

Drug Information Table
Tricyclic antidepressants – amitriptyline

Therapeutic Use	Administration	
<ul style="list-style-type: none"> • Treatment of major depression 	<ul style="list-style-type: none"> • Give orally at bedtime. • Monitor for therapeutic effects after 1 to 3 weeks. • Expect long-term use to control depression. 	
Side/Adverse Effects	Interventions	Patient Instructions
<ul style="list-style-type: none"> • Drowsiness, sedation 	<ul style="list-style-type: none"> • Monitor for sedation and, if it occurs, take measures to prevent falls. 	<ul style="list-style-type: none"> • Take at bedtime to prevent daytime drowsiness. • Do not drive or perform hazardous activities if drowsy.
<ul style="list-style-type: none"> • Orthostatic hypotension 	<ul style="list-style-type: none"> • Monitor orthostatic vital signs. 	<ul style="list-style-type: none"> • Do not drive or perform hazardous activities if drowsy. • Move slowly from lying to sitting or standing.
<ul style="list-style-type: none"> • Anticholinergic effects (dry mouth, constipation, urinary retention, blurred vision) 	<ul style="list-style-type: none"> • Monitor for these effects. 	<ul style="list-style-type: none"> • Urinate before taking the daily dose. • Increase fiber and fluids to prevent constipation. • Chew gum, suck on hard candy, or sip water to prevent dry mouth.
<ul style="list-style-type: none"> • Increased risk for suicide (especially in children, adolescents) 	<ul style="list-style-type: none"> • Monitor for increases in depression and suicidal ideation. • Initiate suicide precautions when appropriate. 	<ul style="list-style-type: none"> • Report any feelings of self-harm or worsening of depression.
<ul style="list-style-type: none"> • Withdrawal symptoms with abrupt discontinuation (anxiety, headache, muscle pain, nausea) 	<ul style="list-style-type: none"> • Taper the drug over 2 weeks to prevent or minimize withdrawal. 	<ul style="list-style-type: none"> • Do not stop taking the drug abruptly.
<ul style="list-style-type: none"> • High risk for overdose (life-threatening dysrhythmias, confusion, seizures) 	<ul style="list-style-type: none"> • Assure that patients have no more than a 1-week supply of the drug. • For overdose, prepare for gastric lavage and administer physostigmine to treat anticholinergic effects and lidocaine to treat ventricular dysrhythmias. 	<ul style="list-style-type: none"> • Take the drug exactly as prescribed.
Contraindication	Precautions	Interactions
<ul style="list-style-type: none"> • Allergy to TCAs • Children younger than 12 years • Recent acute myocardial infarction • Cardiac dysrhythmias • Seizure disorder history • Concurrent use with MAOIs 	<ul style="list-style-type: none"> • Angle closure glaucoma • Prostatic hypertrophy • History of urinary retention • Liver or renal disorders • Suicidal ideation • History of electroconvulsive therapy • Schizophrenia • Hematologic or respiratory disorders • Diabetes mellitus • Alcohol use disorder 	<ul style="list-style-type: none"> • CNS depressants increase sedation. • Levodopa/carbidopa and sympathomimetic drugs may cause increased effects of those drugs, such as hypertension. • Administration within 2 weeks of MAOIs may cause hypertensive crisis. • Cimetidine (Tagamet) and methylphenidate (Ritalin) increase amitriptyline levels. • Anticholinergic drugs (such as antihistamines) increase anticholinergic effects. • Ginkgo biloba increases the risk of seizures; St. John's wort may cause serotonin syndrome.