

Vepdegestrant (ARV-471), a PROTAC[®] ER degrader, in people with ER+/HER2- advanced breast cancer

This summary contains information from the scientific poster:

Updated Results From VERITAC Evaluating Vepdegestrant, a PROteolysis TArgeting Chimera (PROTAC) Estrogen Receptor (ER) Degradator, in ER-Positive/Human Epidermal Growth Factor Receptor 2 (HER2)-Negative Advanced Breast Cancer

[CLICK HERE TO VIEW THE SCIENTIFIC POSTER](#)

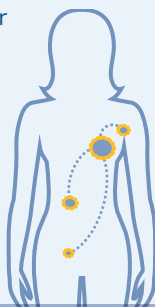
Copies of this poster obtained through this link are for personal use only and may not be reproduced without written permission from SABCS[®] and the author of this poster

What is ER+/HER2- advanced breast cancer?

ER+/HER2- breast cancer is one type of breast cancer

- Certain types of breast cancer grow in response to **estrogen**, a hormone (or **chemical messenger**) in your body. This is called **estrogen receptor-positive (ER+)** breast cancer
- Some types of breast cancer have a lot of a protein called **human epidermal growth factor receptor 2 (HER2)** and are called **HER2-positive (HER2+)**. Other breast cancer types have low levels or no HER2 and are called **HER2-negative (HER2-)**

Advanced breast cancer is cancer that has spread from the breast to nearby tissue (**locally advanced cancer**) or from the breast to more distant parts of the body (**metastatic cancer**)



What are some common treatments for ER+/HER2- advanced breast cancer?

Some treatments, called **endocrine therapies**, work by either blocking the body's ability to produce hormones, such as estrogen, or blocking the activity of these hormones in cancer cells. This may slow or stop cancer growth

- **Aromatase inhibitors**, such as letrozole, anastrozole, or exemestane, are endocrine therapies that reduce the production of estrogen
- **Fulvestrant** is an endocrine therapy that binds estrogen receptors leading to their degradation, which reduces estrogen's effects on tumors

Chemotherapy is a treatment that damages cancer cells. Sometimes people get chemotherapy prior to surgery to shrink the size of their tumor, after surgery to kill lingering cancer cells, or if their cancer has spread beyond the breast

CDK4/6 inhibitors are another type of treatment and work by blocking certain proteins that cause cancer cells to grow

What is vepdegestrant?

Vepdegestrant, also called **ARV-471**, is an orally administered investigational drug that is being evaluated as a treatment for ER+ breast cancer. It is a **PROteolysis TArgeting Chimera (PROTAC) estrogen receptor degrader**

- PROTAC protein degraders are designed to bind specific proteins of interest in cells, which causes those proteins to be **marked for elimination** by a natural protein disposal system in the body
- Vepdegestrant works by causing **estrogen receptors to be eliminated**, which blocks the activity of estrogen and could potentially stop ER+ breast cancer tumors from growing or cause the tumors to shrink

In the first part of a **clinical study that tested different doses of vepdegestrant** in people with ER+/HER2- advanced breast cancer who received prior treatments for their advanced cancer:

- During the study, **36%** of people had tumors that shrank or remained stable (neither grew nor shrank) **for at least 24 weeks following vepdegestrant treatment**
- The **side effects of vepdegestrant were mostly mild or moderate**

In the second part of the clinical study, researchers tested 2 doses of **vepedegestrant** in people with advanced breast cancer who received prior treatments for their advanced cancer

This summary describes updated results from the **35 participants who received vepedegestrant 200 mg once daily**, after following those patients for 12 months longer than previously described

The main aims of this study are to evaluate

- If vepedegestrant can cause tumors to stop growing or shrink in people with ER+/HER2- advanced breast cancer
- The side effects people who take vepedegestrant may experience

This summary describes

- How well vepedegestrant caused tumors to stop growing or shrink in people with ER+/HER2- advanced breast cancer and the side effects they experienced while taking the lower dose of vepedegestrant

Study Population

WHO PARTICIPATED IN THIS STUDY?

35 people with ER+/HER2- locally advanced or metastatic breast cancer enrolled in this study and were assigned to receive vepedegestrant 200 mg once daily



Before the study



had received a CDK4/6 inhibitor



had received an aromatase inhibitor



had received fulvestrant



had received chemotherapy

During the study

Participants took vepedegestrant 200 mg as pills by mouth once daily

Results

WHAT WERE THE RESULTS OF THE STUDY?

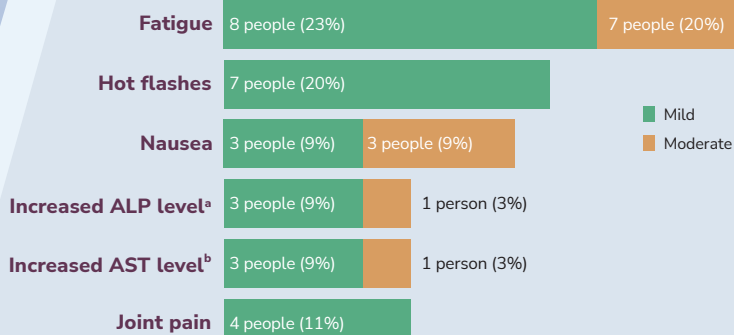


During the study, **tumors shrank or stopped growing for at least 24 weeks** in 37% of people



Half of the people lived without their cancer getting worse for **3.5 months or longer**

People taking vepedegestrant experienced **mostly mild or moderate side effects**. The most common side effects were:



^aALP is alkaline phosphatase, a substance produced by the liver and bones. ^bAST is aspartate aminotransferase, a substance produced by the liver

TAKE-HOME MESSAGES

Treatment with vepedegestrant 200 mg showed **clinical benefits for people with ER+/HER2- advanced breast cancer**

- Most of the side effects with vepedegestrant were **mild or moderate**
- A **larger study comparing vepedegestrant vs fulvestrant** in people with ER+/HER2- advanced breast cancer is ongoing
 - The **200 mg daily dose of vepedegestrant** was selected to use in the larger study based on the results from this study

Who sponsored this study?

This study is sponsored by **Arvinas Estrogen Receptor, Inc.**
5 Science Park
395 Winchester Ave.
New Haven, CT 06511
Phone (United States): +1 203-535-1456

Arvinas thanks the **people who volunteered to participate in this study and their caregivers**, as well as the **investigators, researchers, and coordinators** who contributed to this study

Writing and editorial support for this summary was provided by Justine Lempart, PhD, and Melissa Austin of Apollo Medical Communications, part of Helios Global Group, and funded by Arvinas Operations, Inc.

Where can I find more information?

For more information on **this study**

[VIEW CLINICAL TRIAL RECORD](#)

For more information on **clinical studies in general**

[VIEW INFORMATION](#)