



# Study Assessing the Efficacy and Safety of Alpelisib + Nab-paclitaxel in Subjects With Advanced TNBC Who Carry Either a PIK3CA Mutation or Have PTEN Loss Without PIK3CA Mutation

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## Main Information

### Primary registry identifying number

LBCTR2021044784

### Protocol number

CBYL719H12301

### 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 Pharma Services inc.

### Primary sponsor: Country of origin

Novartis Pharmaceuticals

### Date of registration in primary registry

29/07/2021

### Date of registration in national regulatory agency

### Public title

Study Assessing the Efficacy and Safety of Alpelisib + Nab-paclitaxel in Subjects With Advanced TNBC Who Carry Either a PIK3CA Mutation or Have PTEN Loss Without PIK3CA Mutation

### Acronym

### Scientific title

A Phase III, Multicenter, Randomized, Double-blind, Placebo-controlled Study to Assess the Efficacy and Safety of Alpelisib (BYL719) in Combination With Nab-paclitaxel in Patients With Advanced Triple Negative Breast Cancer With Either Phosphoinositide-3-kinase Catalytic Subunit Alpha (PIK3CA) Mutation or Phosphatase and Tensin Homolog Protein (PTEN) Loss Without PIK3CA Mutation

### Acronym

### Brief summary of the study: English

The purpose of this study is to determine whether treatment with alpelisib in combination with nab-paclitaxel is safe and effective in subjects with advanced triple negative breast cancer (aTNBC) who carry either a PIK3CA mutation (Study Part A) or have PTEN loss without PIK3CA mutation (Study Parts B1 and B2)

### Brief summary of the study: Arabic

دراسة متعددة المراكز، عشوائية التوزيع، مزدوجة التعمية، مرتكزة على المقارنة بدواء وهمي في المرحلة الثالثة، لتقييم فعالية وسلامة ألبيليسيب (BYL719) لدى المرضى المصابين بسرطان الثدي الثلاثي السلبي المتقدم إما مع طفرة (nab-paclitaxel) بالاشتراك مع ناب-باكليتاكسيل (BYL719) PIK3CA بدون طفرة في جين PTEN أو مع فقدان البروتين مماثل الفوسفاتاز والتسسين PIK3CA في حين

### Health conditions/problem studied: Specify

Triple Negative Breast Neoplasms

### Interventions: Specify



Drug: alpelisib  
300 mg orally once per day (QD)  
Other Name: BYL719  
Drug: placebo  
300 mg orally once per day (QD)  
Other Name: alpelisib matching placebo  
Drug: nab-paclitaxel  
100 mg/m<sup>2</sup> as IV infusion on Days 1, 8 and 15 of a 28-day cycle  
Other Name: abraxane

## Key inclusion and exclusion criteria: Inclusion criteria

Subject has histologically confirmed diagnosis of advanced (loco-regionally recurrent and not amenable to curative therapy, or metastatic (stage IV)) TNBC  
Subject has either a measurable disease per RECIST 1.1 criteria or, if no measurable disease is present, then at least one predominantly lytic bone lesion or mixed lytic-blastic bone lesion with identifiable soft tissue component (that can be evaluated by CT/MRI) must be present Part B1: patients must have measurable disease  
Subject has adequate tumor tissue to identify the PIK3CA mutation status (either carrying a mutation or without a mutation) and the PTEN loss status; both of which will determine whether the subject can be allocated to Part A - PIK3CA mutation regardless of PTEN status; or to Part B - PTEN loss without a PIK3CA mutation  
Subject has an Eastern Cooperative Oncology Group (ECOG) performance status of 0 or 1  
Subject has received no more than one line of therapy for metastatic disease.  
Subject has adequate bone marrow and organ function

