



CMBG453B12301 Study of Efficacy and Safety of MBG453 in Combination With Azacitidine in Subjects With Intermediate, High or Very High Risk Myelodysplastic Syndrome (MDS) as Per IPSS-R, or Chronic Myelomonocytic Leukemia-2 (CMML-2)

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Main Information

Primary registry identifying number

LBCTR2020094590

Protocol number

CMBG453B12301

MOH registration number

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 Pharmaceuticals

Primary sponsor: Country of origin

Novartis Pharmaceuticals

Date of registration in primary registry

15/07/2021

Date of registration in national regulatory agency

Public title

CMBG453B12301 Study of Efficacy and Safety of MBG453 in Combination With Azacitidine in Subjects With Intermediate, High or Very High Risk Myelodysplastic Syndrome (MDS) as Per IPSS-R, or Chronic Myelomonocytic Leukemia-2 (CMML-2)

Acronym

CMBG453B12301

Scientific title

A Randomized, Double-blind, Placebo-controlled Phase III Multi-center Study of Azacitidine With or Without MBG453 for the Treatment of Patients With Intermediate, High or Very High Risk Myelodysplastic Syndrome (MDS) as Per IPSS-R, or Chronic Myelomonocytic Leukemia-2 (CMML-2)

Acronym

Brief summary of the study: English

This is a Phase III multi-center, randomized, two-arm parallel-group, double-blind, placebo-controlled study of MBG453 or placebo added to azacitidine in adult subjects with intermediate, high or very high risk myelodysplastic syndrome (MDS) as per IPSS-R, or Chronic Myelomonocytic Leukemia-2 (CMML-2) who have an indication for treatment with azacitidine in first-line setting and are not eligible for intensive chemotherapy or hematopoietic stem cell transplantation (HSCT) according to medical judgment by the investigator.

The purpose of the current study is to assess clinical effects of MBG453 in combination with azacitidine in adult subjects with IPSS-R intermediate, high, very high risk MDS and CMML-2.

Brief summary of the study: Arabic





دراسة متعددة المراكز، عشوائية التوزيع، مزدوجة التعمية، مرتكزة على المقارنة بدواء وهمي، في المرحلة الثالثة حول دواء أزاسيتيدين مع أو MBG453 لعلاج المرضى المصابين بمتلازمة خلل التنسج النقوي العالية الخطورة أو ذات الخطورة العالية جداً وفقاً للنظام الدولي المنفتح لتسجيل النتائج (IPSS-R) 2 أو بسرطان الدم الوحيدّي النقوي المزمن-

Health conditions/problem studied: Specify

- Myelodysplastic Syndromes
- Leukemia, Myelomonocytic, Chronic

Interventions: Specify

- Drug: MBG453

A dose of MBG453 800 mg will be administered intravenously (IV) every 4 weeks (Q4W).

- Drug: Azacitidine

A dose of Azacitidine 75 mg/m² will be administered IV or subcutaneously (SC) on Day 1-7, or Day 1-5, 8 and 9.

- Drug: Placebo

A dose of Placebo 800 mg will be administered intravenously every 4 weeks (Q4W).

Key inclusion and exclusion criteria: Inclusion criteria

Inclusion Criteria:

- Signed informed consent must be obtained prior to participation in the study
- Age ≥ 18 years at the date of signing the informed consent form
- Morphologically confirmed diagnosis of myelodysplastic syndrome (MDS) based on WHO 2016 classification (Arber et al 2016) by local investigator assessment with one of the following Prognostic Risk Categories, based on the revised International Prognostic Scoring System (IPSS-R):
 - Very high (> 6 points)
 - High (> 4.5 - ≤ 6 points)
 - Intermediate (> 3 - ≤ 4.5 points) Or Morphologically confirmed diagnosis of Chronic Myelomonocytic Leukemia -2 based on WHO 2016 classification (Arber et al 2016) by local investigator assessment with WBC < 13 x 10⁹/L
- Indication for azacitidine treatment according to the investigator, based on local standard medical practice and institutional guidelines for treatment decisions
- Not eligible at time of screening for intensive chemotherapy according to the investigator, based on local standard medical practice and institutional guidelines for treatment decisions, including assessment of individual clinical factors such as age, comorbidities and performance status
- Not eligible at time of screening for hematopoietic stem cell transplantation (HSCT) according to the investigator, based on local standard medical practice and institutional guidelines for treatment decisions, including assessment of individual clinical factors such as age, comorbidities, performance status, and donor availability
- Eastern Cooperative Oncology Group (ECOG) performance status of 0, 1 or 2

Key inclusion and exclusion criteria: Gender

Both

Key inclusion and exclusion criteria: Specify gender

Key inclusion and exclusion criteria: Age minimum

18

Key inclusion and exclusion criteria: Age maximum

99

Key inclusion and exclusion criteria: Exclusion criteria

- Prior exposure to TIM-3 directed therapy at any time. Prior therapy with immune checkpoint inhibitors (e.g. anti-CTLA4, anti-PD-1, anti-PD-L1, or anti-PD-L2), cancer vaccines is allowed except if the drug was administered within 4 months prior to randomization
- Previous first-line treatment for intermediate, high, very high risk myelodysplastic syndromes (based on IPSS-R) or CMML with any antineoplastic agents including for example chemotherapy, lenalidomide and hypomethylating agents (HMAs) such as decitabine and azacitidine. However, previous treatment with hydroxyurea or leukopheresis to reduce WBC count is allowed prior to randomization.
- Investigational treatment received within 4 weeks or 5 half-lives of this investigational treatment, whatever is longer, prior to randomization. In case of a checkpoint inhibitor: a minimal interval of 4 months prior to randomization is necessary to allow randomization.
- Subjects with Myelodysplastic syndrome (MDS) based on 2016 WHO classification (Arber et al 2016) with revised International Prognostic Scoring System (IPSS-R) ≤ 3
- Diagnosis of acute myeloid leukemia (AML) including acute promyelocytic leukemia and extra-medullary acute myeloid leukemia, primary or secondary myelofibrosis based on WHO 2016 classification (Arber et al 2016)
- Diagnosis of therapy related myeloid neoplasms based on WHO 2016 classification (Arber et al 2016)
- History of organ or allogeneic hematopoietic stem cell transplant

Other protocol-defined Inclusion/Exclusion Criteria may apply.

