



Study of Two Doses of Crizanlizumab Versus Placebo in Adolescent and Adult Sickle Cell Disease Patients (STAND)

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Main Information

Primary registry identifying number

LBCTR2019060244

Protocol number

CSEG101A2301

MOH registration number

24334/2019

Study registered at the country of origin

Yes

Study registered at the country of origin: Specify**Type of registration**

Prospective

Type of registration: Justify

N/A

Date of registration in national regulatory agency**Primary sponsor**

Novartis Pharma Services Inc.

Primary sponsor: Country of origin

Novartis Pharmaceuticals

Date of registration in primary registry

25/04/2021

Date of registration in national regulatory agency**Public title**

Study of Two Doses of Crizanlizumab Versus Placebo in Adolescent and Adult Sickle Cell Disease Patients (STAND)

Acronym**Scientific title**

A Phase III, Multicenter, Randomized, Double-blind Study to Assess Efficacy and Safety of Two Doses of Crizanlizumab Versus Placebo, With or Without Hydroxyurea/ Hydroxycarbamide Therapy, in Adolescent and Adult Sickle Cell Disease Patients With Vaso-Occlusive Crises (STAND)

Acronym**Brief summary of the study: English**

The purpose of this study is to compare the efficacy and safety of 2 doses of crizanlizumab (5.0 mg/kg and 7.5 mg/kg) versus placebo in adolescent and adult sickle cell disease (SCD) patients with history of vaso-occlusive crisis (VOC) leading to healthcare visit.

Brief summary of the study: Arabic

دراسة مرحلة ثالثة، متعددة المراكز، عشوائية التوزيع ومزدوجة التعمية لتقييم فعالية وسلامة جرعتين من دواء كريزانليزوماب مقابل الدواء الوهمي، مع أو بدون علاج هيدروكسي يوريا / هيدروكسي كارباميد، لدى المرضى المراهقين والبالغين المصابين ببدء الكريات المنجلية مع نوبات انسداد وعائي (STAND)

Health conditions/problem studied: Specify

Sickle Cell Disease

Interventions: Specify

•Drug: Crizanlizumab (SEG101)

Crizanlizumab will be supplied in single use 10 mL glass vials at a concentration of 10 mg/mL. One vial contains 100 mg of crizanlizumab. This is a concentrate for solution for infusion IV.

Other Name: SEG101





•Drug: Placebo

Placebo will be supplied in single use 10 mL glass vials at a concentration of 10 mg/mL. One vial contains 100 mg of placebo. This is a concentrate for solution for infusion IV.

Key inclusion and exclusion criteria: Inclusion criteria

1. Written informed consent must be obtained prior to any screening procedures
2. Male or female patients aged 12 years and older on the day of signing informed consent. Adolescent include patients aged 12 to 17 years old and adults ≥ 18 years and older
3. Confirmed diagnosis of SCD by hemoglobin electrophoresis or high performance liquid chromatography (HPLC) [performed locally]. All SCD genotypes are eligible, genotyping is not required for study entry
4. Experienced at least 2 VOCs leading to healthcare visit within the 12 months prior to screening visit as determined by medical history. Prior VOC leading to healthcare visit must include:
 - a. Pain crisis defined as an acute onset of pain for which there is no other medically determined explanation other than vaso- occlusion -
 - b. a visit to a medical facility and/or healthcare professional,
 - c. and receipt of oral/parenteral opioids or parenteral nonsteroidal anti-inflammatory drug (NSAID) analgesia As well as other complicated crises, such as acute chest syndrome, priapism, and hepatic or splenic sequestration
5. If receiving HU/HC or erythropoietin stimulating agent or L-glutamine, must have been receiving the drug for at least 6 months prior to Screening visit and plan to continue taking at the same dose and schedule until the subject has reached one year of study treatment
6. Patients must meet the following central laboratory values at the screening visit:
 - Absolute Neutrophil Count $\geq 1.0 \times 10^9/L$
 - Platelet count $\geq 75 \times 10^9/L$
 - Hemoglobin: for adults (Hb) ≥ 4.0 g/dL and for adolescents (Hb) ≥ 5.5 g/dL
 - Glomerular filtration rate ≥ 45 mL/min/1.73 m² using CKD-EPI formula in adults, and Schwartz formula in adolescents
 - Direct (conjugated) bilirubin $< 2.0 \times ULN$
 - Alanine transaminase (ALT) $< 3.0 \times ULN$
7. ECOG performance status ≤ 2.0 for adults and Karnofsky $\geq 50\%$ for adolescents

