for all bleedings.Blinding by an independent third party nurse or pharmacist unblinded who prepares undistinguishable syringes with patient dosing and labelling

Participants/Inclusion and exclusion criteria

Inclusion:Adult and children more than12 years with confirmed diagnosis of severe Factor VII Deficiency(less than 1%)with more than 2 episodes of bleeding per year requiring treatment with FVII infusions Exclusion:inhibitor against FVII is positive at time of study entry.Active hepatitis and liver cirrhosis

Intervention groups

Intervention group are congenital FVII deficient patients who receive AryoSeven 30 mcg per kg Control group are

congenital FVII deficient patients who receive NovoSeven 30 mcg per kg

Main outcome variables

similar pharmacokinetic profile

General information

Reason for update

Trial completion date is recorded wrong and needs correction.

Acronym

IRCT registration information

IRCT registration number: **IRCT2016120231193N1**

Registration date: **2016-12-25, 1395/10/05**

Registration timing: **prospective**

Last update: **2022-02-27, 1400/12/08**

Update count: **2**

Registration date

2016-12-25, 1395/10/05

Registrant information

Name

Amirhossein Saadatirad

Name of organization / entity

AryoGen Pharmed

Country

Iran (Islamic Republic of)

Phone

+98 26 3610 6480

Email address

saadatirada@aryogen.com

Recruitment status

Recruitment complete

Funding source

Aryogen Pharmed Company

Expected recruitment start date

2016-12-21, 1395/10/01
Expected recruitment end date
2017-09-23, 1396/07/01
Actual recruitment start date
2017-06-24, 1396/04/03
Actual recruitment end date
2018-09-01, 1397/06/10
Trial completion date
2019-09-14, 1398/06/23

Scientific title

A Randomized, Multi-center, Single Dose, Cross over Study Comparing the Pharmacokinetic of Bio-similar EPTACOG Alfa with NOVOSEVEN®, in Patients with Congenital Factor VII Deficiency

Public title

The pharmacokinetic study of AryoSeven with Novoseven in Patients with Congenital Factor VII Deficiency

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Patients with a confirmed diagnosis of congenital, severe Factor VII deficiency (FVII <1%), with >2 episodes of bleeding/year requiring treatment with FVII infusions, in non-bleeding status. Male and female subjects Adult and children (>12 years) Patients to be enrolled must also provide voluntary written informed consent to the protocol to be eligible for the study. For minor patients, parent/legal guardian will provide consent and, when possible, patient assent will also be obtained. For compromised patients, their designated proxy must provide informed consent. Patients in the Pharmacokinetic (PK) phase will be hospitalized at time of study medication administration and plasma sampling (2 times during the study).

Exclusion criteria:

Any other type of congenital or acquired coagulopathy (except congenital Factor VII deficiency), such as: liver disease (hepatitis), vitamin k deficiency, uremia, malignancy. Antibodies against Factor VII Patients entering the PK Phase who have not suspended prophylactic regime with Novoseven or AryoSeven 3 days before starting the trial (receiving first dose of study medication). Platelet count less than 100.000 platelets/mcL (at screening visit) Patients who have received routine (prophylactic) treatment with rFVIIa in the period between screening visit (visit 1) and visit 2 of this study (first dose administration) Any clinical sign or known history of arterial thrombotic event or deep venous- thrombosis or pulmonary embolism HIV positive with current CD4+ count of less than 200/μL Liver cirrhosis Known hypersensitivity to the study medication Parallel participation in another experimental drug trial. Parallel participation in another marketed drug trial that may affect the primary end point of the study.

Age

From **12 years** old to **99 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data and Safety Monitoring Board

Sample size

Target sample size: **24**

Actual sample size reached: **24**

Randomization (investigator's opinion)

Randomized

Randomization description

The eligible patients are administered randomly in a 1:1 manner with either AryoSeven or NovoSeven. Allocation of treatment in this study is performed randomly using block randomization. Randomization sequence is prepared using R-CAN software version 4.0.1 in a blocks of 2 or 4 for 24 patients. Randomization list is prepared by an independent statisticians based on a balanced Latin Square design and the code is allocated to a patient by a third party in the study site based on the order of enrollment.

Blinding (investigator's opinion)

Double blinded

Blinding description

Blinding is performed by an independent third party operator (nurse/pharmacist, unblinded), who will prepared undistinguishable syringes with patient's dosing and labelling.

Placebo

Not used

Assignment

Crossover

Other design features

Secondary Ids

1

Registry name

clinicaltrials.gov

Secondary trial Id

NCT03079063

Registration date

2017-03-14, 1395/12/24

Ethics committees

1

Ethics committee

Name of ethics committee

Iran University of Medical science

Street address

Iran University of Medical Sciences ,Shahid Hemmat Highway

City

Tehran

Province

Tehran

Postal code

1449614535

Approval date

2016-09-25, 1395/07/04

Ethics committee reference number

IR.IUMS.REC.1395.28155

2**Ethics committee****Name of ethics committee**

Added at 2017-01-07: Shiraz University of Medical Sciences

Street address

Added at 2017-01-07: Shiraz University of Medical Sciences

City

Added at 2017-01-07: Shiraz

Province

Fars

Postal code

Added at 2017-01-07:

Approval date

2016-12-14, 1395/09/24

Ethics committee reference number

IR.SUMS.REC.1395.158

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

Hereditary deficiency of factor VII

ICD-10 code

D68.2

ICD-10 code description

Hereditary deficiency of other clotting factors

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

PK-parameters: the area under the plasma activity-time curve from time 0 to last quantifiable activity (AUClast)

Timepoint

10 min before drug administration, 10 min, 20 min, 1 h, 3 h, 6 h, 8 h and 12 h and 24 h after AryoSeven or NovoSeven injection.