Type of study

Interventional

**Type of intervention**

Pharmaceutical

Type of intervention: Specify type

N/A

Trial scope

Therapy

Trial scope: Specify scope

N/A

Study design: Allocation

Randomized controlled trial

Study design: Masking

Blinded (masking used)

Study design: Control

Placebo

Study phase

3

Study design: Purpose

Treatment

Study design: Specify purpose

N/A

Study design: Assignment

Parallel

Study design: Specify assignment

N/A

IMP has market authorization

No

IMP has market authorization: Specify**Name of IMP**

MBG453

Year of authorization**Month of authorization****Type of IMP**

Immunological

Pharmaceutical class

humanized monoclonal antibody against human TIM-3

Therapeutic indication

Patients with:
Myelodysplastic Syndromes or with chronic Leukemia Myelomonocytic

Therapeutic benefit

The primary objective of this study is to compare overall survival (OS) in the MBG453 plus azacitidine arm versus placebo plus azacitidine arm where OS is the time from randomization until death due to any cause.

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

Samples will be shipped to covance central laboratory and BMA will be

Target sample size

4

Actual enrollment target size

Date of first enrollment: Type

Anticipated

Date of first enrollment: Date

29/10/2020

Date of study closure: Type

Anticipated

Date of study closure: Date

25/11/2027

Recruitment status

Pending

Recruitment status: Specify

Date of completion

29/07/2022

IPD sharing statement plan

Yes

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 expert 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.

Additional data URL

<https://clinicaltrials.gov/ct2/show/record/NCT04266301?term=MBG&draw=2&rank=2&view=record>

Admin comments

Trial status

Approved

Secondary Identifying Numbers

| Full name of issuing authority | Secondary identifying number |
|--------------------------------|------------------------------|
| Clinical trials.gov | NCT04266301 |



Sources of Monetary or Material Support

Name

Novartis Pharmaceuticals

Secondary Sponsors

Name

NA

Contact for Public/Scientific Queries

| Contact type | Contact full name | Address | Country | Telephone | Email | Affiliation |
|--------------|-------------------|---------|---------|-----------------------|-------------------------------|--|
| Public | Ali Bazarbachi | Beirut | Lebanon | 961-1-350000 ext 5800 | bazarbac@aub.edu.lb | American University of Beirut Medical Center |
| Scientific | Hind Khairallah | Beirut | Lebanon | 961151200 2 ext 271 | hind.khairallah@fattal.com.lb | Khalil Fattal et Fils |

Centers/Hospitals Involved in the Study

| Center/Hospital name | Name of principles investigator | Principles investigator speciality | Ethical approval |
|--|---------------------------------|------------------------------------|------------------|
| American UNiversity of Beirut medical Center | Ali Bazarbachi | Hematology | Approved |

Ethics Review

| Ethics approval obtained | Approval date | Contact name | Contact email | Contact phone |
|--|---------------|--------------|-----------------|---------------------|
| American University of Beirut Medical Center | 31/08/2020 | Fuad Ziyadeh | fz05@aub.edu.lb | 9611350000 ext 5445 |



Countries of Recruitment

| Name |
|----------------|
| Lebanon |
| Australia |
| Austria |
| Belgium |
| Oman |
| Saudi Arabia |
| Czech Republic |
| Finland |
| France |
| Japan |
| Singapore |
| Spain |
| Switzerland |
| Taiwan |
| Thailand |

Health Conditions or Problems Studied

| Condition | Code | Keyword |
|-----------|---|---------|
| MDS | Myelodysplastic syndrome, unspecified (D46.9) | MDS |
| CMML 2 | Chronic monocytic leukaemia (C93.1) | CMML2 |

Interventions

| Intervention | Description | Keyword |
|---|---|---|
| Informed consent, IMP administration, questionnaire , Bone marrow aspirate, Lab tests | Informed consent, IMP administration, questionnaire , Bone marrow aspirate, Lab tests | Informed consent, IMP administration, questionnaire , Bone marrow aspirate, Lab tests |



Primary Outcomes

| Name | Time Points | Measure |
|------------------|-------------|---------|
| Overall Survival | 5 years | 5 years |

Key Secondary Outcomes

| Name | Time Points | Measure |
|--|-------------|---------|
| Time to definitive deterioration of fatigue using Functional Assessment of Cancer Therapy (FACIT)-Fatigue score | 5 years | 5 years |
| Red Blood Cell transfusion-free intervals | 5 years | 5 years |
| Percent of subjects with at least 3 point confirmed improvement from baseline in FACIT-fatigue scorescore | 5 years | 5 years |
| Percent of subjects with at least 10 point confirmed improvement from baseline in physical functioning using European Organization for Research and Treatment of Cancer's Core Quality of Life Questionnaire | 5 years | 5 years |
| Percentage of subjects with either CR, or mCR, or PR, or HI in each treatment arm according to International Working Group for MDS | 5 years | 5 years |
| Progression Free Survival (PFS) | 5 years | 5 years |
| Percentage of subjects with stable disease in each treatment arm according to International Working Group for MDS (IWG-MDS) as per investigator assessment | 5 years | 5 years |



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