

YOU NEVER THOUGHT YOU'D SEE THE DAY.

BEING THERE IS POSSIBLE™



WHAT IS ERLEADA®?

ERLEADA® (apalutamide) is a prescription medicine used for the treatment of prostate cancer:

- that has spread to other parts of the body and still responds to a medical or surgical treatment that lowers testosterone,

OR

- that has not spread to other parts of the body and no longer responds to a medical or surgical treatment that lowers testosterone.

It is not known if ERLEADA® is safe and effective in females.

It is not known if ERLEADA® is safe and effective in children.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

IMPORTANT SAFETY INFORMATION

ERLEADA® may cause serious side effects including:

- Heart disease, stroke, or mini-stroke; fractures and falls; seizure; and severe skin reactions
- The most common side effects of ERLEADA® include feeling very tired, joint pain, rash—tell your healthcare provider if you get a rash, decreased appetite, fall, weight loss, high blood pressure, hot flash, diarrhea, and fracture



IMPORTANT SAFETY INFORMATION (CONTINUED)

Before taking ERLEADA[®], tell your healthcare provider about all your medical conditions, including if you:

- have a history of heart disease
- have high blood pressure
- have diabetes
- have abnormal amounts of fat or cholesterol in your blood (dyslipidemia)
- have a history of seizures, brain injury, stroke, or brain tumors

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

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BEING THERE IS POSSIBLE™

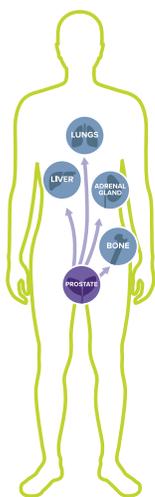
Click to Learn More About
Patient Support From *Janssen Compass*®



LIVING YOUR LIFE WITH PROSTATE CANCER

Prostate cancer is common and may be diagnosed early

It's important to understand your disease to help make treatment decisions with your healthcare provider. There are many factors to consider, from the type of cancer you have to the clinical study results for your medication.



mCSPC

WHAT IS METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (mCSPC)?

Prostate cancer that **HAS SPREAD** to other parts of the body and **STILL RESPONDS** to medical or surgical treatment that lowers testosterone.



nmCRPC

WHAT IS NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (nmCRPC)?

Prostate cancer that **HAS NOT SPREAD** to other parts of the body and **NO LONGER** responds to medical or surgical treatment that lowers testosterone.

IMPORTANT SAFETY INFORMATION (CONTINUED)

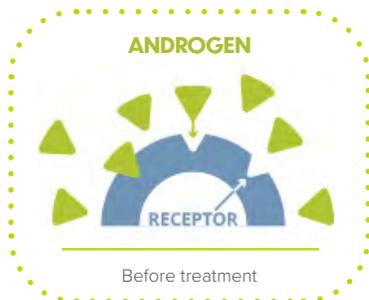
Before taking ERLEADA[®], tell your healthcare provider about all your medical conditions, including if you:

- are pregnant or plan to become pregnant. ERLEADA[®] can cause harm to your unborn baby and loss of pregnancy (miscarriage).
- have a partner who is pregnant or may become pregnant.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

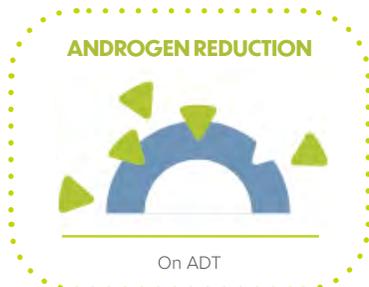
HOW ERLEADA® WORKS

ERLEADA® + ADT work together to lower androgens that can help fuel prostate cancer



How androgens help fuel prostate cancer

- Androgens are male hormones, primarily testosterone, that are needed for the prostate to function normally
- However, when androgens attach to androgen receptors, they can help fuel prostate cancer cell growth



The goal of ADT* is to lower androgen[†] levels

- Medical or surgical treatments that lower testosterone are also referred to as androgen deprivation therapy (ADT)
- ADT includes treatment to suppress or block the production or action of male hormones called androgens



ERLEADA® + ADT fight prostate cancer together

- ERLEADA® blocks androgens from attaching to receptors to help prevent cancer cells from growing

*ADT: Androgen deprivation therapy (ADT) includes medical or surgical treatment that lowers testosterone.

