



Study of Efficacy and Safety of LEE011 in Postmenopausal Women With Advanced Breast Cancer.(MONALEESA-2)

05/04/2025 08:00:26

Main Information

Primary registry identifying number

LBCTR2019050229

Protocol number

CLEE011A2301

MOH registration number

9695/ص

Study registered at the country of origin

Yes

Study registered at the country of origin: Specify

Type of registration

Retrospective

Type of registration: Justify

LCTR was recently initiated, original file was previously submitted by Paper

Date of registration in national regulatory agency

05/11/2014

Primary sponsor

Novartis Pharma Services Inc.

Primary sponsor: Country of origin

Novartis Pharmaceuticals

Date of registration in primary registry

09/12/2022

Date of registration in national regulatory agency

05/11/2014

Public title

Study of Efficacy and Safety of LEE011 in Postmenopausal Women With Advanced Breast Cancer.(MONALEESA-2)

Acronym

Scientific title

A Randomized Double-blind, Placebo-controlled Study of LEE011 in Combination With Letrozole for the Treatment of Postmenopausal Women With Hormone Receptor Positive, HER2 Negative, Advanced Breast Cancer Who Received no Prior Therapy for Advanced Disease

Acronym

Brief summary of the study: English

This is a multi-center, randomized, double-blinded, placebo controlled trial. The primary purpose of this study was to assess the efficacy of LEE011, as measured by progression free survival (PFS), in postmenopausal women with HR positive, HER2 negative advanced breast cancer who received no prior treatment for advanced disease.

Brief summary of the study: Arabic

وفعاليتها في معالجة Letrozole مع ليتروزول LEE011 دراسة عشوائية مزدوجة التعمية ومضبوطة بدواء وهمي حول سلامة استخدام الدواء والواتي لم يتلقين علاجاً سابقاً للمرض في2النساء بعد انقطاع الطمث المصابايات بسرطان ثدي متقدم إيجابي مستقبلات الهرمون وسليبي الهير مراحلها المتقدمة

Health conditions/problem studied: Specify

Advanced Breast Cancer

Interventions: Specify

•Drug: LEE011
Ribociclib was administered orally at a dose of 600 mg once daily (three 200 mg capsules).

•Drug: Letrozole



Letrozole 2.5 mg tablets taken orally.

•Drug: LEE011 Placebo

Matching ribociclib placebo was the control drug and was administered orally once daily.

Key inclusion and exclusion criteria: Inclusion criteria

- 1.Women with advanced (locoregionally recurrent or metastatic) breast cancer not amenable to curative therapy.
- 2.Patient is postmenopausal. Postmenopausal status is defined either by:
 - Prior bilateral oophorectomy
 - Age ≥ 60
 - Age < 60 and amenorrhea for 12 or more months (in the absence of chemotherapy, tamoxifen, toremifen, or ovarian suppression) and FSH and estradiol in the postmenopausal range per local normal range Note: For women with therapy-induced amenorrhea, serial measurements of FSH and/or estradiol are needed to ensure postmenopausal status. Ovarian radiation or treatment with a luteinizing hormone-releasing hormone agonist (LH-RHa) (goserelin acetate or leuprolide acetate) is not permitted for induction of ovarian suppression in this trial.
- 3.No prior systemic anti-cancer therapy for advanced disease.
- 4.Patient has a histologically and/or cytologically confirmed diagnosis of estrogen-receptor positive and/or progesterone receptor positive breast cancer by local laboratory.
- 5.Patient has HER2-negative breast cancer defined as a negative in situ hybridization test or an IHC status of 0, 1+ or 2+. If IHC is 2+, a negative in situ hybridization (FISH, CISH, or SISH) test is required by local laboratory testing.
- 6.Patient must have either:
 - Measurable disease, i.e., at least one measurable lesion as per RECIST 1.1 criteria (Tumor lesions previously irradiated or subjected to other locoregional therapy will only be considered measurable if disease progression at the treated site after completion of therapy is clearly documented).

OR

- If no measurable disease is present, then at least one predominantly lytic bone lesion must be present (Patients with no measurable disease and only one predominantly lytic bone lesion that has been previously irradiated are eligible if there is documented evidence of disease progression of the bone lesion after irradiation).

7.Patient has an Eastern Cooperative Oncology Group (ECOG) performance status 0 or 1

Key inclusion and exclusion criteria: Gender

Female

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

- 1.Patient who received any CDK4/6 inhibitor.
- 2.Patient who received any prior systemic anti-cancer therapy (including hormonal therapy and chemotherapy) for advanced breast cancer

Note:

- Patients who received (neo) adjuvant therapy for breast cancer are eligible. If the prior neo (adjuvant) therapy included letrozole or anastrozole the disease free interval must be greater than 12 months from the completion of treatment until randomization.
 - Patients who received ≤ 14 days of letrozole or anastrozole for advanced disease prior to randomization are eligible.
 - Any prior (neo) adjuvant anti-cancer therapy must be stopped at least 5 half-lives or 7 days, whichever is longer, before randomization
- 3.Patient is concurrently using other anti-cancer therapy.
 - 4.Patient has a concurrent malignancy or malignancy within 3 years of randomization, with the exception of adequately treated, basal or squamous cell carcinoma, non-melanomatous skin cancer or curatively resected cervical cancer.
 - 5.Patient has active cardiac disease or a history of cardiac dysfunction including any of the following:
 - History of angina pectoris, symptomatic pericarditis, or myocardial infarction within 12 months prior to study entry
 - History of documented congestive heart failure (New York Heart Association functional classification III-IV)
 - Documented cardiomyopathy
 - Patient has a Left Ventricular Ejection Fraction (LVEF) $< 50\%$ as determined by Multiple Gated acquisition (MUGA) scan or echocardiogram (ECHO)
 - History of any cardiac arrhythmias, e.g., ventricular, supraventricular, nodal arrhythmias, or conduction abnormality in the previous 12 months.
 - On screening, any of the following cardiac parameters:

bradycardia (heart rate < 50 at rest), tachycardia (heart rate > 90 at rest), PR interval > 220 msec, QRS interval > 109 msec, or QTcF > 450 msec.

