



DON'T LET CUSHING'S DISEASE
HOLD YOU BACK

START A NEW CHAPTER WITH ISTURISA



Model shown is not
an actual patient.

ISTURISA® (osilodrostat) Indication and Important Safety Information

Indications and Usage

ISTURISA (osilodrostat) is used to treat adults with Cushing's disease who cannot have pituitary surgery, or who have had pituitary surgery, but the surgery did not cure their Cushing's disease.

Important Safety Information:

- **Hypocortisolism:** Treatment with ISTURISA may cause symptoms associated with low levels of cortisol in your blood (hypocortisolism). Tell your healthcare provider right away if you experience more than one of the following symptoms, as these may be symptoms of very low cortisol level, known as adrenal insufficiency: nausea, vomiting, tiredness (fatigue), low blood pressure, stomach (abdominal) pain, loss of appetite, dizziness. If you get symptoms of hypocortisolism while taking ISTURISA, your healthcare provider may change your dose or ask you to stop taking it.

Please see additional Important Safety Information throughout and the enclosed Patient Information found at the end of the full Prescribing Information.



Isturisa[®]
(osilodrostat)

ISTURISA HELPS BRING CORTISOL LEVELS DOWN TO NORMAL^a

ISTURISA[®] (is tur ee' sah) (osilodrostat) is a prescription medicine used to treat adult patients with Cushing's disease who can't have pituitary surgery, or who have had pituitary surgery but were not cured of their Cushing's disease.

ISTURISA may help you reach and maintain normal cortisol levels^a

- The symptoms of Cushing's disease are caused by too much cortisol
- ISTURISA blocks an enzyme that helps make cortisol, so your adrenal glands don't make as much. Because less cortisol is made, less cortisol is released into your bloodstream
- ISTURISA is the first FDA-approved medicine for Cushing's disease that works this way

^aThe safety and effectiveness of ISTURISA were studied in a 48-week clinical trial with 137 patients with Cushing's disease that involved a dose titration period, a maintenance dose period, a randomized withdrawal period, and an open-label period during which all patients received ISTURISA. Period 3 (weeks 26 to 34) was the 8-week, randomized, double-blind withdrawal period where patients were randomly assigned to receive ISTURISA or placebo. At the end of Period 3, 31 out of 36 patients who received ISTURISA maintained normal cortisol levels vs 10 out of 34 patients who received placebo. Individual results may vary.

Important Safety Information (cont'd)

- **Heart Problem or Heart Rhythm Problem:** ISTURISA may cause an irregular heartbeat which could be a sign of a heart problem called QT prolongation. Call your healthcare provider right away if you have irregular heartbeats.

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START A NEW CHAPTER

We're here
to support
you during
your ISTURISA
treatment.

ISTURISA® (osilodrostat) offers:

- Proven results in patients with Cushing's disease
- Dosing that can be individualized to your specific needs
- Tablets that can be taken with or without food
- Dedicated nurse, pharmacist, and insurance specialist support through the Recordati Access, Resources, and Engagement (R.A.R.E.®) program

Personalized treatment for you

ISTURISA comes in 2 different tablet strengths: 1 mg and 5 mg. You may take a combination of different tablets based on how you respond. This allows your doctor to determine the precise dose of ISTURISA that works best for you.

The most common side effects of ISTURISA include:

- very low cortisol levels (adrenal insufficiency)
- tiredness (fatigue)
- nausea
- headache
- swelling of the legs, ankles, or other signs of fluid retention (edema)

These are not all of the possible side effects of ISTURISA.

Call your healthcare provider for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.



Important Safety Information (cont'd)

- **Increase in Other Adrenal Hormone Levels:** Your other adrenal hormones may increase when you take ISTURISA. Your healthcare provider may monitor you for the symptoms associated with these hormonal changes while you are taking ISTURISA such as low potassium (hypokalemia), high blood pressure (hypertension), swelling (edema) in the legs, ankles, or other signs of fluid retention, excessive facial or body hair growth (hirsutism), acne (in women). Call your healthcare provider if you have any of these side effects.

CORTISOL AND CUSHING'S DISEASE

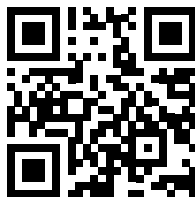
Cushing's disease can be a confusing and challenging journey—but knowing what you can about it can help along the way.

Cushing's disease is caused by a noncancerous tumor on the pituitary gland. One of the jobs of the pituitary gland is to tell your adrenal glands to make a hormone called cortisol. Cortisol is sometimes called the "stress hormone" because it helps regulate the body's response to stress. The tumor causes too much cortisol to be made.