†Androgens: Male hormones, primarily testosterone, that are needed for the prostate to function normally but can help fuel prostate cancer cell growth.

IMPORTANT SAFETY INFORMATION (CONTINUED)

- Males who have female partners who are able to become pregnant should use effective birth control (contraception) during treatment and for 3 months after the last dose of ERLEADA®.
- Males should use a condom during sex with a pregnant female. Talk with your healthcare provider if you have questions about birth control.
- are breastfeeding or plan to breastfeed. It is not known if ERLEADA® passes into breast milk.



FOR MEN
WITH mCSPC,

BEING THERE LONGER IS POSSIBLE™

ERLEADA® + ADT was compared with placebo + ADT in a clinical study of 1,052 men with mCSPC

- In this study, men received either ERLEADA® 240 mg once daily or placebo. All men in the study received ADT

In a clinical study,
**ERLEADA® + ADT reduced the risk
of death by 35% vs placebo + ADT*†**



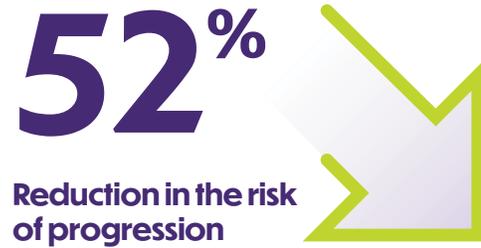
**Here's another way to look
at the results:**

At 4 years, approximately 65%
of men taking ERLEADA® + ADT
were alive vs 52% of men
taking placebo + ADT*

*Median (middle) follow-up time was 44.0 months. Median (middle) data has not been reached for ERLEADA®.

† In an earlier analysis from the study, the reduction in the risk of death was 33%.

In a clinical study,
**ERLEADA® + ADT reduced the risk of
prostate cancer getting worse by 52%
vs placebo + ADT‡**



**Here's another way to look
at the results:**

At 2 years, approximately 68%
of men taking ERLEADA® + ADT lived
without their disease getting worse
vs 48% of men taking placebo‡

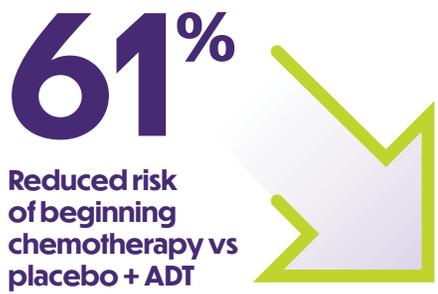
‡Median (middle) data has not been reached for ERLEADA®.

IMPORTANT SAFETY INFORMATION (CONTINUED)

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. ERLEADA® can interact with many other medicines.



Many of your best moments may still be ahead of you. ERLEADA® may help you be there.



Reduced risk of beginning chemotherapy vs placebo + ADT

The clinical study for men with mCSPC also evaluated time to chemotherapy:

ERLEADA® + ADT reduced the risk of beginning chemotherapy by 61% vs placebo + ADT[§]

[§]Median (middle) data has not been reached.

ADT: Androgen deprivation therapy (ADT) includes medical or surgical treatment that lowers testosterone.

Chemotherapy: Type of drug(s) that kill cancer cells.

Median: The middle number in a set of numbers. For 50% of people, this value was larger, and for 50% of people, it was smaller.

mCSPC: Metastatic castration-sensitive prostate cancer (mCSPC) is prostate cancer that HAS SPREAD to other parts of the body and STILL responds to medical or surgical treatment that lowers testosterone.

Placebo: Pronounced “pluh-see-bow”: a pill that looks like “real” medicine but contains nothing to affect health.

Progression: Disease spreading further as measured by imaging studies or dying.

Time to chemotherapy: Length of time from when patients began study to starting chemotherapy.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.



FOR MEN
WITH nmCRPC,

BEING THERE LONGER IS POSSIBLE™

ERLEADA® + ADT was compared with placebo + ADT in a clinical study of 1,207 men with nmCRPC

- In this study, men either received ERLEADA® 240 mg once daily or placebo. All men in the study received ADT

ERLEADA® HELPED SOME MEN WITH nmCRPC LIVE LONGER WITHOUT THE SPREAD OF CANCER.