## 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

Subject has received prior treatment with any PI3K, mTOR or AKT inhibitor  
Subject has a known hypersensitivity to alpelisib, nab-paclitaxel or to any of their excipients  
Subject has not recovered from all toxicities related to prior anticancer therapies to NCI CTCAE version 4.03 Grade ≤1; with the exception of alopecia  
Subject has central nervous system (CNS) involvement  
Subject with an established diagnosis of diabetes mellitus type I or uncontrolled type II based on Fasting Plasma Glucose and HbA1c  
Subject has impairment of gastrointestinal (GI) function or GI disease that may significantly alter the absorption of the study drugs (e.g., ulcerative diseases, uncontrolled nausea, vomiting, diarrhea, malabsorption syndrome, or small bowel resection) based on investigator discretion  
Subject has a history of acute pancreatitis within 1 year of screening or past medical history of chronic pancreatitis  
Subject has currently documented pneumonitis/interstitial lung disease  
Subject has a history of severe cutaneous reactions, such as Steven-Johnson Syndrome (SJS), erythema multiforme (EM), Toxic Epidermal Necrolysis (TEN) or Drug Reaction with Eosinophilia and Systemic Syndrome (DRESS)  
Subject with unresolved osteonecrosis of the jaw  
Other protocol-defined inclusion/exclusion criteria apply.

## 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: Masking

Blinded (masking used)

## Study design: Control

Placebo

## Study phase

3

## Study design: Purpose

Treatment

## Study design: Specify purpose

N/A

## Study design: Assignment

## Study design: Specify assignment

Parallel	N/A	
<b>IMP has market authorization</b>	<b>IMP has market authorization: Specify</b>	
Yes, Worldwide	US, EMEA, Australia, Lebanon	
<b>Name of IMP</b>	<b>Year of authorization</b>	<b>Month of authorization</b>
Alpelisib	2020	12
<b>Type of IMP</b>		
Gene therapy		
<b>Pharmaceutical class</b>		
Class I $\alpha$ -specific PI3K inhibitor		
<b>Therapeutic indication</b>		
Advanced Triple Negative Breast Cancer		
<b>Therapeutic benefit</b>		
to determine whether treatment with alpelisib in combination with nab-paclitaxel is safe and effective in subjects with advanced triple negative breast cancer (aTNBC) who carry either a PIK3CA mutation (Study Part A) or have PTEN loss without PIK3CA mutation (Study Parts B1 and B2)		
<b>Study model</b>	<b>Study model: Explain model</b>	
N/A	N/A	
<b>Study model: Specify model</b>		
N/A		
<b>Time perspective</b>	<b>Time perspective: Explain time perspective</b>	
N/A	N/A	
<b>Time perspective: Specify perspective</b>		
N/A		
<b>Target follow-up duration</b>	<b>Target follow-up duration: Unit</b>	
<b>Number of groups/cohorts</b>		
<b>Biospecimen retention</b>	<b>Biospecimen description</b>	
Samples with DNA**	Standard laboratory samples and biomarkers will be shipped to central labs : Navigate pharma in US and to Q2 solutions in UK	
<b>Target sample size</b>	<b>Actual enrollment target size</b>	
5		
<b>Date of first enrollment: Type</b>	<b>Date of first enrollment: Date</b>	
Anticipated	17/05/2021	

**Date of study closure: Type**

Anticipated

**Date of study closure: Date**

09/01/2026

**Recruitment status**

Pending

**Recruitment status: Specify****Date of completion**

24/08/2023

**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 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](http://www.clinicalstudydatarequest.com)

**Additional data URL**

<https://www.clinicaltrials.gov/ct2/show/study/NCT04251533?term=CBYL719H12301&draw=2&rank=1>

**Admin comments****Trial status**

Approved

## Secondary Identifying Numbers

Full name of issuing authority	Secondary identifying number
Clinical trials. gov	NCT04251533

## 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	Jawad Makarem	Al Chouf	Lebanon	03484288	Jawad.Makarem@awmedicalvillage.org	Ain Wazein Medical Village
Scientific	Hind Khairallah	Sinelfil	Lebanon	01512002#271	Hind.khairallah@fattal.com.lb	Khalil Fattal et Fils s.a.l.

## Centers/Hospitals Involved in the Study

Center/Hospital name	Name of principles investigator	Principles investigator speciality	Ethical approval
Ain Wazein Medical Village	Jawad Makarem	Hematology oncology	Approved

## Ethics Review

Ethics approval obtained	Approval date	Contact name	Contact email	Contact phone
Ain w Zein Medical Village	14/01/2021	Hayat Kamaledine	irb@awmedicalvillage.org	+961 (0) 5 509 001 ext 2014



Countries of Recruitment	
Name	
Lebanon	
Australia	
Austria	
Brazil	
Bulgaria	
Colombia	
Croatia	
France	
Germany	
Hungary	
India	
Italy	
Republic of Korea	
Malaysia	
Norway	
Poland	
Russian Federation	
Slovakia	
Spain	
Switzerland	
Taiwan	
Turkey	
United Kingdom	
United States of America	



