PRODUCT MONOGRAPH

[Pr]AURO-OLANZAPINE ODT

olanzapine orally disintegrating tablets

5 mg, 10 mg, 15 mg and 20 mg

House Standard

Antipsychotic Agent

Auro Pharma Inc.
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CANADA

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PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	All Nonmedicinal Ingredients
oral	orally disintegrating tablets / 5 mg, 10 mg, 15 mg and 20 mg	Aspartame, mannitol, polacrilin potassium, crospovidone, colloidal silica anhydrous, microcrystalline cellulose, sodium stearyl fumarate and artificial pineapple flavor.

INDICATIONS AND CLINICAL USE

Adults:

Schizophrenia and Related Disorders

AURO-OLANZAPINE ODT (olanzapine) is indicated for the acute and maintenance treatment of schizophrenia and related psychotic disorders. In controlled clinical trials, olanzapine was found to improve both positive and negative symptoms.

Olanzapine has been shown to be effective in maintaining clinical improvement during 1-year of continuation therapy in patients who had shown an initial treatment response.

Bipolar Disorder

AURO-OLANZAPINE ODT (olanzapine) is indicated for the acute treatment of manic or mixed episodes in bipolar I disorder. Olanzapine may be used as monotherapy or cotherapy with agents commonly used in the treatment of acute bipolar disorder (e.g., lithium or divalproex sodium).

The efficacy of olanzapine as monotherapy maintenance treatment in bipolar patients with manic or mixed episodes who responded to acute treatment with olanzapine was demonstrated in two 1-year “time to relapse” trials (see Part II: CLINICAL TRIALS section).

The physician who elects to use AURO-OLANZAPINE ODT for extended periods should periodically re-evaluate the long-term usefulness of the drug for the individual patient (see DOSAGE AND ADMINISTRATION section).

Geriatrics (≥ 65 years): AURO-OLANZAPINE ODT is not indicated in elderly patients with dementia. See **WARNINGS AND PRECAUTIONS-Serious Warnings and Precautions Box** and **Special Populations**. Caution should be used when treating geriatric patients with AURO-OLANZAPINE ODT. See ACTION AND CLINICAL PHARMACOLOGY,

WARNINGS AND PRECAUTIONS, Special Populations, and DOSAGE AND ADMINISTRATION sections.

Pediatrics (< 18 years of age): The safety and efficacy of olanzapine have not been established in pediatric populations and its use is not recommended. See also WARNINGS and PRECAUTIONS, Pediatrics (< 18 years of age) and ADVERSE REACTIONS, Other Investigational Trials, Adverse Events in Adolescent Patients (ages 13-17 years).

CONTRAINDICATIONS

AURO-OLANZAPINE ODT (olanzapine) is contraindicated in those patients with a known hypersensitivity to the drug or the excipients of the product. For a complete listing, see DOSAGE FORMS, COMPOSITION and PACKAGING section.

WARNINGS AND PRECAUTIONS

SERIOUS WARNINGS AND PRECAUTIONS

Increased Mortality in Elderly Patients with Dementia:

Elderly patients with dementia treated with atypical antipsychotic drugs are at an increased risk of death compared to placebo. Analyses of thirteen placebo controlled trials with various atypical antipsychotics (modal duration of 10 weeks) in these patients showed a mean 1.6-fold increase in death rate in the drug-treated patients. Although the causes of death were varied, most of the deaths appeared to be either cardiovascular (e.g., heart failure, sudden death) or infectious (e.g., pneumonia) in nature. (See WARNINGS AND PRECAUTIONS, Special Populations, Use in Geriatric Patients with Dementia).

General

Neuroleptic Malignant Syndrome:

Neuroleptic malignant syndrome (NMS) is a potentially fatal symptom complex that has been reported in association with antipsychotic drugs, including olanzapine.

Clinical manifestations of NMS are hyperpyrexia, muscle rigidity, altered mental status, and evidence of autonomic instability (irregular pulse or blood pressure, tachycardia, diaphoresis, and cardiac dysrhythmia). Additional signs may include elevated creatine phosphokinase, myoglobinuria (rhabdomyolysis), and acute renal failure.

In arriving at a diagnosis, it is important to identify cases where the clinical presentation includes both serious medical illness (e.g., pneumonia, systemic infection, etc.), and untreated or inadequately treated extrapyramidal signs and symptoms (EPS). Other important considerations in the differential diagnosis include central anticholinergic toxicity, heat stroke, drug fever, and primary central nervous system pathology.

The management of NMS should include 1) immediate discontinuation of all antipsychotic drugs including AURO-OLANZAPINE ODT and other drugs not essential to therapy; 2) intensive symptomatic treatment and medical monitoring; and 3) treatment of any concomitant serious

medical problems for which specific treatments are available. There is no general agreement about specific pharmacological treatment for uncomplicated NMS.

If a patient requires antipsychotic drug treatment after recovery from NMS, the potential re-introduction of therapy should be very carefully considered. The patient should be carefully monitored since recurrences of NMS have been reported.

Weight Gain:
Olanzapine was associated with weight gain during clinical trials. Clinically significant weight gain was observed across all baseline body mass index (BMI) categories (see ADVERSE REACTIONS, Other Adverse Events Observed During Clinical Trials with Oral and Intramuscular Olanzapine Across All Indications, Weight Changes). Using pooled data from patients treated with olanzapine over the dosage range of 5 mg to 20 mg per day mean gain was 5.4 kg. The mean change in weight was comparable for patients with schizophrenia and bipolar mania. A retrospective analysis of 573 patients receiving olanzapine for up to 3 years found that dose was not a significant predictor of greater long-term changes in weight.