In a clinical study, ERLEADA® + ADT delayed the spread of cancer to other parts of the body or death by a median of 24.3 months compared with placebo + ADT.

40.5 months

WITH ERLEADA® + ADT

vs

16.2 months

WITH PLACEBO + ADT

Additional study results in nmCRPC

THE CLINICAL STUDY FOR MEN WITH nmCRPC ALSO EVALUATED:

Overall survival: living longer



Men taking ERLEADA® + ADT

lived 14 months longer

than men taking placebo + ADT

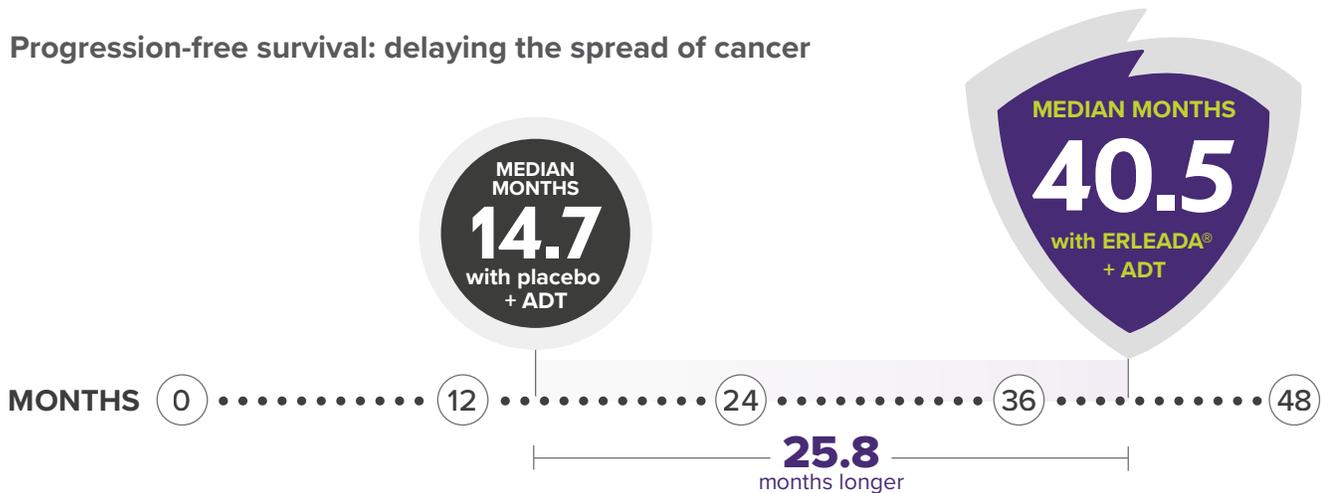
(73.9 months vs 59.9 months)

IMPORTANT SAFETY INFORMATION (CONTINUED)

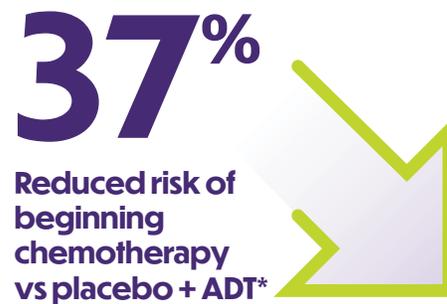
You should not start or stop any medicine before you talk with the healthcare provider that prescribed ERLEADA®.

Know the medicines you take. Keep a list of them with you to show to your healthcare provider and pharmacist when you get a new medicine.

Progression-free survival: delaying the spread of cancer



Time to chemotherapy: delaying the start of chemotherapy



*Median data has not been reached.

ADT: Androgen deprivation therapy (ADT) includes medical or surgical treatment that lowers testosterone.

Chemotherapy: Type of drug(s) that kill cancer cells.

Median: The middle number in a set of numbers. For 50% of people, this value was larger, and for 50% of people, it was smaller.

nmCRPC: Non-metastatic castration-resistant prostate cancer (nmCRPC) is prostate cancer that HAS NOT SPREAD to other parts of the body and NO LONGER responds to medical or surgical treatment that lowers testosterone.

Placebo: Pronounced "pluh-see-bow": a pill that looks like "real" medicine but contains nothing to affect health.