- Systolic blood pressure > 160 or < 90 mmHg



6. Patient is currently receiving any of the following medications and cannot be discontinued 7 days prior start if the treatment:

- That are known strong inducers or inhibitors of CYP3A4.
- That have a known risk to prolong the QT interval or induce Torsades de Pointes.
- That have a narrow therapeutic window and are predominantly metabolized through CYP3A4.
- Herbal preparations/medications

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

Yes, Lebanon and Worldwide

IMP has market authorization: Specify

Lebanon and Worldwide

Name of IMP

Ribociclib

Year of authorization

2017

Month of authorization

8

Type of IMP

Others

Pharmaceutical class

Orally bioavailable, highly selective small molecule inhibitor of cyclin-dependent kinases 4 and 6 (CDK4/6).

Therapeutic indication

postmenopausal women with hormone receptor positive, HER2-negative, advanced breast cancer who received no prior therapy for advanced disease.

Therapeutic benefit

increase OS & PFS

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



Time perspective: Specify perspective

N/A

N/A

Target follow-up duration

Target follow-up duration: Unit

Number of groups/cohorts

Biospecimen retention

Samples with DNA**

Biospecimen description

Samples with circulating tumor DNA will be done, in addition to safety Labs (CBC, chemistry) sent to central lab : Covance located in Switzerland.

Target sample size

15

Actual enrollment target size

15

Date of first enrollment: Type

Actual

Date of first enrollment: Date

29/04/2014

Date of study closure: Type

Actual

Date of study closure: Date

28/03/2023

Recruitment status

Complete

Recruitment status: Specify

Date of completion

12/02/2015

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.

This trial data is currently available according to the process described on www.clinicalstudydatarequest.com.

URL: <http://www.clinicalstudydatarequest.com>

Additional data URL

<https://clinicaltrials.gov/ct2/show/record/NCT01958021?id=CLEE011A2301&rank=1>

Admin comments

Trial status

Approved



Secondary Identifying Numbers

| Full name of issuing authority | Secondary identifying number |
|--------------------------------|------------------------------|
| Clinicaltrials.gov | NCT01958021 |

Sources of Monetary or Material Support

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

Secondary Sponsors

No Sponsors

Contact for Public/Scientific Queries

| Contact type | Contact full name | Address | Country | Telephone | Email | Affiliation |
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| Public | Ziad Salem | Beirut | Lebanon | 9611347263 | zs04@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 |
|--|---------------------------------|------------------------------------|------------------|
| Hotel Dieu De France | Dr Joseph Kattan | Hematology Oncology | Approved |
| Hammoud Hospital University Medical Center | Dr Fadi Farhat | Hematology Oncology | Approved |
| Bellevue Medical Center | Dr Fadi El Karak | Hematology Oncology | Approved |
| American University of Beirut Medical Center | Dr Ziad Salem | Hematology Oncology | Approved |

Ethics Review

| Ethics approval obtained | Approval date | Contact name | Contact email | Contact phone |
|--|---------------|-----------------|-----------------------------|-----------------------------|
| American University of Beirut Medical Center | 05/09/2014 | Fuad Ziyadeh | fz05@aub.edu.lb | +961 (0) 1 350 000 ext:5445 |
| Hotel Dieu de France | 20/09/2013 | Sami Richa | cue@usj.edu.lb | 961421229 |
| Bellevue Medical Center | 20/02/2014 | Ghassan Maalouf | gmaalouf@bmc.com.lb | +961 (0) 1 682666 ext 5006 |
| Hammoud Hospital University Medical Center | 30/10/2013 | Ahmad Zaatari | zaatari@hammoudhospital.com | +961 (0) 7 723111 ext 1160 |



| Countries of Recruitment | |
|--------------------------|--|
| Name | |
| Lebanon | |
| Argentina | |
| Australia | |
| Austria | |
| Belgium | |
| Brazil | |
| Canada | |
| Denmark | |
| Finland | |
| France | |
| Germany | |
| Hungary | |
| Ireland | |
| Italy | |
| Netherlands | |
| Norway | |
| Singapore | |
| Spain | |
| Sweden | |
| Turkey | |
| United Kingdom | |
| United States of America | |



Health Conditions or Problems Studied

| Condition | Code | Keyword |
|------------------------|------------------------------------|---------|
| Advanced Breast Cancer | Malignant neoplasm of breast (C50) | ABC |

Interventions

| Intervention | Description | Keyword |
|--|--|--------------------------|
| Physical Exam, Vital signs, ECG, Echocardiography, Urinalysis, Serum/ urine pregnancy test, lab test, completion of QoL questionnaires | Physical Exam, Vital signs, ECG, Echocardiography, Urinalysis, Serum/ urine pregnancy test, lab test, completion of QoL questionnaires | ICF, Lab, IMP, radiology |

Primary Outcomes

| Name | Time Points | Measure |
|---------------------------|-------------|-----------------------------------|
| Progression Free Survival | 20 months | PFS up to approximately 20 months |

Key Secondary Outcomes

| Name | Time Points | Measure |
|------------------------------|-------------|-----------|
| •Overall Response Rate (ORR) | 20 months | 20 months |
| Overall survival | 65 months | 65 months |



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