In long-term studies (at least 48 weeks), both the magnitude of weight gain and the proportion of olanzapine-treated patients who had clinically significant weight gain were greater than in short-term studies. The percentage of patients who gained ≥ 25% of their baseline body weight with long-term exposure was very common (≥ 10%).

Body Temperature Regulation:
Disruption of the body's ability to reduce core body temperature has been attributed to antipsychotic agents. Appropriate care is advised when prescribing AURO-OLANZAPINE ODT for patients who will be experiencing conditions which may contribute to an elevation of core temperature, e.g. exercising strenuously, exposure to extreme heat, receiving concomitant medication with anticholinergic activity, or being subject to dehydration.

Potential Effect on Cognitive and Motor Performance:
Because AURO-OLANZAPINE ODT may cause somnolence, patients should be cautioned about operating hazardous machinery, including motor vehicles, until they are reasonably certain that AURO-OLANZAPINE ODT therapy does not affect them adversely.

Falls:
AURO-OLANZAPINE ODT may cause somnolence, postural (orthostatic) hypotension, motor and sensory instability, which may lead to falls and, consequently, fractures or other injuries. For patients with diseases, conditions, or medications that could exacerbate these effects, complete fall risk assessments when initiating antipsychotic treatment and recurrently for patients on long-term antipsychotic therapy.

Carcinogenesis and Mutagenesis
For animal data, see Part II: TOXICOLOGY section.

Cardiovascular

Hypotension and Syncope:

As with other drugs that have high alpha-1 adrenergic receptor blocking activity, AURO-OLANZAPINE ODT may induce orthostatic hypotension, tachycardia, dizziness, and sometimes syncope, especially at the initiation of treatment. In a clinical trial database of 2500 patients treated with oral olanzapine, syncope was reported in 0.6% (15/2500). The risk of orthostatic hypotension and syncope may be minimized by initiating therapy with 5 mg QD (see DOSAGE AND ADMINISTRATION section). A more gradual titration to the target dose should be considered if hypotension occurs.

AURO-OLANZAPINE ODT should be used with particular caution in patients with known cardiovascular disease (history of myocardial infarction or ischemia, heart failure, or conduction abnormalities), cerebrovascular disease, and conditions which would predispose patients to hypotension (dehydration, hypovolemia, and treatment with antihypertensive medications).

Venous Thromboembolism:

Venous thromboembolism (VTE), including fatal pulmonary embolism, has been reported in temporal association with antipsychotic drugs, including olanzapine, in case reports and/or observational studies. When prescribing AURO-OLANZAPINE ODT, all potential risk factors for VTE should be identified and preventative measures undertaken, particularly since patients with schizophrenia often present with risk factors for VTE. Very rare cases of VTE have been reported in olanzapine-treated patients during the post-marketing period.

QT Interval:

In clinical trials, clinically meaningful QTc prolongations (Fridericia QT correction [QTcF] ≥ 500 milliseconds [msec] at any time post baseline in patients with baseline QTcF < 500 msec) were uncommon (0.1% to 1%) in patients treated with olanzapine, with no significant differences in associated cardiac events compared to placebo. However, as with other antipsychotics, caution should be exercised when olanzapine is prescribed with medicines known to increase QTc interval, especially in the elderly, in patients with congenital long QT syndrome, congestive heart failure, heart hypertrophy, hypokalaemia or hypomagnesaemia (see ADVERSE REACTIONS, Post-Market Adverse Drug Reactions).

Cardiac Death:

In a retrospective observational study, patients treated with atypical antipsychotics (including olanzapine) or typical antipsychotics had a similar dose-related increase of presumed sudden cardiac death (SCD) compared to non-users of antipsychotics (almost twice the risk than that for non-users). In postmarketing reports with olanzapine, the event of SCD has been reported very rarely.

Endocrine and Metabolism

Hyperglycaemia:

As with some other antipsychotics, exacerbation of pre-existing diabetes and hyperglycaemia have been reported rarely and diabetic ketoacidosis and diabetic coma including some fatal cases have been reported very rarely during the use of olanzapine, sometimes in patients with no reported history of hyperglycaemia (see ADVERSE REACTIONS; Post-Market Adverse Drug

Reactions section). In some cases, a prior increase in body weight has been reported which may be a predisposing factor. Patients should have baseline and periodic monitoring of blood glucose and body weight.

In clinical trials (up to 52 weeks) olanzapine was associated with a greater mean change in glucose relative to placebo. Treatment-emergent clinically significant changes in fasting glucose were observed in patients with or without evidence of glucose dysregulation at baseline (see ADVERSE REACTIONS, Other Adverse Events Observed During Clinical Trials with Oral and Intramuscular Olanzapine Across All Indications, Glucose Changes).

Assessment of the relationship between atypical antipsychotic use and glucose abnormalities is complicated by the possibility of an increased background risk of diabetes mellitus in patients with schizophrenia and the increasing incidence of diabetes mellitus in the general population. Given these confounders, the relationship between atypical antipsychotic use and hyperglycemia-related adverse events is not completely understood. However, epidemiological studies suggest an increased risk of treatment-emergent hyperglycemia-related adverse events in patients treated with the atypical antipsychotics. Precise risk estimates for hyperglycemia-related adverse events in patients treated with atypical antipsychotics are not available.