Progression-free survival: Length of time patients lived without their prostate cancer spreading to local or distant parts of the body or death.

Time to chemotherapy: Length of time from when patients began study to starting chemotherapy.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.



SIDE EFFECTS

ERLEADA® may cause serious side effects including:



Heart Disease, Stroke, or Mini-Stroke. Bleeding in the brain or blockage of the arteries in the heart or in part of the brain have happened in some people during treatment with ERLEADA® and can lead to death. Your healthcare provider will monitor you for signs and symptoms of heart or brain problems during your treatment with ERLEADA®. Call your healthcare provider or get medical help right away if you get:

- chest pain or discomfort at rest or with activity
- shortness of breath
- numbness or weakness of the face, arm, or leg, especially on one side of the body
- trouble talking or understanding
- trouble seeing in one or both eyes
- dizziness, loss of balance or coordination, or trouble walking



Fractures and Falls. ERLEADA® treatment can cause bones and muscles to weaken and may increase your risk for falls and fractures. Falls and fractures have happened in men during treatment with ERLEADA®. Your healthcare provider will monitor your risks for falls and fractures during treatment with ERLEADA®.



Seizure. Treatment with ERLEADA® may increase your risk of having a seizure. You should avoid activities where a sudden loss of consciousness could cause serious harm to yourself or others. Tell your healthcare provider right away if you have a loss of consciousness or seizure. Your healthcare provider will stop ERLEADA® if you have a seizure during treatment.



Severe Skin Reactions. Treatment with ERLEADA® may cause severe skin reactions that can be life-threatening or may lead to death. Stop taking ERLEADA® and get medical help right away if you develop any of these signs or symptoms of a severe skin reaction:

- severe rash or rash that continues to get worse
- fever or flu-like symptoms
- swollen lymph nodes
- blisters or sores in the mouth, throat, nose, eyes, or genital area
- blistering or peeling of the skin

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

SIDE EFFECTS (CONTINUED)

THE MOST COMMON SIDE EFFECTS OF ERLEADA® INCLUDE:



feeling very tired



weight loss



joint pain



high blood pressure



rash. Tell your healthcare provider if you get a rash



hot flash



decreased appetite



diarrhea



fall



fracture

Your healthcare provider may reduce your dose, temporarily stop, or permanently stop treatment with ERLEADA® if you have certain side effects.

A NONCHEMOTHERAPY TREATMENT YOU CAN TAKE AT HOME

Take ERLEADA® as prescribed

TWO FLEXIBLE DOSING OPTIONS

ERLEADA® is taken orally, once daily, with or without food

| | | |
|--|------------------|--|
| <p>FOUR 60 MG TABLETS,* ONCE DAILY</p>  <p>(~17 mm x 9 mm)</p> | <p>OR</p> | <p>ONE 240 MG TABLET,* ONCE DAILY</p>  <p>(~21 mm x 10 mm)</p> |
|--|------------------|--|

*Tablets shown are not actual size.

HOW SHOULD I TAKE ERLEADA®?



- Take ERLEADA® exactly as your healthcare provider tells you
- Do not stop taking your prescribed dose of ERLEADA® without talking with your healthcare provider first



- Take your prescribed dose of ERLEADA® 1 time a day, at the same time each day



- Swallow ERLEADA® tablets whole. **Do not** crush or split the tablets. If you cannot swallow ERLEADA® tablets whole, see the **INSTRUCTIONS FOR USE for detailed instructions** on how to prepare and take a dose of ERLEADA® by mouth. ERLEADA® comes in 2 different strengths (60 mg and 240 mg). Follow the instructions for your prescribed strength of ERLEADA®
- If you have a feeding tube, the ERLEADA® 240 mg tablet can be given through a feeding tube. See the **INSTRUCTIONS FOR USE for detailed instructions** on how to prepare and give a dose of ERLEADA® through a feeding tube (8 French or larger)



