



QUESTIONS TO ASK YOUR DOCTOR

When you visit your doctor, it can help to know in advance what kinds of questions you may want to ask about treatment with SEROQUEL XR. This guide is designed to make it easier for you to prepare.

- :: What aspects of bipolar disorder is SEROQUEL XR indicated for?
- :: Does SEROQUEL XR treat bipolar depression?
- :: How does SEROQUEL XR work?
- :: Is SEROQUEL XR approved for maintenance therapy?
- :: How often should SEROQUEL XR be taken?
- :: When should SEROQUEL XR be taken?
- :: What things should I consider when starting SEROQUEL XR?
- :: What are the potential side effects associated with SEROQUEL XR?
- :: In addition to medication, what are other parts of my treatment plan?
- :: Are there any patient support programs available with free information, resources, and practical advice?
- :: How can I save on my prescription with the SEROQUEL XR Savings Card?

For additional information regarding the SEROQUEL XR Savings Card, please contact AstraZeneca at **1-800-236-9933**.



Notes:

Important Safety Information for SEROQUEL XR

Elderly patients with dementia-related psychosis (having lost touch with reality due to confusion and memory loss) treated with this type of medicine are at an increased risk of death, compared to placebo (sugar pill). SEROQUEL XR is not approved for treating these patients.

Antidepressants have increased the risk of suicidal thoughts and actions in some children, teenagers, and young adults. Patients of all ages starting treatment should be watched closely for worsening of depression, suicidal thoughts or actions, unusual changes in behavior, agitation, and irritability. Families and caregivers should watch patients daily and report these symptoms immediately to the physician. SEROQUEL XR is not approved for patients under the age of 18 years.


Important Safety Information and Indications continues on page 2.

Please print the [Prescribing Information](#), including Boxed Warnings and Medication Guide, and discuss it with your health care provider.



Important Safety Information and Indications for SEROQUEL XR (continued)

- High blood sugar and diabetes have been reported with SEROQUEL XR and medicines like it. If you have diabetes or risk factors such as obesity or a family history of diabetes, ask your doctor about checking your blood sugar before starting SEROQUEL XR and regularly throughout treatment. If you develop symptoms of high blood sugar or diabetes, such as excessive thirst or hunger, increased urination, or weakness, contact your doctor. Complications from diabetes can be serious and even life threatening
- Increases in triglycerides and in LDL (bad) cholesterol and decreases in HDL (good) cholesterol have been reported with SEROQUEL XR. Your doctor should check your cholesterol levels before you start SEROQUEL XR and during therapy
- Weight gain has been reported with SEROQUEL XR. Your doctor should check your weight regularly
- A rare, but potentially fatal, side effect reported with SEROQUEL XR and medicines like it is neuroleptic malignant syndrome (NMS). Tell your doctor if you have very high fever; rigid muscles; shaking; confusion; sweating; changes in pulse, heart rate, or blood pressure; or muscle pain and weakness because treatment should be stopped if you have NMS
- Another serious side effect reported with SEROQUEL XR and medicines like it is tardive dyskinesia (TD)— uncontrollable movements of the face, tongue, or other parts of the body. TD may become permanent, and the risk of TD is believed to increase as the length of time on and the amount of these medications increase. While TD can develop in patients taking low doses for short periods, this is much less common. There is no known treatment for TD, but it may go away partially or completely if treatment is stopped
- Before starting treatment, tell your doctor if you have high prolactin levels or have a history of, or are at risk for, seizures or a low white blood cell (WBC) count. An eye exam for cataracts is recommended at the beginning of treatment and every 6 months thereafter
- Other risks include feeling dizzy or lightheaded upon standing, or having trouble swallowing. Tell your doctor if you experience any of these
- Suicidal thoughts or actions may occur; tell your doctor if you have thoughts about death or suicide
- Since drowsiness has been reported with SEROQUEL XR, you should not participate in activities such as driving or operating machinery until you know that you can do so safely. Avoid drinking alcohol while taking SEROQUEL XR because SEROQUEL XR increases the effects of alcohol. Avoid becoming overheated or dehydrated while taking SEROQUEL XR
- Common side effects: The most common side effects are drowsiness, dry mouth, increases in cholesterol and triglycerides, constipation, upset stomach, dizziness, a sudden drop in blood pressure upon standing, weight gain, increased hunger, tiredness, increases in blood sugar, difficulty speaking, and stuffy nose

This is not a complete summary of safety information. Please discuss the full [Prescribing Information](#)  with your health care provider.

Indications

SEROQUEL XR is a once-daily tablet approved to treat acute depressive episodes in bipolar disorder; acute manic or mixed episodes in bipolar disorder alone or when added to lithium or divalproex; and long-term maintenance of bipolar disorder when added to lithium or divalproex.

Please print the [Prescribing Information](#) , including Boxed Warnings and Medication Guide, and discuss it with your health care provider.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.FDA.gov/medwatch or call 1-800-FDA-1088.

Patient photos are intended to be representative of typical patients with bipolar disorder and are not of actual patients.

Message for Health Care Professionals:

In order to keep effective drugs available on the market for use by you and your patients, the FDA relies on the voluntary reporting of serious adverse events that you suspect are associated with the use of an FDA-regulated drug. In the interest of patient safety, please be sure to notify the FDA or the manufacturer of any such events that you become aware of when discussing this patient questionnaire with your patients or otherwise.