Any patient treated with atypical antipsychotics should be monitored for symptoms of hyperglycemia including polydipsia, polyuria, polyphagia, and weakness. Patients who develop symptoms of hyperglycemia during treatment with atypical antipsychotics should undergo fasting blood glucose testing. In some cases, hyperglycemia has resolved when the atypical antipsychotic was discontinued; however, some patients required continuation of anti-diabetic treatment despite discontinuation of the suspect drug. Patients with risk factors for diabetes mellitus (e.g., obesity, family history of diabetes) who are starting treatment with atypical antipsychotics should undergo fasting blood glucose testing at the beginning of treatment and periodically during treatment. Patients with an established diagnosis of diabetes mellitus who are started on atypical antipsychotics should be monitored regularly for worsening of glucose control.

Hyperprolactinemia:
As with other drugs that block dopamine D_2 and/or serotonin 5-HT_2 receptors, olanzapine may elevate prolactin levels. Elevations associated with olanzapine treatment are generally mild, and may decline during continued administration.

Since tissue culture experiments indicate that approximately one-third of human breast cancers are prolactin-dependent *in vitro*, AURO-OLANZAPINE ODT should only be administered to patients with previously detected breast cancer if the benefits outweigh the potential risks. Caution should also be exercised when considering AURO-OLANZAPINE ODT treatment in patients with pituitary tumours. Possible manifestations associated with elevated prolactin levels are amenorrhea, galactorrhea, and menorrhagia.

As is common with compounds which stimulate prolactin release, the administration of olanzapine resulted in an increase in the incidence of mammary neoplasms in both rats and mice.

The physiological differences between rats and humans with regard to prolactin make the clinical significance of these findings unclear. To date, neither clinical nor epidemiological studies have shown an association between chronic administration of these drugs and mammary tumorigenesis.

Long-standing hyperprolactinemia when associated with hypogonadism may lead to decreased bone mineral density in both female and male subjects.

Lipids:
Increases in lipids have been observed in olanzapine-treated patients in placebo-controlled clinical trials. Treatment-emergent clinically significant changes in fasting lipids were observed in patients with or without evidence of dyslipidemia at baseline (see ADVERSE REACTIONS; Other Adverse Events Observed During Clinical Trials with Oral and Intramuscular Olanzapine Across All Indications, Lipids subsection). Appropriate clinical monitoring is recommended, including baseline and follow-up lipid evaluations.

Gastrointestinal
Antiemetic Effect:
Consistent with its dopamine antagonist effects, olanzapine may have an antiemetic effect. Such an effect may mask signs of toxicity due to overdosage of other drugs, or may mask symptoms of disease such as brain tumour or intestinal obstruction.

Genitourinary
Priapism:
Rare cases of priapism have been reported with antipsychotic use, such as olanzapine. This adverse reaction, as with other psychotropic drugs, did not appear to be dose-dependent and did not correlate with the duration of treatment. The most likely mechanism of action of priapism is a relative decrease in sympathetic tone.

Urinary Retention:
Olanzapine possesses anticholinergic properties, which can lead to adverse drug reactions such as urinary retention. There have been several serious post-marketing reports of urinary retention in olanzapine-treated patients and in some cases, catheterization was required. AURO-OLANZAPINE ODT should be prescribed with caution in patients with a current diagnosis or prior history of urinary retention and in patients with other risk factors for urinary retention (e.g. benign prostatic hyperplasia). AURO-OLANZAPINE ODT should also be prescribed with caution in patients receiving medications with anticholinergic activity that can affect voiding.

Hematologic
Hematologic Indices:
In oral olanzapine clinical trials, there were no data to suggest olanzapine adversely affected bone marrow function, even in patients with a history of clozapine-associated neutropenia or leukopenia. Olanzapine was associated with a 5.7% incidence of mainly transient treatment-emergent elevations of eosinophil counts above the normal range. Elevations were not associated with any symptoms, identifiable allergic phenomena, or changes in other hematologic indices.

Rare cases of leukopenia have been reported with olanzapine. In case of symptoms of infection, WBC count and differential count should be considered.

Neutropenia, granulocytopenia and agranulocytosis have been reported during antipsychotic use. Therefore, it is recommended that patients have their complete blood count (CBC) tested prior to starting AURO-OLANZAPINE ODT and then periodically throughout treatment.

Hepatic

Aminotransferase Elevations:

During pre-marketing clinical trials, therapy with olanzapine was associated with elevation of hepatic aminotransferases, primarily ALT (SGPT). Within a clinical trial database of 2280 olanzapine-treated patients, with baseline ALT (SGPT) levels ≤ 60 IU/L, 5.9% (134/2280) had treatment-emergent ALT (SGPT) elevations to > 120 IU/L, 1.9% (44/2280) had elevations to > 200 IU/L, and 0.2% (5/2280) had elevations to > 400 IU/L. No patients had values in excess of 700 IU/L. None of the olanzapine-treated patients who had elevated aminotransferase values manifested clinical symptomatology associated with liver impairment. The majority of aminotransferase elevations were seen during the first six weeks of treatment. Most elevations were transient (66%) while patients continued on olanzapine therapy, or falling (11%) at the last available measurement. Of the 134 olanzapine-treated patients whose enzyme levels increased to > 120 IU/L, 20 discontinued treatment (6 for hepatic, 14 for other reasons) while their ALT (SGPT) values were still rising. In 38 olanzapine-treated patients with baseline ALT (SGPT) > 90 IU/L, none experienced an elevation to > 400 IU/L.