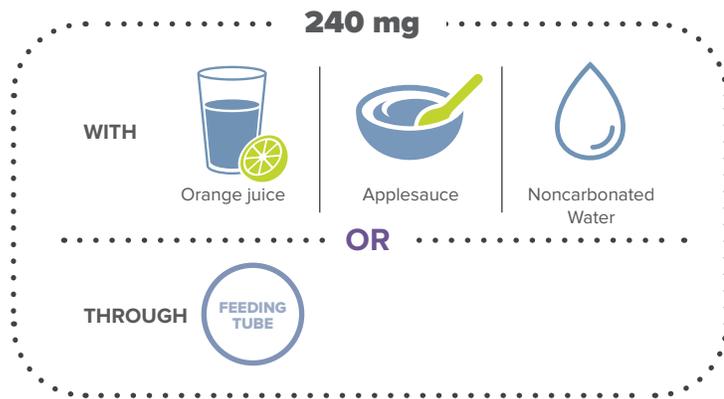
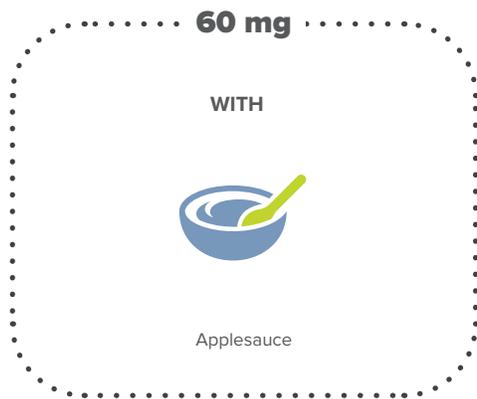
- If you miss a dose of ERLEADA®, take your normal dose as soon as possible on the same day. Return to your normal schedule on the following day. You should not take extra tablets to make up the missed dose
- You should start or continue a gonadotropin-releasing hormone (GnRH) analog therapy during your treatment with ERLEADA® unless you have had a surgery to lower the amount of testosterone in your body (surgical castration)



- If you take too much ERLEADA®, call your healthcare provider or go to the nearest hospital emergency room

HAVE TROUBLE SWALLOWING TABLETS WHOLE OR HAVE A FEEDING TUBE?†

Learn about alternate ways to take ERLEADA®



INSTRUCTIONS FOR USE†

The INSTRUCTIONS FOR USE contain information on how to prepare and give a dose of ERLEADA® tablets if you cannot swallow ERLEADA® tablets whole or if you have a feeding tube. Read the INSTRUCTIONS FOR USE before you prepare and take or give the first dose of ERLEADA®, and each time you get a refill. Ask your healthcare provider or pharmacist if you have any questions.

Important information you need to know before preparing a dose of ERLEADA®

- ERLEADA® comes in 2 different strengths, 60 mg tablets and 240 mg tablets. Please find the instructions on page 14 that refer to your prescribed ERLEADA® strength for how to prepare and take ERLEADA® tablets if you cannot swallow tablets whole
- ERLEADA® 240 mg tablet can be prepared and given through a feeding tube 8 French or larger

†Please see the full INSTRUCTIONS FOR USE on the next page.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.



PREPARING AND TAKING ERLEADA® IF YOU CANNOT SWALLOW TABLETS WHOLE

Preparing and taking ERLEADA® 60 mg tablets by mixing with applesauce:



Please read the complete directions below for mixing ERLEADA® with applesauce.

1. Place your dose of ERLEADA® in a container that contains 4 ounces (120 mL) of applesauce and stir. **Do not crush or split the tablets**
2. Wait 15 minutes and stir the mixture
3. Wait another 15 minutes and stir the mixture until tablets are well mixed with no chunks remaining
4. Swallow the mixture right away using a spoon
5. Rinse the container with 2 ounces (60 mL) of water and drink the water mixture right away
6. Repeat the rinse with 2 ounces (60 mL) of water one more time to make sure that you take your full dose of ERLEADA®
7. Swallow all the applesauce and medicine mixture within 1 hour of preparation. **DO NOT** store ERLEADA® that is mixed with applesauce

Preparing and taking ERLEADA® 240 mg tablet by placing the tablet in noncarbonated water and then mixing with orange juice, applesauce, or more noncarbonated water:



Please read the complete directions below for mixing ERLEADA® with orange juice, applesauce, or noncarbonated water.