## Health Conditions or Problems Studied

Condition	Code	Keyword
Triple Negative Breast Neoplasms	Malignant neoplasm of breast (C50)	Triple Negative Breast Neoplasms

## Interventions

Intervention	Description	Keyword
IMP administration , ICF, visit assessment and schedule	IMP administration , ICF, visit assessment and schedule	IMP administration , ICF, visit assessment and schedule

## Primary Outcomes

Name	Time Points	Measure
Progression-free Survival (PFS) Per Investigator Assessment in Study part A	Once approximately 192 PFS events in Study Part A had been observed	up to 35 months
Progression-free Survival (PFS) Per Investigator Assessment in Study part B2	Once approximately 192 PFS events in Study Part B2 had been observed	up to 22 months
Overall Response Rate (ORR) based on local radiology assessments in subjects with measurable disease at baseline in study Part B1	Up to 6 months	Up to 6 months

## Key Secondary Outcomes

Name	Time Points	Measure
Overall Survival (OS) in Study Part A	Up to 66 months	Up to 66 months
Overall Survival (OS) in Study Part B2	Up to 41 months	Up to 41 months
Overall response rate (ORR) with confirmed response in Study Part A	Up to 35 months	Up to 35 months
Overall response rate (ORR) with confirmed response in Study Part B2	Up to 22 months	Up to 22 months
Clinical benefit rate (CBR) with confirmed response in Study Part A	Up to 35 months	Up to 35 months
Clinical benefit rate (CBR) with confirmed response in Study Part B1	Up to 6 months	Up to 6 months
Clinical benefit rate (CBR) with confirmed response in Study Part B2	Up to 22 months	Up to 22 months
Time to response (TTR) in Study Part A	Up to 35 months	Up to 35 months
Time to response (TTR) in Study Part B1	Up to 6 months	Up to 6 months
Time to response (TTR) in Study Part B2	Up to 22 months	Up to 22 months
Duration of Response (DOR) with confirmed response in Study Part A	Up to 35 months	Up to 35 months



Duration of Response (DOR) with confirmed response in Study Part B1	Up to 6 months	Up to 6 months
Duration of Response (DOR) with confirmed response in Study Part B2	Up to 22 months	Up to 22 months
Overall Survival (OS) in Study Part B1	Up to 6 months	Up to 6 months
Progression-free Survival (PFS) Per Investigator Assessment in Study part B1	Up to 6 months	Up to 6 months
Plasma concentrations of alpelisib - Part A	Up to 35 months	Up to 35 months
Plasma concentrations of alpelisib - Part B1	Up to 6 months	Up to 6 months
Plasma concentrations of alpelisib -Part B2	up to 22 months	up to 22 months
Plasma concentrations of paclitaxel - Part A	Up to 35 months	Up to 35 months
Plasma concentrations of paclitaxel - Part B1	up to 6 months	up to 6 months
Change from baseline in the global health status/Quality of life (QoL) scale score of the European Organization for Research and Treatment of Cancer Quality of Life Questionnaire 30-item Core Module (EORTC QLQ-C30) in study Part A	Up to 35 months	Up to 35 months
Change from baseline in the global health status/QoL scale score of the EORTC QLQ-C30 in study Part B2	Up to 22 months	Up to 22 months
Time to 10% definitive deterioration in the global health status/QOL scale score of the EORTC QLQ-C30 in study Part A	Up to 35 months	Up to 35 months
Time to 10% definitive deterioration in the global health status/QOL scale score of the EORTC QLQ-C30 in study Part B2	Up to 22 months	Up to 22 months
PFS based on local radiology assessments using RECIST 1.1 criteria for subjects by PIK3CA mutation status measured in baseline ctDNA in study Part A	Up to 35 months	Up to 35 months
PFS based on local radiology assessments using RECIST 1.1 criteria for subjects by PIK3CA mutation status measured in baseline ctDNA in study Part B2	Up to 22 months	Up to 22 months
Time to definitive deterioration of the Eastern Cooperative Oncology Group (ECOG) performance status (PS) from baseline in Study Part A	Up to 35 months	Up to 35 months
Time to definitive deterioration of the ECOG performance status from baseline in Study Part B2	Up to 22 months	Up to 22 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