Prozac/Prozac Weekly/Sarafem (fluoxetine)

Generic name: Fluoxetine
Available strengths: 10 mg, 20 mg, 40 mg capsules or tablets;
   90 mg delayed-release capsule (Prozac Weekly);
   20 mg/5 mL oral solution
Available in generic: Yes
Drug class: Selective serotonin reuptake inhibitor antidepressant

General Information

Prozac (fluoxetine) was approved by the U.S. Food and Drug Administration (FDA) for the treatment of major depressive disorder, obsessive-compulsive disorder (OCD), bulimia nervosa (a binge eating and vomiting disorder), and premenstrual dysphoric disorder (PMDD). The use of a medication for its approved indications is called its labeled use. In clinical practice, however, physicians often prescribe medications for unlabeled (“off-label”) uses when published clinical studies, case reports, or their own clinical experiences support the efficacy and safety of these medications for these unapproved indications. Prozac may be used to treat other psychiatric disorders, including panic disorder, generalized anxiety disorder, social anxiety disorder, and posttraumatic stress disorder.

Prozac was the first selective serotonin reuptake inhibitor (SSRI) approved by the FDA for the treatment of PMDD. The symptoms occur during a specific phase of the menstrual cycle just prior to menstrual bleeding, and the woman typically presents with labile mood, anger, irritability, and depression. Prozac is marketed under the brand name Sarafem specifically for PMDD.

Prozac is a serotonin-specific medication that works by blocking the reuptake of the neurotransmitter serotonin back into brain cells, thereby increasing its levels in the brain. Depression and other mental disorders may be caused by abnormally low levels of serotonin. The presumed action of Prozac and other SSRIs is to increase serotonin levels, which may help to restore those areas of the brain to normal function.
Dosing Information

The usual starting dose of Prozac in depression is 20 mg (capsule or tablet), taken once a day in the morning. If no improvement is seen after 3–4 weeks, the dosage may be increased in increments of 10 mg to a maximum dosage of 60 mg/day. If the patient’s symptoms are stable with a 20 mg/day dosage, the dosage may be switched to 90 mg of the delayed-release Prozac Weekly once a week for dosing convenience, with similar therapeutic effect. Seniors and people with severe or chronic medical conditions may require a lower starting dosage. Generally, higher dosages of Prozac are required for treatment of other psychiatric disorders. Treatment of OCD, for example, may require dosages between 80 mg and 100 mg daily. For treatment of PMDD, Prozac is marketed under the brand name Sarafem, which merely contains Prozac. For patients who cannot take a tablet or capsule, Prozac also comes in liquid form.

For most people, it may take as long as 3–4 weeks to experience the optimal effects of the antidepressant. The duration of medication treatment depends on the individual's personal psychiatric history and family history. For instance, the length of medication treatment will be longer for those who have had two or more previous episodes of major depressive disorder. For most people, the medication may be tapered 6 months after their depression responds to treatment. However, a small percentage of patients will continue to have depressive symptoms after their antidepressant is reduced or stopped. These individuals may benefit from continuing to take Prozac for 1 year or longer.

Common Side Effects

The most common side effects reported with Prozac are nervousness, jitteriness, nausea, and insomnia. There is a high rate of sexual dysfunction in individuals taking Prozac. The most frequent sexual side effects reported were delayed or lack of orgasm in women and retarded ejaculation in men. Some people may experience decreased desire or lack of interest in sexual drive. Occasionally, individuals report headaches, sleepiness, changes in appetite, excessive sweating, stomach cramps, diarrhea, and constipation with Prozac.

Patients should discuss these side effects with their physician, especially if they continue to be bothersome 3–4 weeks after the medication is started. If a rash or any other severe symptoms develop, patients should contact their physician immediately.

Adverse Reactions and Precautions

Prozac may cause drowsiness in some people. Patients should not drive or operate machinery until they are certain that their alertness or coordination is not affected by the medication. Patients with a known allergy to Prozac or who have experienced a severe reaction after taking it should not take Prozac.

Use in Pregnancy and Breastfeeding: Pregnancy Category C

Prozac has not been tested in women to determine its safety in pregnancy. The effects of Prozac on the developing fetus in pregnant women are unknown. Women who are pregnant or may become pregnant should discuss this with their physician. Some women may experience a recurrence of their depression when they stop their antidepressant. In these circumstances it may be necessary to restart the medication or seek an alternative medication or treatment.
Nursing mothers should not take Prozac because small amounts will pass into breast milk and be ingested by the baby. If stopping the antidepressant is not an alternative, breastfeeding should not be started or should be discontinued.