1. Place the whole ERLEADA® 240 mg tablet in a cup. **Do not crush or split the tablet.**
2. Add about 2 teaspoons (10 mL) of noncarbonated water to make sure that the tablet is completely covered in water.
3. Wait 2 minutes until the tablet is broken up and spread out, and then stir the mixture.
4. Add 2 tablespoons (30 mL) of orange juice, applesauce, or noncarbonated water to the cup and stir the mixture.
5. Swallow the mixture right away.
6. Rinse the cup with enough noncarbonated water to make sure that you take your full dose of ERLEADA® and drink it right away. **DO NOT** store ERLEADA® that is mixed with noncarbonated water, orange juice, or applesauce for later use.

If you have a feeding tube, please [click](#) to see INSTRUCTIONS FOR USE and ask your healthcare provider or pharmacist if you have questions.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

TALKING TO YOUR DOCTOR ABOUT GETTING STARTED

Always remember that your healthcare team should be your main source of treatment direction and information. You should have an open and honest conversation with them.

TO HELP GET THE MOST OUT OF YOUR TREATMENT WITH ERLEADA®:



Take ERLEADA® exactly as your doctor tells you



Tell your doctor if you have any side effect that does not go away or that bothers you



Call your doctor for medical advice about side effects



Do not stop taking your prescribed dose of ERLEADA® without talking with your doctor first

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

 **Erleada**®
(apalutamide) 60 mg tablets
240 mg tablet

THERE ARE 2 WAYS TO RECEIVE YOUR ERLEADA® (apalutamide) PRESCRIPTION

Doctor's office pharmacy

If your doctor's office has its own pharmacy, your prescription may be available for pickup there.

OR

Specialty pharmacy

Your doctor will most likely send your prescription to a specialty pharmacy. Specialty pharmacies carry certain medications that are not available at retail pharmacies.

THE SPECIALTY PHARMACY PATH



When you receive your prescription:

- Your doctor's office will complete the necessary paperwork and then submit on your behalf to the specialty pharmacy. **Before you leave the doctor's office, make sure your paperwork has been submitted**



Within a few days:

- The specialty pharmacy will call you to discuss your prescription costs and arrange delivery by mail of your ERLEADA® right to your front door. **If you don't receive a call, contact your doctor's office**



Within 2 weeks:

- Your 30-day ERLEADA® prescription bottle should arrive by mail within 14 days. If it does not, contact your doctor's office

EVERY MONTH, YOU WILL BE CONTACTED TO COORDINATE YOUR MONTHLY REFILL.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

Once you and your doctor have decided that ERLEADA® is right for you, Janssen has resources to help support your treatment journey



GET THE SUPPORT YOU MAY NEED TO HELP YOU FEEL CONFIDENT THROUGHOUT YOUR TREATMENT JOURNEY

EASY-TO-ACCESS RESOURCES TO HELP YOU GET STARTED AND STAY ON TRACK WITH YOUR TREATMENT.



Explore options that may help you pay for your medication



Learn about your cancer, your Janssen medication, and get resources to support your practical and emotional needs during your treatment journey



Connect 1-on-1 with a **Janssen Compass® Care Navigator at 844-628-1234**, Monday through Friday, 8:30 AM–8:30 PM ET. A Care Navigator can help simplify and personalize your support throughout your treatment journey. Spanish-speaking Care Navigators are available.

CONNECT WITH US!

Your personal Care Navigator is just a phone call away.

Request a Call



Call us at 844-628-1234,
Monday through Friday, 8:30 AM–8:30 PM ET



Patient Support from
Janssen Compass®

Janssen Compass® is limited to education about your Janssen therapy, its administration, and/or your disease. It is intended to supplement your understanding of your therapy and is not intended to provide medical advice, replace a treatment plan from your doctor or nurse, or serve as a reason for you to start or stay on this medication.

PAYING FOR ERLEADA®

FOR COMMERCIALLY INSURED PATIENTS: SAVINGS PROGRAM

ELIGIBLE PATIENTS PAY AS LITTLE AS **\$0** PER MONTH

There is a limit to savings each year. Participate without sharing your income information. See [program requirements](#).



Call a Care Navigator to learn more about program requirements and enroll over the phone.