Rare post-marketing reports of hepatitis have been received. Very rare cases of cholestatic or mixed liver injury have also been reported in the post-marketing period. Hepatic failure, including fatalities has also been reported very rarely in the post-marketing period. See POST-MARKET ADVERSE DRUG REACTIONS section.

Precaution should be exercised when using AURO-OLANZAPINE ODT in patients with pre-existing hepatic disorders, in patients who are being treated with potentially hepatotoxic drugs, or if treatment-emergent signs or symptoms of hepatic impairment appear.

For patients who have known or suspected abnormal hepatic function prior to starting AURO-OLANZAPINE ODT, standard clinical assessment including measurement of aminotransferase levels is recommended. Periodic clinical reassessment with aminotransferase levels is recommended for such patients, as well as for patients who develop any signs and symptoms suggestive of a new onset liver disorder during AURO-OLANZAPINE ODT therapy.

Neurologic

Tardive Dyskinesia:

Tardive dyskinesia (TD), a syndrome consisting of potentially irreversible involuntary dyskinetic movements, is associated with the use of antipsychotic drugs. Tardive dyskinesia occurs more frequently in elderly patients; however, patients of any age can be affected. It is unknown whether antipsychotic drugs may differ in their potential to cause TD. However, during long-term, double-blind, extension schizophrenia maintenance trials (894 olanzapine-treated patients; median olanzapine treatment, 237 days), olanzapine was associated with a statistically

significantly lower incidence of treatment-emergent dyskinesia compared to haloperidol. During long-term, double-blind, monotherapy extension bipolar maintenance trials (567 olanzapine-treated patients, for up to 1 year), there were no cases of TD in the olanzapine arms, as determined by either reported adverse events or the Abnormal Involuntary Movement Scale (AIMS). TD has been reported very rarely (≤ 0.0025%) in post-market surveillance.

The risk of developing tardive dyskinesia and the chance of it becoming irreversible are believed to increase as the duration of treatment and the cumulative dose of antipsychotic drug increase. However, the syndrome can develop, although less commonly, after relatively brief periods of treatment at low doses. There is no known treatment for established cases of TD. The syndrome may remit, partially or completely, if antipsychotic drug treatment is withdrawn. Antipsychotic drug treatment itself, however, may suppress the signs and symptoms of tardive dyskinesia, thereby masking the underlying process.

Given these considerations, olanzapine should be prescribed in a manner that is most likely to minimize the risk of tardive dyskinesia. As with any antipsychotic drug, AURO-OLANZAPINE ODT should be reserved for patients who appear to be receiving substantial benefit from the drug. In such patients the lowest effective dose and the shortest duration of treatment should be sought. The need for continued treatment should be reassessed periodically.

If signs or symptoms of tardive dyskinesia appear in a patient on AURO-OLANZAPINE ODT, drug discontinuation should be considered. However, some patients may benefit from continued treatment with AURO-OLANZAPINE ODT despite the presence of the syndrome.

Seizures:
Conventional neuroleptics are known to lower seizure threshold. In clinical trials, seizures have occurred in a small number (0.9%, 22/2500) of olanzapine-treated patients. There were confounding factors that may have contributed to the occurrence of seizures in many of these cases. AURO-OLANZAPINE ODT should be used cautiously in patients who have a history of seizures or have conditions associated with seizures or have a lowered seizure threshold.

Psychiatric
Suicide:
The possibility of suicide or attempted suicide is inherent in psychosis, and thus close supervision and appropriate clinical management of high-risk patients should accompany drug therapy.

Renal
Uric Acid:
In the pre-marketing clinical trial database, olanzapine was associated with mild elevations of uric acid in some patients. However, only one olanzapine-treated patient experienced treatment-emergent gout, and the baseline uric acid concentration for this patient was at least as large as all concentrations observed while the patient was receiving olanzapine.

Skin
Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS):

Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS) has been reported with olanzapine exposure. DRESS consists of a combination of three or more of the following: cutaneous reaction (such as rash or exfoliative dermatitis), eosinophilia, fever, lymphadenopathy and one or more systemic complications such as hepatitis, nephritis, pneumonitis, myocarditis, and pericarditis. Severe cutaneous adverse reactions are sometimes fatal. Discontinue olanzapine if DRESS is suspected.

Special Populations

Pregnant Women:

There are no adequate and well-controlled studies in pregnant women. Patients should be advised to notify their physician if they become pregnant or intend to become pregnant during treatment with AURO-OLANZAPINE ODT. Because human experience in pregnant females is limited, this drug should be used in pregnancy only if the potential benefit justifies the potential risk to the fetus.

Non-Teratogenic Effects:

Neonates exposed to antipsychotic drugs (including AURO-OLANZAPINE ODT) during the third trimester of pregnancy are at risk for extrapyramidal and/or withdrawal symptoms following delivery. There have been reports of agitation, hypertonia, hypotonia, tremor, somnolence, respiratory distress and feeding disorder in these neonates. These complications have varied in severity; while in some cases symptoms have been self-limited, in other cases neonates have required intensive care unit support and prolonged hospitalization.

AURO-OLANZAPINE ODT should not be used during pregnancy unless the expected benefits to the mother markedly outweigh the potential risks to the fetus.

Labour and Delivery:

Parturition in rats was not affected by olanzapine. The effect of olanzapine on labour and delivery in humans is not known.