Key inclusion and exclusion criteria: Gender

Both

Key inclusion and exclusion criteria: Specify gender

Key inclusion and exclusion criteria: Age minimum

12

Key inclusion and exclusion criteria: Age maximum

99

Key inclusion and exclusion criteria: Exclusion criteria

1. History of stem cell transplant.
2. Participating in a chronic transfusion program (pre-planned series of transfusions for prophylactic purposes) and/or planning on undergoing an exchange transfusion during the duration of the study; episodic transfusion in response to worsened anemia or VOC is permitted.
3. Contraindication or hypersensitivity to any drug or metabolites from similar class as study drug or to any excipients of the study drug formulation. History of severe hypersensitivity reaction to other monoclonal antibodies, which in the opinion of the investigator may pose an increased risk of serious infusion reaction.
4. Received active treatment on another investigational trial within 30 days (or 5 half-lives of that agent, whichever is greater) prior to Screening visit or plans to participate in another investigational drug trial.
5. Women of child-bearing potential, defined as all women physiologically capable of becoming pregnant unless they are using highly effective methods of contraception during dosing and for 15 weeks after stopping treatment.
6. Concurrent severe and/or uncontrolled medical conditions which, in the opinion of the Investigator, could cause unacceptable safety risks or compromise participation in the study.
7. History or current diagnosis of ECG abnormalities indicating significant risk of safety such as:
 - Resting QTcF ≥ 470 msec at pretreatment (baseline) for both male and female or instability to determine QTc
 - Concomitant clinically significant cardiac arrhythmias (e.g ventricular tachycardia), and clinically significant second or third degree AV block without a pacemaker
 - History of familial long QT syndrome or know family history of Torsades de Pointes
8. Not able to understand and to comply with study instructions and requirements.

Type of study

Interventional

Type of intervention

Pharmaceutical

Type of intervention: Specify type

N/A

Trial scope

Safety

Trial scope: Specify scope

N/A

**Study design: Allocation**

Randomized controlled trial

Study design: Control

Placebo

Study design: Purpose

Prevention

Study design: Assignment

Parallel

IMP has market authorization

No

Name of IMP

SEG101 - Crizanlizumab

Type of IMP

Immunological

Pharmaceutical class

anti-human P-selectin antibody G1

Therapeutic indication

prevention of vaso-occlusive crises (VOCs) in patients of all genotypes with sickle cell disease (SCD)

Therapeutic benefit

To compare the efficacy of 5.0 mg/kg versus placebo and 7.5 mg/kg of crizanlizumab versus placebo in addition to standard of care.

To compare the efficacy of 7.5 mg/kg versus placebo on the annualized rate of all VOCs (managed at home + leading to healthcare visit), based on documentation by health care provider following contact with participant.

To compare the efficacy of 5.0 mg/kg versus placebo on the annualized rate of all VOCs (managed at home + leading to healthcare visit)

Study model

N/A

Study model: Explain model

N/A

Study model: Specify model

N/A

Time perspective

N/A

Time perspective: Explain time perspective

N/A

Time perspective: Specify perspective

N/A

Target follow-up duration**Target follow-up duration: Unit****Number of groups/cohorts**

Biospecimen retention

Samples without DNA

Biospecimen description

All Blood samples and Urine Samples will be shipped to Covance Geneva Central Lab

Target sample size

10

Actual enrollment target size

10

Date of first enrollment: Type

Actual

Date of first enrollment: Date

07/08/2019

Date of study closure: Type

Actual

Date of study closure: Date

31/12/2021

Recruitment status

Recruiting

Recruitment status: Specify

Date of completion

30/06/2021

IPD sharing statement plan

No

IPD sharing statement description

Novartis is committed to sharing with qualified external researchers, access to patient-level data and supporting clinical documents from eligible studies. These requests are reviewed and approved by an independent review panel on the basis of scientific merit. All data provided is anonymized to respect the privacy of patients who have participated in the trial in line with applicable laws and regulations.