Request a Call



You can also call us at 844-628-1234,
Monday through Friday, 8:30 AM–8:30 PM ET

FOR MEDICARE PART D PATIENTS: *EXTRA HELP* AND *LOW-INCOME SUBSIDY*

ELIGIBLE PATIENTS PAY UP TO **\$10.35** PER MONTH

Call a Care Navigator to learn more about program requirements and how to enroll.

[Download Medicare Resource Guide Medicare Part D Extra Help — Low-Income Subsidy](#)

- You may be eligible for the Extra Help Program provided by the Social Security Administration
- The Low-Income Subsidy (LIS) program provides financial assistance for patients who may otherwise be unable to afford the costs associated with their Medicare Part D plan
- Eligible patients may be able to access brand-name drugs for up to \$10.35 per month

STATE HEALTH INSURANCE ASSISTANCE PROGRAMS (SHIPS)

A state program that gets money from the federal government to give free local health insurance counseling to people with Medicare.

Visit shiptacenter.org if you use government-funded healthcare programs such as Medicare, Medicaid, or need supplemental assistance with paying for your Janssen medication.

Visit shiptacenter.org



IF YOU DO NOT HAVE INSURANCE

A Care Navigator can be your guide to non-profit organizations, patient advocacy groups and state programs that may help with financial assistance.

Visit janssencompass.com to request your first call and learn more about how *Janssen Compass*[®] can be here for you.

You can also call us at 844-628-1234, Monday through Friday, 8:30 AM–8:30 PM ET.

Or text “CALL” to 844-628-1234* to request your first call!

*The personal information you provide about yourself through text message will be used by Janssen Biotech, Inc., and its affiliates and the program’s service providers (collectively Janssen) to contact you by phone to describe the *Janssen Compass*[®] program and complete the enrollment process. The [Privacy Policy](#) further explains how Janssen processes the personal information you provide. By texting “CALL,” you indicate that you have read, understand, and agree to these terms.

By texting “CALL” to 844-628-1234, you consent to receive automated text messages from the *Janssen Compass*[®] program to schedule a call with a dedicated Care Navigator and to learn more about the program. Please note, text message and data rates may apply. You may opt out at any time by texting “STOP” to 844-628-1234. Please do not share any unprompted personal or medical information as we are not able to reply to messages other than “CALL.” If you have specific questions, please contact a *Janssen Compass*[®] Care Navigator at 844-628-1234, Monday through Friday, 8:30 AM–8:30 PM ET. Call your doctor for medical advice about side effects. You may report side effects to the Janssen Medical Information Center by calling 800-526-7736.

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.



YOUR HEALTHCARE TEAM

Use this space to keep contact information for your healthcare team



Primary care physician's name: _____

Primary care physician's address: _____

Primary care physician's phone number: _____

Oncologist's name: _____

Oncologist's address: _____

Oncologist's phone number: _____

Urologist's name: _____

Urologist's address: _____

Urologist's phone number: _____

Specialty pharmacy's name: _____

Specialty pharmacy's address: _____

Specialty pharmacy's phone number: _____

Please see Important Safety Information throughout and on [pages 23-24](#), and [click here](#) for the full Prescribing Information.

STOP AND REFLECT

Use this space to write about hopes and goals for the journey ahead of you



What is one thing you are actively trying to change or maintain this month?

What is something you are looking forward to in the near future?

What is something new you'd like to try?

How can your friends and loved ones help support you?

Additional thoughts:



IMPORTANT SAFETY INFORMATION

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Before taking ERLEADA[®], tell your healthcare provider about all your medical conditions, including if you:

- have a history of heart disease
- have high blood pressure
- have diabetes
- have abnormal amounts of fat or cholesterol in your blood (dyslipidemia)
- have a history of seizures, brain injury, stroke, or brain tumors
- are pregnant or plan to become pregnant. ERLEADA[®] can cause harm to your unborn baby and loss of pregnancy (miscarriage).
- have a partner who is pregnant or may become pregnant.
 - Males who have female partners who are able to become pregnant should use effective birth control (contraception) during treatment and for 3 months after the last dose of ERLEADA[®].
 - Males should use a condom during sex with a pregnant female.Talk with your healthcare provider if you have questions about birth control.
- are breastfeeding or plan to breastfeed. It is not known if ERLEADA[®] passes into breast milk.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. ERLEADA[®] can interact with many other medicines.

