Ask if QUZYTTIR® is right for you

Quzyttir®
Cetirizine HCl Injection
10 mg/mL

When talking with your doctor, use your answers to the following questions to help guide your discussion:

1. What treatments have you received for short-term hives? What has your experience with those treatments been like?

2. Do you have any concerns about your current treatment? If so, please describe.

3. Have you experienced any side effects after receiving treatment for short-term hives? If yes, how would you describe these side effects?

4. What are your treatment goals when you are experiencing short-term hives?

What is QUZYTTIR?

QUZYTTIR is an FDA-approved prescription medicine for the treatment of short-term hives (known as acute urticaria) in adults and children 6 months of age and older.

QUZYTTIR is not recommended in children younger than 6 years old with kidney or liver problems.

SELECT SAFETY INFORMATION

- You should not receive QUZYTTIR if you are allergic to any of its ingredients or if you are allergic to other medicines that contain levocetirizine or hydroxyzine.

Please see additional Important Safety Information on the next page.
Please see accompanying Full Prescribing Information.

QUZYTTIR has you covered with

the next-generation IV antihistamine that lasts 24 hours

Quzyttir®
Cetirizine HCl Injection
10 mg/mL

Reasons to ask about QUZYTTIR

- **Treats hives from allergic reactions**
- **Shown to cause less drowsiness***
- **Lasts for 24 hours**
- **Preservative-free solution for IV use**
- **No major interactions with other medications**

QUZYTTIR is an IV medicine that must be administered by a medical professional.

*Based on a clinical study comparing QUZYTTIR to IV diphenhydramine.

SELECT SAFETY INFORMATION

- QUZYTTIR may cause sleepiness. Be careful when driving a car or operating potentially dangerous machinery. Avoid alcohol or taking certain other medicines with QUZYTTIR, as this may raise your risk of having problems with alertness or thinking. Tell the healthcare provider treating you about all the medicines you are taking, including prescription or over-the-counter medicines, vitamins, or herbal supplements.

Support is available when you're ready

TerSera SupportSource

QUZYTTIR Reimbursement Resources
for both providers and patients.
To learn more, visit QUZYTTIR.com.

Not actual card

QUZYTTIR Co-Pay Program
Eligible patients could lower their co-pay **to as little as $0.†**
Learn more at QUZYTTIR.com.

†For eligible, commercially insured patients, the co-pay card is subject to an annual limit.. Patients are not eligible if prescriptions are paid by any state or federally funded program, including, but not limited to, Medicare or Medicaid, Medigap, VA, DOD, or TRICARE, or where prohibited by law. Additional terms and conditions apply.

IMPORTANT SAFETY INFORMATION

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- QUZYTTIR may cause sleepiness. Be careful when driving a car or operating potentially dangerous machinery. Avoid alcohol or taking certain other medicines with QUZYTTIR, as this may raise your risk of having problems with alertness or thinking. Tell the healthcare provider treating you about all the medicines you are taking, including prescription or over-the-counter medicines, vitamins, or herbal supplements.
- QUZYTTIR was not studied in pregnant or nursing women. Tell your healthcare provider if you are pregnant, plan to become pregnant, or are breastfeeding.
- The most common side effects (which occurred in <1% of patients) are: a change in sense of taste, headache, a tingling sensation, feeling faint, indigestion, feeling hot, and excessive sweating.
- The most common side effects observed in studies of an oral form of the active ingredient in QUZYTTIR taken by adults for a prolonged amount of time were sleepiness, tiredness, dry mouth, sore throat, and dizziness. In similar studies with children, the most common side effects were headache, sore throat, abdominal pain, sleepiness, diarrhea, nosebleed, bronchial spasm, nausea, and vomiting.
- Call your healthcare provider about any side effects that you may experience.

You are encouraged to report negative side effects of prescription drugs to the FDA. To report suspected adverse reactions, contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch. You may also contact TerSera Therapeutics at 1-844-334-4035 or medicalinformation@tersera.com.

Please see accompanying Full Prescribing Information.

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QYR-P-0161 v2 (08/2024)

TerSera® therapeutics