Results from a phase 1a clinical trial about using pamiparib together with tislelizumab to treat people with advanced solid tumors

Date of summary: December 9, 2022

results from recent research. • Pamiparib together with tislelizumab is not approved to treat cancer.

The purpose of this plain language summary is to help you understand the

- Researchers must look at the results of many types of studies to understand if a study drug works,
- how it works, and whether it is safe to prescribe to patients. • This summary reports the results of only 1 study. The results of this study may be different from the
- results of other studies.



Pamiparib: PAM-ih-per-rib

How to say:

the tissue.

Tislelizumab: Tiss-leh-LIZ-yu-mab

• Cancer is a disease where abnormal cells in a

person's body keep growing when they should not. These cells can form a tumor. Cells from

the tumor can invade nearby tissue and organs

What did this study look at?

- or spread through the blood system or lymph system to other parts of the body. Cancer is called advanced when it has spread from where it started to nearby tissue, lymph nodes, or distant parts of the body. Treatment for advanced cancer could
- help to make the tumor smaller, slow the growth of cancer cells, or relieve symptoms. Treatment is unlikely to completely cure advanced cancer.
- are potential cancer treatments. The people who took part in this study took both drugs (called a combination treatment).

• Cancer can start in the tissue or in the blood.

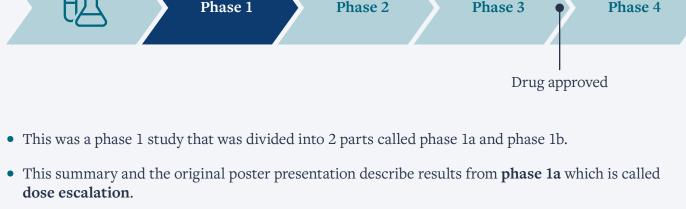
Cancer is called a **solid tumor** when it is in

• Pamiparib and tislelizumab are 2 drugs that

- The researchers wanted to see if pamiparib together with tislelizumab could be a future treatment for people with advanced solid tumors. • The study looked at side effects when people
- took different doses of pamiparib and tislelizumab to decide which dose should be used in larger studies.
- Where is this study in the drug development timeline? Studies in people

Lab studies

Phase 1 Phase 2 Phase 3



• Phase 1b is called dose expansion – these results will be described separately.

- Who took part in the study?



5 centers in Australia

The study included

Their average age was

took part

49 people.

63 years



women or assigned female at birth

7 out of 10 people had ovarian, fallopian tube, or peritoneal cancer

8 out of 10 people were



Researchers

recorded how

many side effects

happened during

the first 3 weeks

and how severe they were



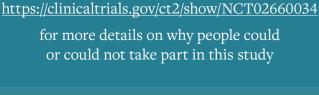
Who COULD

take part

People aged



Please visit the study website:



This carried on

joined the study

with the doses

shown below

Vomiting

Group 1:

until 5 groups had

Who COULD NOT take part

People who had

not taken at least 1

previous treatment

for their cancer

Study enrolment start date: January 22, 2016 Study enrolment end date: May 16, 2017

The next 3 people

who joined took

either the same

higher dose of

the drugs

dose or a slightly



The first 3 people

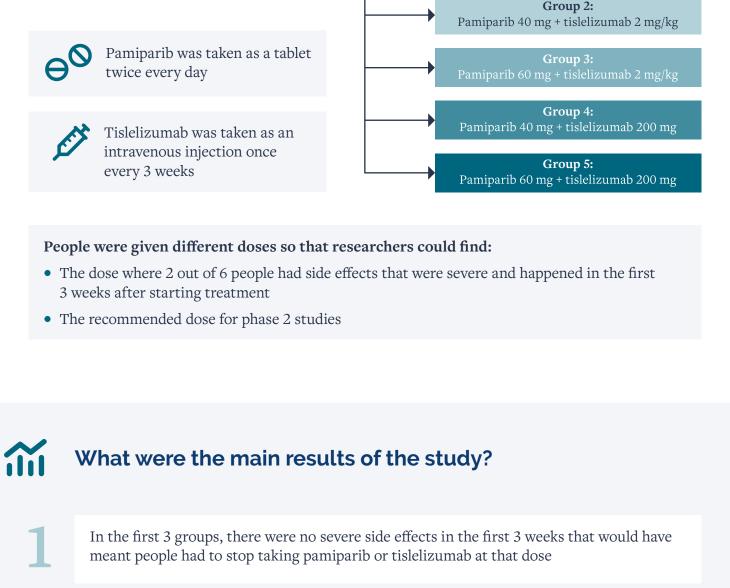
lowest doses of the

who joined the

study took the

2 drugs

Pamiparib 20 mg + tislelizumab 2 mg/kg 49 people



In group 4, there were 2 severe side effects in the first 6 patients. These side effects were:

More people joined group 4 but there were no more severe side effects in the first 3 weeks.

Recommended phase 2 dose =

pamiparib 40 mg twice every day + tislelizumab 200 mg every 3 weeks

Across all groups, 5 out of 10 people had a side effect that involved their immune system. Half of these side effects involved the liver, and they went away after steroid treatment

As a result, the dose for group 4 has been recommended for use in larger studies

If the group did

the next group

not have too many

severe side effects,

could join the study



A severe rash

Feeling sick

(nausea)

Everyone in the study had at least 1 side effect. Most sides effects were mild or moderate The most common side effects were:

Feeling tired

(fatigue)

Moderate nausea that did not go away with medication

What were some of the other results of the study?

Tumors became smaller for 2 out of 10 people

ovarian, fallopian tube, or peritoneal cancer

plus tislelizumab 200 mg every 3 weeks.

• Most side effects at this dose were mild or moderate.



Tumors stayed the same size for 3 out of 10 people

Tumors became smaller or stayed the same size for 5 out 10 people with

What were the main conclusions reported by the researchers?

• The dose to be used in larger phase 2 studies was pamiparib 40 mg twice every day



• As some side effects involved the immune system, researchers should look out for and treat these types of side effects in future studies. • More studies in people with specific tumor types will help researchers understand if pamiparib together with tislelizumab would work as a treatment for those tumors.

- Are there any plans for future studies? This phase of the study is completed. Phase 1b There are also other studies that are looking
 - Who sponsored this study?

at pamiparib or tislelizumab on their own

or combined with other treatments such

as chemotherapy.

Phone (USA): +1 (781) 801-1800 BeiGene would like to thank everyone who took part in this study.

of the study is ongoing and results will be

reported separately.

BeiGene

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Further information Study ID: NCT02660034

Website:

Chicago, IL, USA.

involved in preparing this summary.

The full title of the poster presentation is: Pamiparib in combination with tislelizumab in patients with advanced solid tumors: results

https://clinicaltrials.gov/ct2/show/NCT02660034

from the dose-escalation stage of a multicenter, open-label, phase 1a/b trial

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This summary was prepared by Envision Pharma Group. The original authors of the full poster presentation were not