Method of measurement

Pharmacokinetic assessment by measurement of plasma level of factor VII clotting activity (FVII:C) determined by commercial Staclot® VIIa-recombinant tissue factor assay (Diagnostica Stago, Asnières sur Seine, France), performed by a central lab blinded to the patient's treatment.

2**Description**

PK-parameters: maximum plasma activity Cmax

Timepoint

10 min before drug administration, 10 min, 20 min, 1 h, 3 h, 6 h, 8 h and 12 h and 24 h after AryoSeven or NovoSeven injection.

Method of measurement

Pharmacokinetic assessment by measurement of plasma level of factor VII clotting activity (FVII:C) determined by commercial Staclot® VIIa-recombinant tissue factor assay (Diagnostica Stago, Asnières sur Seine, France), performed by a central lab blinded to the patient's treatment.

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

Secondary Pharmacokinetic parameters: AUCinf, Vd, Thalf, Tmax, Clearance, Mean Residence Time, λz.

Timepoint

For secondary PK parameters: 10 min before drug administration, 10 min, 20 min, 1 h, 3 h, 6 h, 8 h and 12 h and 24 h after AryoSeven or NovoSeven injection.

Method of measurement

PK Parameters: Measurement of plasma level of factor VII clotting activity (FVII:C) performed by a central lab.

2**Description**

Immunogenicity assessment.

Timepoint

At screening visit, after the second dose/second drug administration (visit 3) and then every 3 months for a year.

Method of measurement

Immunogenicity by PT-based Bethesda assay by local lab and confirmatory test by the modified Nijmegen method of the Bethesda assay by central lab

3**Description**

Clinical response in treatment of bleeding

Timepoint

2h, 6h and 12 h after last dose of Aryoseven injection at every bleeding

Method of measurement

4 point scale (Excellent, Good, Moderate, None) by investigator

4**Description**

Adverse events

Timepoint

at any time during the study

Method of measurement

Adverse events grading for severity, seriousness, expected or unexpected, relationship to the study drug, action taken, outcome.

Intervention groups

1

Description

Biosimilar eptacog alfa (AryoSeven) product of AryoGen Pharmed, intravenous, single dose of 30 mcg per kg

Category

Treatment - Drugs

2

Description

Novo Nordisk eptacog alfa (Novoseven), intravenous, single dose of 30 mcg per kg

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Iran Comprehensive Hemophilia Care Center

Full name of responsible person

Dr. Mohammadreza Baghaipour

Street address

No. 3, Cross Zartosht and Felestin St, Tehran, Iran

City

Tehran

Province

Tehran

Postal code

1415863675

Phone

+98 21 8891 0001

Email

mrbaghaipour@yahoo.com

2

Recruitment center

Name of recruitment center

Added at 2017-01-07: Hematology research center

Full name of responsible person

Added at 2017-01-07: Dr. Mohammad Reza Bordbar

Street address

Added at 2017-01-07: Hematology research center, 6th floor, Mohammad Rasoulallah research building, Khalili street, Shiraz, Iran

City

Added at 2017-01-07: Shiraz

Province

Fars

Postal code

۷۱۴۵۶۸۳۷۶۹

Phone

+98 71 3228 9308

Email

bordbarm@sums.ac.ir

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

AryoGen Pharmed

Full name of responsible person

Amrhossein Saadatirad

Street address

No 140, Corner of Tajbakhsh street, 24th Km of Tehran Karaj Makhsous road

City

Garmdareh

Province

Alborz

Postal code

3164819712

Phone

+98 26 3610 6480

Email

saadatirada@aryogen.com

Grant name

Grant code / Reference number

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

Yes

Title of funding source

AryoGen Pharmed

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

AryoGen Pharmed

Full name of responsible person

Amirhossein Saadatirad

Position

Project Manager of rFVIIa registration in Europe

Latest degree

Ph.D.

Other areas of specialty/work

Medical Pharmacy

Street address

No 140, corner of Tajbakhsh street, 24th Km of Tehran Karaj Makhsous road

City

Garmdareh

Province

Alborz

Postal code

3164819712

Phone

+98 26 3610 6480

Fax

Email

saadatirada@aryogen.com

Web page address

Person responsible for scientific inquiries

Contact

Name of organization / entity

Baqiyatallah Hospital

Full name of responsible person

Dr. Hasan Abolghasemi

Position

Full professor

Latest degree

Subspecialist

Other areas of specialty/work

Pediatrics

Street address

Baqiatallah hospital clinic, Mollasadra st, south
Sheykh Bahaii st, Tehran, Iran

City

Tehran

Province

Tehran

Postal code

1435915371

Phone

0098218882742

Fax

Email

Abolghasem@bmsu.ac.ir

Web page address

Person responsible for updating data

Contact

Name of organization / entity

AryoGen Pharmed

Full name of responsible person

Amirhossein Saadatirad

Position

Project Manager of rFVIIa registration in Europe

Latest degree

Ph.D.

Other areas of specialty/work

Medical Pharmacy

Street address

No 140, corner of Tajbakhsh street, 24th Km of
Tehran Karaj Makhsoos road

City

Garmdareh

Province

Alborz

Postal code

3164819712

Phone

+98 26 3610 6480

Fax

Email

saadatirada@aryogen.com

Web page address

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