You should not start or stop any medicine before you talk with the healthcare provider that prescribed ERLEADA[®].

Know the medicines you take. Keep a list of them with you to show to your healthcare provider and pharmacist when you get a new medicine.

How should I take ERLEADA[®]?

- Take ERLEADA[®] exactly as your healthcare provider tells you.
- Do not stop taking your prescribed dose of ERLEADA[®] without talking with your healthcare provider first.
- Take your prescribed dose of ERLEADA[®] 1 time a day, at the same time each day.
- Take ERLEADA[®] with or without food.
- Swallow ERLEADA[®] tablets whole.
- If you miss a dose of ERLEADA[®], take your normal dose as soon as possible on the same day. Return to your normal schedule on the following day. You should not take extra tablets to make up the missed dose.
- You should start or continue a gonadotropin-releasing hormone (GnRH) analog therapy during your treatment with ERLEADA[®] unless you have had a surgery to lower the amount of testosterone in your body (surgical castration).
- If you take too much ERLEADA[®], call your healthcare provider or go to the nearest hospital emergency room.

Please see Important Safety Information throughout, and [click here](#) for the full Prescribing Information.

IMPORTANT SAFETY INFORMATION (CONT'D)

What are the possible side effects of ERLEADA®?

ERLEADA® may cause serious side effects including:

- **Heart Disease, Stroke, or Mini-Stroke.** Bleeding in the brain or blockage of the arteries in the heart or in part of the brain have happened in some people during treatment with ERLEADA® and can lead to death. Your healthcare provider will monitor you for signs and symptoms of heart or brain problems during your treatment with ERLEADA®. Call your healthcare provider or get medical help right away if you get:
 - chest pain or discomfort at rest or with activity
 - shortness of breath
 - numbness or weakness of the face, arm, or leg, especially on one side of the body
 - trouble talking or understanding
 - trouble seeing in one or both eyes
 - dizziness, loss of balance or coordination, or trouble walking
- **Fractures and Falls.** ERLEADA® treatment can cause bones and muscles to weaken and may increase your risk for falls and fractures. Falls and fractures have happened in people during treatment with ERLEADA®. Your healthcare provider will monitor your risks for falls and fractures during treatment with ERLEADA®.
- **Seizure.** Treatment with ERLEADA® may increase your risk of having a seizure. You should avoid activities where a sudden loss of consciousness could cause serious harm to yourself or others. Tell your healthcare provider right away if you have a loss of consciousness or seizure. Your healthcare provider will stop ERLEADA® if you have a seizure during treatment.
- **Severe skin reactions.** Treatment with ERLEADA® may cause severe skin reactions that can be life-threatening or may lead to death. Stop taking ERLEADA® and get medical help right away if you develop any of these signs or symptoms of a severe skin reaction:
 - severe rash or rash that continues to get worse
 - fever or flu-like symptoms
 - swollen lymph nodes
 - blisters or sores in the mouth, throat, nose, eyes, or genital area
 - blistering or peeling of the skin

The most common side effects of ERLEADA® include:

- feeling very tired
- joint pain
- rash. Tell your healthcare provider if you get a rash
- decreased appetite
- fall
- weight loss
- high blood pressure
- hot flash
- diarrhea
- fracture

Your healthcare provider may reduce your dose, temporarily stop, or permanently stop treatment with ERLEADA® if you have certain side effects.

ERLEADA® may cause fertility problems in males, which may affect the ability to father children. Talk to your healthcare provider if you have concerns about fertility. **Do not** donate sperm during treatment with ERLEADA® and for 3 months after the last dose of ERLEADA®.

Tell your healthcare provider if you have any side effect that bothers you or that does not go away.

These are not all the possible side effects of ERLEADA®.

Call your doctor for medical advice about side effects. You may report side effects to FDA at [1-800-FDA-1088](https://www.fda.gov/medwatch).

[Click here](#) to see the full Prescribing Information for ERLEADA®.



cp-50508v5

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HELP YOU BE THERE.**



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 **Erleada**[®]
(apalutamide) 60 mg tablets
240 mg tablet

Please see Important Safety Information throughout and on [pages 23-24](#),
and [click here](#) for the full Prescribing Information.

Janssen Biotech, Inc.
800 Ridgeview Drive
Horsham, PA 19044

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