Nursing Women:

Lactation:

In a study in lactating, healthy women, olanzapine was excreted in breast milk. Mean infant exposure (mg/kg) at steady state was estimated to be 1.8% of the maternal olanzapine dose (mg/kg). Patients should be advised not to breast feed an infant if they are taking AURO-OLANZAPINE ODT.

Pediatrics (< *18* years of age):

The safety and efficacy of olanzapine in children under the age of 18 years have not been established and its use is not recommended.

Weight gain has been observed with atypical antipsychotic use in pediatric and adolescent patient populations. Independent of any changes during treatment with atypical antipsychotics, weight gain can be associated with adverse changes in other metabolic parameters (e.g., glucose and lipid metabolism). Abnormal childhood weight and metabolic status have been associated with adverse cardiovascular outcomes in adulthood. Weight gain and changes in other metabolic

parameters associated with atypical antipsychotics can be more frequent or more severe in pediatric and adolescent patients than in the adult patients.

The long-term safety of atypical antipsychotics, including potential cardiometabolic effects and effects on growth, maturation and behavioural development in patients under 18 years of age has not been systematically evaluated.

A greater magnitude of weight gain and lipid alterations has been reported in adolescents compared with adults. Adolescents treated with olanzapine experienced a significantly higher incidence of elevated prolactin levels and significantly higher mean increases in prolactin levels compared with adults. Hepatic aminotransferase elevations are more common in adolescents as compared to adults. Sedation-related events are more common in adolescents as compared to adults.

See also Adverse Reactions/Other Investigational Trials/Adverse Events in Adolescent Patients (ages 13-17 years).

Geriatrics (≥ *65* years of age):
The number of patients 65 years of age or over with schizophrenia or related disorders exposed to olanzapine during clinical trials was limited (N = 44). Caution should thus be exercised with the use of AURO-OLANZAPINE ODT in the elderly patient, recognizing the more frequent hepatic, renal, central nervous system, and cardiovascular dysfunctions, and more frequent use of concomitant medication in this population (see DOSAGE AND ADMINISTRATION section).

Use in Geriatric Patients with Dementia
Overall Mortality:
Elderly patients with dementia treated with atypical antipsychotic drugs showed increased mortality compared to placebo in a meta-analysis of 13 controlled trials of various atypical antipsychotic drugs. In five placebo-controlled trials with olanzapine in this population, the incidence of mortality was 3.5% for olanzapine-treated patients compared to 1.5% for placebo-treated patients. AURO-OLANZAPINE ODT is not indicated in elderly patients with dementia.

Dysphagia:
Esophageal dysmotility and aspiration have been associated with antipsychotic drug use. Aspiration pneumonia is a common cause of morbidity and mortality in patients with advanced Alzheimer's disease. Olanzapine and other antipsychotic drugs should be used cautiously in patients at risk for aspiration pneumonia.

Cerebrovascular Adverse Events (CVAEs), Including Stroke, in Elderly Patients with Dementia:
Cerebrovascular adverse events (e.g., stroke, transient ischemic attack), including fatalities, were reported in trials of olanzapine in elderly patients with dementia-related psychosis. In placebo-controlled studies, there was a higher incidence of CVAEs in patients treated with olanzapine compared to patients treated with placebo (1.3% vs. 0.4%, respectively; see ADVERSE REACTIONS section). AURO-OLANZAPINE ODT is not approved for the treatment of elderly patients with dementia.

There is insufficient evidence to determine whether CVAEs in elderly patients with dementia are associated specifically with olanzapine or all antipsychotic agents. Clinical trial data appear to suggest that patients with a dementia diagnosis of vascular or mixed type had a higher likelihood of experiencing CVAEs than other types of dementia.

The risks and benefits of the use of AURO-OLANZAPINE ODT in elderly patients with dementia should be assessed taking into account the risk predictors for CVAEs in the individual patient. Patients/caregivers should be advised to immediately report signs and symptoms of potential CVAEs, such as sudden weakness or numbness in the face, arms, or legs, and speech or vision problems.

Use in Patients with Other Concomitant Illness:
Clinical experience with olanzapine in patients with concomitant illness is limited. Caution is thus advised when using olanzapine in patients with diseases or conditions that could affect the metabolism or the pharmacodynamic activity of olanzapine (see DOSAGE AND ADMINISTRATION section and Part II: DETAILED PHARMACOLOGY).

Use in Patients with Cardiac Disorders:
Olanzapine has not been evaluated in patients with a recent history of myocardial infarction or unstable heart disease. Patients with these conditions were excluded from pre-marketing clinical trials.

Use in Patients with Diabetes and Risk Factors for Development of Diabetes:
As with some other antipsychotics, exacerbation of pre-existing diabetes and hyperglycaemia have been reported rarely, and diabetic ketoacidosis and diabetic coma including some fatal cases have been reported very rarely during the use of olanzapine, sometimes in patients with no reported history of hyperglycaemia (see ADVERSE REACTIONS, Post-Market Adverse Drug Reactions section). In some cases, a prior increase in body weight has been reported which may be a pre-disposing factor. Appropriate clinical monitoring is advisable in diabetic patients and in patients with risk factors for the development of diabetes mellitus.

Use in Patients with Renal and Hepatic Impairment:
Small single-dose clinical pharmacology studies (see Part II: DETAILED PHARMACOLOGY section) did not reveal any major alterations in olanzapine pharmacokinetics in subjects with renal or hepatic impairment. Given the limited clinical experience with olanzapine in patients with these conditions, caution should be exercised (see DOSAGE AND ADMINISTRATION section).