Cortisol is important—but too much cortisol can lead to problems like obesity, diabetes, and high blood pressure. It is also what leads to the symptoms of Cushing's disease.

ISTURISA® (OSILODROSTAT) IS PROVEN TO HELP BRING CORTISOL LEVELS DOWN TO A NORMAL RANGE

Check out the results of our clinical trials on ISTURISA.com, and learn more about how ISTURISA may help you in your Cushing's disease journey.



Scan here to learn more.

Important Safety Information (cont'd)

- **Most common side effects include** very low cortisol levels (adrenal insufficiency), tiredness (fatigue), nausea, headache, swelling of the legs, ankles or other signs of fluid retention (edema). These are not all of the possible side effects of ISTURISA.

To report SUSPECTED SIDE EFFECTS, contact Recordati Rare Diseases Inc. at 1-888-575-8344, or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

We support you right from the start

Your R.A.R.E. Support Team

Our knowledgeable and reliable professionals have been fully trained about ISTURISA[®] (osilodrostat) and are dedicated to supporting your treatment goals. Your team includes:



INSURANCE SPECIALISTS
to help clarify coverage and
financial assistance available



PHARMACISTS who
are available to you 24/7,
every day of the year



NURSE EDUCATORS to
answer your questions and
support you during treatment



PATIENT LIAISONS (PLs)
to be there for you
throughout your journey

These services are provided at no additional cost to help you manage your therapy. Information provided is for educational purposes only and is not intended to replace the advice of your healthcare provider.

Our team will stay in touch throughout your therapy

Your enrollment in R.A.R.E. will be processed along with your prescription. However, if you have any questions, or would like to discuss how the program can help support your treatment, give us a call.



Scan here to get started with a PL.

R.A.R.E.[®]

Recordati Access, Resources, and Engagement

The R.A.R.E. Patient Support Program

1-888-855-RARE (7273)

Monday through Friday 8 AM – 8 PM ET

A pharmacist is always available

Learn more at www.isturisa.com

Please see Important Safety Information throughout and the enclosed Patient Information found at the end of the full Prescribing Information.



DON'T LET CUSHING'S DISEASE HOLD YOU BACK

IT'S TIME TO START A NEW CHAPTER WITH ISTURISA

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- **Heart Problem or Heart Rhythm Problem:** ISTURISA may cause an irregular heartbeat which could be a sign of a heart problem called QT prolongation. Call your healthcare provider right away if you have irregular heartbeats.
- **Increase in Other Adrenal Hormone Levels:** Your other adrenal hormones may increase when you take ISTURISA. Your healthcare provider may monitor you for the symptoms associated with these hormonal changes while you are taking ISTURISA such as low potassium (hypokalemia), high blood pressure (hypertension), swelling (edema) in the legs, ankles, or other signs of fluid retention, excessive facial or body hair growth (hirsutism), acne (in women). Call your healthcare provider if you have any of these side effects.
- **Most common side effects include** very low cortisol levels (adrenal insufficiency), tiredness (fatigue), nausea, headache, swelling of the legs, ankles or other signs of fluid retention (edema). These are not all of the possible side effects of ISTURISA.

To report SUSPECTED SIDE EFFECTS, contact Recordati Rare Diseases Inc. at 1-888-575-8344, or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Before taking ISTURISA, tell your doctor about all of your medical conditions, including if you:

- have or had heart problems, such as an irregular heartbeat, including a condition called prolonged QT syndrome (QT interval prolongation). Your healthcare provider will check the electrical signal of your heart (called an electrocardiogram) before you start taking ISTURISA, 1 week after starting ISTURISA, and as needed after that.
- have a history of low levels of potassium or magnesium in your blood.
- have liver problems.
- are breastfeeding or plan to breastfeed. It is not known if ISTURISA passes into your breast milk. **You should not breastfeed** if you take ISTURISA and for 1 week after stopping treatment.

Tell your healthcare provider about all the medicines you take, including prescription medicines, over-the-counter medicines, vitamins, and herbal supplements. Especially tell your healthcare provider if you take medicines used to treat certain heart problems. Ask your healthcare provider if you are not sure whether your medicine is used to treat heart problems.

Please report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088. You may also contact Recordati Rare Diseases Inc. at 1-888-575-8344 to report side effects.

ISTURISA® (osilodrostat) is available as 1-mg and 5-mg tablets.

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