SEROPLAX XR—Use with caution in patients with diseases or conditions that could affect hemodynamic stability. Use Effexor XR cautiously in patients with diseases or conditions that could affect hemodynamic stability. For patients who experience a rapid increase in blood pressure, or whose blood pressure does not return to baseline within 10-15 minutes after therapy, the patient should be observed for several hours.

LAbORATORY TESTS: There are no specific laboratory tests recommended.

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In Depression or Generalized Anxiety Disorder

The goal is recovery

- Working on both serotonin and norepinephrine, EFFEXOR XR has been shown to offer more patients the ability to achieve recovery.

Visit us at www.EFFEXORXR.com

The efficacy and safety of EFFEXOR XR for pediatric use have not been established.

EFFEXOR XR is contraindicated in patients taking monoamine oxidase inhibitors (MAOIs). EFFEXOR XR should not be used in combination with an MAOI or within at least 14 days of discontinuing treatment with an MAOI; at least 7 days should be allowed after stopping EFFEXOR XR before starting an MAOI.

The most common adverse events reported in EFFEXOR XR placebo-controlled depression trials (incidence ≥10% and ≥2× that of placebo) were nausea, dizziness, somnolence, abnormal ejaculation, sweating, dry mouth, and nervousness; and in GAD trials were nausea, dry mouth, insomnia, abnormal ejaculation, anorexia, constipation, nervousness, and sweating.

Treatment with venlafaxine is associated with sustained increases in blood pressure (BP) in some patients. Three percent of EFFEXOR XR patients in depression studies (doses of 75 to 375 mg/day) and 0.4% in GAD studies (doses of 75 to 225 mg/day) had sustained BP elevations. Less than 1% discontinued treatment because of elevated BP. Regular BP monitoring is recommended.