Other Concomitant Illnesses:
As olanzapine demonstrated anticholinergic activity *in vitro*, caution is advised when prescribing for patients with symptomatic prostatic enlargement, narrow-angle glaucoma or paralytic ileus and related conditions.

In clinical trials, a single case of pre-existing intracranial hypertension was exacerbated.

ADVERSE REACTIONS

The stated frequencies of adverse events represent the proportion of individuals who experienced at least once, a treatment-emergent adverse event of the type listed. An event was considered treatment-emergent if it occurred for the first time or worsened while receiving therapy following baseline evaluation. It is important to emphasize that although the events were reported during therapy, they were not necessarily caused by the therapy.

Clinical Trial Adverse Drug Reactions

The prescriber should be aware that the figures in the tables and tabulations cannot be used to predict the incidence of side effects in the course of usual medical practice where patient characteristics and other factors differ from those that prevailed in the clinical trials. Similarly, the cited frequencies cannot be compared with figures obtained from other clinical investigations involving different treatments, uses, and investigators. The figures cited, however, do provide the prescribing physician with some basis for estimating the relative contribution of drug and nondrug factors to the side effect incidence in the populations studied.

Incidence of Adverse Events Associated with Discontinuation:

Schizophrenia and Related Disorders:

In short-term, placebo-controlled trials, there was no statistically significant difference in rates of discontinuation of olanzapine or placebo attributed to adverse events. Overall, 5% of olanzapine-treated patients discontinued treatment for adverse events compared with 6% of placebo-treated patients. Discontinuations due to ALT (SGPT) elevations, however, were considered to be drug related (2% for olanzapine versus 0% for placebo) (see WARNINGS AND PRECAUTIONS, Renal subsection).

Bipolar Disorder:

Bipolar Mania

In short-term, placebo-controlled clinical trials, there was no difference overall in the incidence of discontinuation due to adverse events (2% for olanzapine versus 2% for placebo).

Bipolar Maintenance

In the long-term (1-year), placebo-controlled clinical trial, of the 225 olanzapine-treated patients, 16% (n = 35) discontinued due to an adverse event, compared with 9% (n = 12) of 136 placebo-treated patients.

In the long-term (1-year), active-controlled clinical trial, of the 217 olanzapine-treated patients, 19% (n = 41) discontinued due to an adverse event, compared with 26% (n = 55) of 214 lithium-treated patients.

All Short-Term Trials - Schizophrenia and Bipolar Mania Trials:

In short-term, active-controlled clinical trials, of the 1796 olanzapine-treated patients in comparative clinical trials with haloperidol, 98 (5%) discontinued treatment for adverse events compared with 66 of 810 (8%) haloperidol-treated patients.

All Short-Term Trials - Overall Integrated Safety Database:
In a pre-marketing clinical trial database of 2500 olanzapine-treated patients, 14.9% (372/2500) discontinued due to an adverse event. About half (183/372) of these discontinuations were associated with the underlying psychopathology. Other adverse events most commonly (incidence of 0.5% - 0.6%) reported as the reason for discontinuation among olanzapine-treated patients were: ALT (SGPT) increased, unintended pregnancy, creatine phosphokinase increased, and convulsion.

Incidence of Commonly Observed Adverse Events:
Schizophrenia and Related Disorders:
In the schizophrenia placebo-controlled trials, the most commonly observed adverse events associated with the use of olanzapine (incidence of ≥ 5% and at least twice placebo) were: dizziness (11% for olanzapine vs 4% for placebo), constipation (9% vs 3%), ALT (SGPT) increased (8% vs 3%), personality disorder (8% vs 4%), weight gain (6% vs 1%), akathisia (5% vs 1%), and postural hypotension (5% vs 2%).

Bipolar Disorder:
Bipolar Mania
In the bipolar mania monotherapy placebo-controlled trials, the most commonly observed adverse events associated with the use of olanzapine (incidence of ≥ 5% and at least twice placebo) were: somnolence (35% vs 13%), dry mouth (22% vs 7%), dizziness (18% vs 6%), asthenia (15% vs 6%), constipation (11% vs 5%), dyspepsia (11% vs 5%), increased appetite (6% vs 3%), and tremor (6% vs 3%).

In bipolar mania combination placebo-controlled trials, the most commonly observed adverse events associated with the combination of olanzapine and lithium or valproate (incidence of ≥ 5% and at least twice placebo) were: dry mouth (32% for olanzapine combination vs 9% for placebo), weight gain (26% vs 7%), increased appetite (24% vs 8%), dizziness (14% vs 7%), back pain (8% vs 4%), constipation (8% vs 4%), speech disorder (7% vs 1%), increased salivation (6% vs 2%), amnesia (5% vs 2%), and paresthesia (5% vs 2%). In addition to the latter list of adverse events identified during bipolar mania combination clinical trials tremor (≥ 10%) has also been identified.

Bipolar Maintenance
In the one-year ‘time to relapse’ placebo-controlled clinical trial in bipolar disorder, the most commonly observed adverse events associated with olanzapine (incidence of ≥ 5% and at least twice placebo) were: weight increased (8% for olanzapine vs 1.5% for placebo), headache NOS (6.7% vs 2.9%), fatigue (6.2% vs 1.5%), depression (5.8% vs 2.9%).

Other Indication Trials:
Abnormal gait and falls have been observed very commonly (≥ 10%) in clinical trials with elderly patients with dementia-related psychosis. Also, urinary incontinence and pneumonia were commonly reported (≥ 1% and < 10%) in these patients.

