

הודעה על תכשירים גנריים חדשים

חברת נוברטיס ישראל בע"מ מבקשת לעדכן על אישור משרד הבריאות אודות רישומם של התכשירים הגנריים
החדשים הבאים:

- Aripiprazole Sandoz 5** (aripiprazole 5mg tablets)
- Aripiprazole Sandoz 10** (aripiprazole 10mg tablets)
- Aripiprazole Sandoz 15** (aripiprazole 15mg tablets)
- Aripiprazole Sandoz 20** (aripiprazole 20mg tablets)
- Aripiprazole Sandoz 30** (aripiprazole 30mg tablets)

ההתוויות כפי שאושרו בתעודת הרישום:**5mg; 10mg; 15mg; 20mg:**

Aripiprazole Sandoz is indicated for the treatment of schizophrenia, for the treatment of moderate to severe manic episodes in Bipolar I disorder and for the prevention of a new manic episode in patient who experienced predominantly manic episodes and whose manic episodes responded to aripiprazole treatment. Aripiprazole Sandoz is indicated for use as an adjunctive therapy to antidepressants for the treatment of major depressive disorder (MDD). Efficacy was established in two 6-week trials in adults with MDD who had an inadequate response to antidepressant therapy during the current episode. Aripiprazole Sandoz is indicated for the treatment of irritability associated with autistic disorder. Efficacy was established in two 8-week trials in pediatric patients (aged 6 to 17 years) with irritability associated with autistic disorder (including symptoms of aggression towards others, deliberate self-injuriousness, temper tantrums, and quickly changing moods).

30mg:

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העלון לרופא נשלח לפרסום במאגר התרופות שבאתר משרד הבריאות:

<https://www.health.gov.il/Subjects/PharmAndCosmetics/DrugsRegistryDB/Pages/DrugsDatabase.aspx>

כמו כן ניתן לקבל את העלון המודפס על ידי פניה לבעל הרישום – נוברטיס ישראל בע"מ, בטלפון 03-9201111

בברכה,

אבי ילצינדג
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נוברטיס ישראל בע"מ



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