Restoril (temazepam)

Generic name: Temazepam  
Available strengths: 7.5 mg, 15 mg, 30 mg capsules  
Available in generic: Yes  
Drug class: Benzodiazepine/sedative-hypnotic

General Information

Restoril (temazepam) is a benzodiazepine sedative-hypnotic medication approved for short-term treatment of insomnia. Similar to other benzodiazepines, Restoril has anxiolytic effects (i.e., relieves anxiety), but it is seldom prescribed for this use. It has an intermediate duration of action (i.e., half-life around 8–12 hours) and no active metabolite. Patients with early morning awakening may find Restoril beneficial because of its intermediate duration of action with little or no daytime drowsiness or grogginess. Generally, Restoril should not be used for longer than 1 week. However, longer use occasionally may be necessary for some patients; in such cases, careful monitoring is needed to prevent physical or psychological dependence. As with other benzodiazepines, Restoril is associated with dependence and abuse and is therefore regulated as a controlled substance by federal and state laws.

Dosing Information

The usual dose of Restoril is 15 mg at bedtime. The dose may be increased to 30 mg if needed but should not exceed this amount. Seniors may require a lower dose of 7.5–15 mg at bedtime. Restoril should be taken about 1 hour before retiring, allowing for the absorption of the medication.

Common Side Effects

The common side effects of Restoril are daytime drowsiness and sedation, especially shortly after initiating therapy. Other frequent complaints are impaired concentration and memory, feeling of dissociation ("spacey"), and impaired coordination.
**Adverse Reactions and Precautions**

Restoril may affect alertness and coordination the next day after taking a single bedtime dose. Patients should exercise caution when driving or performing other tasks requiring alertness while taking this medication. Seniors may be more adversely affected, because it may affect their coordination and reflexes and lead to falls and injury. Taking Restoril with other central nervous system (CNS) depressants, such as alcohol, narcotics, and barbiturates, may compound these CNS effects.

Prolonged use of benzodiazepines can lead to dependence. When the medication is abruptly withdrawn, symptoms of withdrawal may occur. Withdrawal symptoms include headache, vomiting, impaired concentration, confusion, tremor, muscle cramps, and seizures. Benzodiazepines are centrally acting depressants, and they can depress respiration. This can affect patients with chronic obstructive pulmonary disease and emphysema by decreasing their “respiratory drive” or their ability to breathe. Patients with sleep apnea—a sleep disorder in which respiration is interrupted by long pauses during the sleep cycle—should not take Restoril or other benzodiazepines. The respiratory depressant effect of benzodiazepines may further suppress the respiratory drive in these patients and put them at risk for respiratory depression.

Benzodiazepines may induce paradoxical reactions in susceptible individuals. Instead of the expected depressant effects, the medication stimulates excitement, aggression, anger, uninhibited behavior, and rage in the susceptible person. These reactions are more likely to occur in seniors, people with brain damage, and individuals with personality and impulse-control disorders.

**Possible Drug Interactions**

The potential drug interactions with Restoril are summarized in the table below.

<table>
<thead>
<tr>
<th>Central nervous system (CNS) depressants (e.g., alcohol, narcotics, barbiturates, hypnotics) and antihistamines</th>
<th>Combination of Restoril with another CNS depressant may impair coordination and breathing, increase sedation, and other produce other CNS depressant effects.</th>
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<tr>
<td>Tagamet (cimetidine), Serzone (nefazodone), oral contraceptives, Antabuse (disulfiram), Prozac (fluoxetine), Luvox (fluvoxamine), isoniazid (e.g., INH), Diflucan (fluconazole), Nizoral (ketoconazole), Sporanox (itraconazole), protease inhibitors (e.g., Crixivan, Norvir, Fortovase)</td>
<td>When any of these medications are taken concurrently with Restoril, they can inhibit its metabolism and increase blood levels. This may increase the likelihood of adverse side effects from Restoril (e.g., sedation, drowsiness).</td>
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</table>

Patients taking Restoril should not consume alcohol because the combination may increase sedation and drowsiness.

**Use in Pregnancy and Breastfeeding: Pregnancy Category X**

Benzodiazepines and their metabolites are known to cross the placenta and accumulate in the fetal circulation. Reproduction studies in animals demonstrated that Restoril was absorbed into fetal circulation and increased the occurrence of abnormalities. Restoril should not be used during pregnancy.
Nursing mothers should not take Restoril, because it will pass into breast milk and be ingested by the baby. If stopping the drug is not an alternative, breastfeeding should not be started or should be discontinued.

**Overdose**

Overdose from oral ingestion of benzodiazepines alone is generally not fatal. Most fatalities reported with benzodiazepines implicate multiple medication ingestion, particularly the combination of a benzodiazepine with CNS depressants, including alcohol, narcotics, and barbiturates.

Mild symptoms of benzodiazepine overdose include drowsiness, confusion, somnolence, tiredness, impaired coordination, clumsiness in walking (ataxia), and slow reflexes. Benzodiazepine overdose, when these agents are taken alone, is rarely fatal. When multiple medications are taken in benzodiazepine overdose, severe symptoms include slowing of respiratory and heart rate, low blood pressure, loss of coordination, and loss of consciousness leading to coma and, potentially, death.

Any suspected overdose should be treated as an emergency. The person should be taken to the emergency department for observation and treatment. The prescription bottle of medication (and any other medication suspected in the overdose) should be brought as well, because the information on the prescription label can be helpful to the treating physician in determining the number of pills ingested.

**Special Considerations**

- Restoril should only be taken when needed for sleep. Do not take more than the prescribed dose.
- Restoril may cause daytime sedation and drowsiness, especially during initiation of therapy, and impair your alertness. Use caution when driving or performing tasks that require alertness. Avoid alcohol when taking Restoril. The combination may increase sedation and drowsiness.
- Store the medication in its originally labeled, light-resistant container, away from heat and moisture. Heat and moisture may precipitate breakdown of your medication.
- Keep your medication out of reach of children.

*If you have any questions with your medication, consult your physician or pharmacist.*