In clinical trials in patients with drug-induced (dopamine agonist) psychosis associated with Parkinson's disease, worsening of Parkinsonian symptomatology and hallucinations were reported very commonly and more frequently than with placebo.

Adverse Events Occurring at an Incidence of 1% or More Among Oral Olanzapine-Treated Patients:

Certain portions of the discussion below relating to objective or numeric safety parameters are derived from studies in patients with schizophrenia and have not been duplicated for bipolar disorder trials. However, this information is also generally applicable to bipolar disorder. Table 1 enumerates the incidence of treatment-emergent adverse events, rounded to the nearest percent, that occurred during acute therapy (up to 6 weeks) of schizophrenia in 1% or more of patients treated with olanzapine (doses ≥ 2.5 mg/day) where the incidence in patients treated with olanzapine was greater than the incidence in placebo-treated patients.

Table 1: Schizophrenia Trials: Treatment-Emergent Adverse Events Incidence in Placebo-Controlled Clinical Trials with Oral Olanzapine - Acute Phase[1]

	Percentage of Patients Reporting Event	
Body System/Adverse Event	**Olanzapine (N = 248)**	**Placebo (N = 118)**
Body As a Whole		
Headache	17%	15%
Pain	10%	9%
Fever	5%	3%
Abdominal pain	4%	2%
Back pain	4%	3%
Chest pain	4%	2%
Neck rigidity	2%	1%
Intentional injury	1%	0%
Cardiovascular System		
Postural hypotension	5%	2%
Tachycardia	4%	1%
Hypotension	2%	1%
Digestive System		
Constipation	9%	3%
Dry mouth	7%	4%
Gamma glutamyl transpeptidase increased	2%	1%
Increased appetite	2%	1%
Hemic and Lymphatic		
Leukopenia	1%	0%
Metabolic and Nutritional Disorders		
SGPT increased	8%	3%
Weight gain[2]	6%	1%
Edema	2%	0%

	Percentage of Patients Reporting Event	
Body System/Adverse Event	**Olanzapine (N = 248)**	**Placebo (N = 118)**
Peripheral edema	2%	0%
SGOT increased	2%	0%
Creatine phosphokinase increased	1%	0%
Musculoskeletal System		
Arthralgia	3%	2%
Joint disorder	2%	1%
Twitching	2%	1%
Nervous System		
Somnolence[2]	26%	15%
Agitation	23%	17%
Insomnia	20%	19%
Nervousness	16%	14%
Hostility	15%	14%
Dizziness[2]	11%	4%
Anxiety	9%	8%
Personality disorder	8%	4%
Akathisia[2]	5%	1%
Hypertonia	4%	3%
Speech disorder	4%	1%
Tremor	4%	3%
Amnesia	2%	0%
Drug dependence	2%	0%
Euphoria	2%	0%
Neurosis	1%	0%
Respiratory System		
Rhinitis	10%	6%
Cough increased	5%	3%
Pharyngitis	5%	3%
Skin and Appendages		
Fungal dermatitis	2%	0%
Vesiculobullous rash	2%	1%
Special Senses		
Amblyopia	5%	4%
Blepharitis	2%	1%
Eye disorder	2%	1%
Corneal lesion	1%	0%
Urogenital System		
Menstrual disorder[3]	2%	0%

[1] The following events had an incidence equal to or less than placebo: abnormal dreams, accidental injury, anorexia, apathy, asthenia, cogwheel rigidity, confusion, conjunctivitis, depression, diarrhea, dysmenorrhea[3], dyspepsia, ecchymosis, emotional lability, hallucinations, hyperkinesia, hypertension, hypokinesia, libido

increased, myalgia, nausea, paranoid reaction, paresthesia, pruritus, rash, schizophrenic reaction, sweating, thinking abnormal, tooth caries, vaginitis[3], vomiting.

[2] Statistically significantly more frequent in patients treated with olanzapine than in patients treated with placebo.

[3] Denominator used was for females only (N = 41 Olanzapine; N = 23 Placebo).

Other Adverse Events from Schizophrenia Trials:

Certain portions of the discussion below relating to objective or numeric safety parameters, namely, vital sign changes, weight gain, laboratory changes, and ECG changes are derived from studies in patients with schizophrenia and have not been duplicated for bipolar mania. However, this information is also generally applicable to bipolar mania.

Weight Changes:

During acute therapy (up to 6 weeks) in controlled clinical trials comparing olanzapine with placebo in the treatment of schizophrenia, the percentages of patients with weight gain ≥ 7% of baseline body weight at any time were 29% for olanzapine and 3% for placebo, which was a statistically significant difference. The average weight gain during acute therapy in patients treated with olanzapine was 2.8 kg. Clinically significant weight gain was observed across all baseline body mass index (BMI) categories. In long-term extension schizophrenia trials, there was an average gain of 5.4 kg, and 56% of olanzapine-treated patients with weight gain > 7% of baseline body weight. In long-term extension bipolar maintenance trials, there was a mean weight gain of 3.8 kg, and with 31% of olanzapine-treated patients with weight gain > 7% of baseline body weight (see WARNINGS AND PRECAUTIONS, Endocrine and Metabolism subsection).