### Possible Drug Interactions

Prozac may increase the levels of other medications by inhibiting their metabolism in the liver. This interaction may result in higher levels of the inhibited medication and thus increase its potential for toxicity. The clinically significant drug interactions with Prozac are summarized in the table below.

<table>
<thead>
<tr>
<th>Medication</th>
<th>Interaction</th>
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<tbody>
<tr>
<td>Coumadin (warfarin)</td>
<td>Prozac may increase Coumadin levels and its anticoagulant effects, resulting in bleeding. This interaction is less likely than with other selective serotonin reuptake inhibitors (SSRIs), but Coumadin therapy should be monitored closely when starting any SSRI.</td>
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<tr>
<td>Tricyclic antidepressants (TCAs)</td>
<td>Prozac may elevate TCA levels, increasing the potential for toxicity.</td>
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<tr>
<td>Valium (diazepam)</td>
<td>Prozac may elevate levels of diazepam and diazepam-like medications, enhancing sedation and impairment of coordination.</td>
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<tr>
<td>Anticonvulsants</td>
<td>Prozac may elevate levels of anticonvulsants such as Dilantin (phenytoin), Tegretol (carbamazepine), and Depakote (divalproex sodium), increasing the potential for toxicity.</td>
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<tr>
<td>Antipsychotics</td>
<td>Prozac may elevate levels of antipsychotic medications, including Haldol (haloperidol), Clozaril (clozapine), Mellaril (thioridazine), and Risperdal (risperidone), possibly increasing their side effects.</td>
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Other medications, including herbal supplements (such as St. John’s wort), that boost serotonin can result in excessive levels of the neurotransmitter serotonin when combined with Prozac, producing a toxic syndrome known as serotonin syndrome. The early signs of serotonin syndrome are restlessness, confusion, tremors, flushing, and involuntary muscle jerks. If the medications are not stopped, the individual may develop more life-threatening complications resulting in muscle disorders, high fever, respiratory problems, clotting problems, and destruction of red blood cells that can lead to acute renal failure. Hence, patients taking Prozac should be alert to the possible signs of serotonin syndrome, which require immediate medical attention and discontinuation of the serotonin-boosting medications.

Antidepressants known as monoamine oxidase inhibitors (MAOIs) should not be taken together with Prozac, because the combination may potentially produce a toxic reaction that includes elevated temperature, high blood pressure, and extreme excitation and agitation. Patients should consult their physician or pharmacist before taking any new medications, including over-the-counter medications and herbal supplements, with Prozac.

Patients taking Prozac should avoid alcohol or should consume it in moderation because the combination may worsen depression.
Overdose

Like other SSRIs, Prozac is much safer in overdose than the older tricyclic antidepressants and some of the newer antidepressants. However, unlike the other SSRIs, Prozac has a very long duration of action, and therefore it takes much longer to eliminate Prozac from the body. Deaths from massive overdoses of Prozac have been reported. Usually, Prozac was combined with other medications in cases of fatal outcomes.

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

Most cases of major depression can be treated successfully, usually with medication, psychotherapy, or both. The combination of psychotherapy and antidepressants is very effective in treating moderate to severe depression. The medications improve mood, sleep, energy, and appetite while therapy strengthens coping skills, deals with possible underlying issues, and improves thought patterns and behavior.

In general, antidepressants alone help about 60%–70% of those taking them. Although a few individuals may experience some improvement from antidepressants by the end of the first week, most people do not see significant benefits from their antidepressants until after 3–4 weeks, and it can sometimes take as long as 8 weeks for the medication to produce its full effects. Thus it is critical that patients continue to take their antidepressant long enough for the medication to be beneficial and that patients not get discouraged and stop their medication prematurely if they do not feel better immediately.

The controversial issue of suicide and antidepressants has prompted the FDA to ask manufacturers of some antidepressants, particularly the SSRIs, to provide warnings in their package insert that the risk of suicide may be increased in depressed individuals (especially children) the first several weeks after beginning an antidepressant. However, studies have found that when more people in a community are taking antidepressants, the suicide rate is lower. The risk of suicide is inherent in depression and may persist until the individual responds to treatment. Depressed individuals who are at risk for suicide should be closely watched at the outset of therapy, and any signs of suicidal or violent behavior should be immediately reported to the physician or a mental health provider.

- Do not discontinue Prozac abruptly. Your dosage should be gradually tapered before stopping to prevent any discontinuation symptoms.
- If you miss a dose, take it as soon as possible, within 2–3 hours of the scheduled dosing. If it is close to your next scheduled dose, skip the missed dose and continue on your regular dosing schedule, but do not take double doses.
- Prozac may be taken with or without food.
- 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 about your medication, consult your physician or pharmacist.