

A six week double blind multi center placebo controlled study evaluating the efficacy and safety of flexible doses of oral ziprasidone as add on adjunctive therapy with lithium, valproate or lamotrigine in Bipolar I Depression.

Inclusion Criteria:

- Male or female patients aged 18 or older
- Patients with a diagnosis of Bipolar I Disorder, most recent episode depressed with or without rapid cycling without psychotic features.
- Patients currently receiving or willing to begin lithium, valproate, may currently be on lamotrigine.

Exclusion Criteria:

- Pregnant or nursing (lactating) women
- Any comorbid Axis I or Axis II disorder that is clinically unstable, requires treatment, or that has been a primary focus or treatment within the 6 months prior to screening.
- Attempted suicide within 3 months prior to screening.
- Patients who have had treatment with investigational drugs at the time of enrollment, or within the last 30 days
- History of Chronic Hepatitis, seizures, or elevated TSH.

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