This trial data availability is according to the criteria and process described on www.clinicalstudydatarequest.com

Additional data URL

<https://clinicaltrials.gov/ct2/show/record/NCT03814746?term=cseg101a2301&rank=1>

Admin comments

Trial status

Approved

Secondary Identifying Numbers

| Full name of issuing authority | Secondary identifying number |
|--------------------------------|------------------------------|
| Clinicaltrials.gov | NCT03814746 |

Sources of Monetary or Material Support

| Name |
|-------------------------------|
| Novartis Pharma Services Inc. |



Secondary Sponsors

Name

NA

Contact for Public/Scientific Queries

| Contact type | Contact full name | Address | Country | Telephone | Email | Affiliation |
|--------------|-------------------|------------|---------|------------------------|-------------------------------|--|
| Public | Adlette Inati | Tripoli | Lebanon | 03228033 | adlette.inati@lau.edu.lb | Nini Hospital |
| Scientific | Hind Khairallah | Sin El Fil | Lebanon | +961 1 512002 Ext. 271 | Hind.Khairallah@fattal.com.lb | Khalil Fattal et Fils s.a.l. |
| Public | Miguel Abboud | Beirut | Lebanon | 03534213 | ma56@aub.edu.lb | American University of Beirut Medical Center |

Centers/Hospitals Involved in the Study

| Center/Hospital name | Name of principles investigator | Principles investigator speciality | Ethical approval |
|----------------------|---------------------------------|------------------------------------|------------------|
| Nini Hospital | Adlette Inati | Hematology | Approved |
| AUBMC | Miguel Abboud | Hematology | Approved |

Ethics Review

| Ethics approval obtained | Approval date | Contact name | Contact email | Contact phone |
|--|---------------|---------------|--------------------------------|-----------------------------|
| Nini Hospital | 20/05/2019 | Nabil Kabbara | Nabil.kabbara@hospitalnini.com | +961 (0) 6 431 400 ext 1062 |
| American University of Beirut Medical Center | 30/12/2019 | Fuad Ziyadeh | fz05@aub.edu.lb | +961 (0) 1 350 000 ext:5445 |



Countries of Recruitment

| Name |
|--------------------------|
| Lebanon |
| Belgium |
| Netherlands |
| United Kingdom |
| United States of America |
| Jordan |

Health Conditions or Problems Studied

| Condition | Code | Keyword |
|---------------------|-----------------------------|---------|
| Sickle Cell Disease | Sickle-cell disorders (D57) | SCD |

Interventions

| Intervention | Description | Keyword |
|--------------------------------------|--------------------------------------|--------------------------------------|
| ICF, Lab tests, IMP , Questionnaires | ICF, Lab tests, IMP , Questionnaires | ICF, Lab tests, IMP , Questionnaires |

Primary Outcomes

| Name | Time Points | Measure |
|---|-------------|---------|
| Rate of vaso-occlusive crisis (VOC) events leading to | 1 year | 1 year |
| To compare the efficacy of 5.0 mg/kg versus placebo and 7.5 mg/kg of crizanlizumab versus placebo in addition to standard of care | 1 year | 1 year |

Key Secondary Outcomes

| Name | Time Points | Measure |
|---|-----------------|-----------------|
| •Rate of all VOCs leading to healthcare visit and treated at home | 1 year, 5 years | 1 year, 5 years |
| •Number of days with VOC leading to healthcare visit | 1 year | 1 year |



Trial Results

Summary results

Study results globally

Date of posting of results summaries

Date of first journal publication of results

Results URL link

Baseline characteristics

Participant flow

Adverse events

Outcome measures

URL to protocol files