Vital Sign Changes:

In placebo-controlled clinical trials, orthostatic hypotension (greater than 30 mm decrease in systolic blood pressure) occurred with an incidence of 5% in oral olanzapine-treated patients compared to 2% in placebo-treated patients (vital sign measurements collected only after 3-7 days of olanzapine treatment). Olanzapine was associated with a mean baseline to endpoint increase in heart rate of 2.4 beats per minute compared to no change among placebo-treated patients (see WARNINGS AND PRECAUTIONS, Cardiovascular subsection).

Laboratory Changes:

Olanzapine is associated with asymptomatic increases in SGPT, SGOT, and GGT (see WARNINGS AND PRECAUTIONS, Hepatic subsection). Olanzapine is also associated with generally mild increases in serum prolactin, which usually decreases with continued drug treatment. Olanzapine is also associated with asymptomatic elevations of eosinophils and uric acid (see WARNINGS AND PRECAUTIONS, Renal subsection), and with decreases in serum bicarbonate.

ECG Changes:

Between-group comparisons for pooled placebo-controlled trials revealed no statistically significant olanzapine/placebo differences in the proportions of patients experiencing potentially important changes in ECG parameters, including QT, QTc, and PR intervals.

Other Adverse Events Observed During Clinical Trials with Olanzapine Across All Indications

The following discussion relates primarily to weight gain, lipids, and glucose changes observed during clinical trials across all indications.

Weight Changes:

Weight gain has been very commonly observed in olanzapine-treated patients during clinical trials. In 13 placebo-controlled olanzapine monotherapy studies, 22.2% of olanzapine-treated patients gained ≥ 7% of their baseline body weight versus 3% of placebo-treated patients, with a median exposure to event of 8 weeks; 4.2% of olanzapine-treated patients gained ≥ 15% of their baseline weight versus 0.3% of placebo-treated patients, with a median exposure to event of 12 weeks. Clinically significant weight gain was observed across all baseline body mass index (BMI) categories.

In long-term studies (at least 48 weeks), both the magnitude of weight gain and the proportion of olanzapine-treated patients who had clinically significant weight gain were greater than in short-term studies. The percentage of patients who gained ≥ 25% of their baseline body weight with long-term exposure was very common (≥ 10%).

Lipids:

In placebo-controlled clinical trials of up to 12 weeks in duration, olanzapine-treated patients had a greater mean increase in fasting total cholesterol, LDL cholesterol, and triglycerides compared to placebo-treated patients.

Mean increases in fasting lipid values (total cholesterol, LDL cholesterol, and triglycerides) were greater in patients without evidence of lipid dysregulation at baseline. However, for mean changes in fasting triglycerides, the difference between olanzapine and placebo was greater in patients with evidence of lipid dysregulation at baseline. Elevations in fasting triglyceride levels ≥ 11.3 mmol/L were uncommonly observed with olanzapine use (8 week median duration of exposure).

Table 2: Changes in Fasting Lipids Values from Adult Placebo-Controlled Olanzapine Monotherapy Studies with Treatment Duration up to 12 Weeks

Laboratory Analyte	Olanzapine*	Placebo
Triglycerides: fasting normal to high (< 1.70 mmol/L to ≥ 2.26 mmol/L)	9.2% (N = 457)	4.4% (N = 251)
Triglycerides: fasting borderline to high (≥ 1.70 mmol/L and < 2.26 mmol/L to ≥ 2.26 mmol/L)	39.3% (N = 135)	20.0% (N = 65)
Cholesterol-Total: fasting normal to high (< 5.18 mmol/L to ≥ 6.22 mmol/L)	2.8% (N = 392)	2.4% (N = 207)
Cholesterol-Total: fasting borderline to high (≥ 5.18 mmol/L and < 6.22 mmol/L to ≥ 6.22 mmol/L)	23.0% (N = 222)	12.5% (N = 112)
LDL cholesterol: fasting normal to high (< 2.59 mmol/L to ≥ 4.14 mmol/L)	0% (N = 154)	1.2% (N = 82)
LDL cholesterol: fasting borderline to high (≥ 2.59 mmol/L and < 4.14 mmol/L to ≥ 4.14 mmol/L)	10.6% (N = 302)	8.1% (N = 173)

* Median duration of exposure 8 weeks.

For fasting HDL cholesterol, no statistically significant differences were observed between olanzapine-treated patients and placebo-treated patients (see WARNINGS AND PRECAUTIONS, Endocrine and Metabolism subsection).

The proportion of patients who had changes in total cholesterol, LDL cholesterol or triglycerides from normal or borderline to high, or changes in HDL cholesterol from normal or borderline to low, was greater in long-term studies (at least 48 weeks) as compared with short-term studies. Treatment-emergent clinically significant changes in fasting lipids were observed in patients with or without evidence of dyslipidemia at baseline.

Glucose Changes:
In clinical trials (up to 52 weeks) olanzapine was associated with a greater mean change in glucose relative to placebo.
The difference in mean changes between olanzapine and placebo was greater in patients with evidence of glucose dysregulation at baseline (including those patients diagnosed with diabetes mellitus or who met criteria suggestive of hyperglycemia), and these patients had a greater increase in HbA1c compared to placebo.

In patients with baseline normal fasting glucose levels (< 5.5 mmol/L), 2.2% (N = 543) of those treated with olanzapine (median exposure duration of 8 weeks) were found to have high glucose levels (≥ 6.99 mmol/L) during olanzapine treatment versus 3.4% (N = 293) of those treated with placebo. In patients with baseline borderline fasting glucose levels (≥ 5.5 mmol/L and < 6.99 mmol/L), 17.4% (N = 178) of those treated with olanzapine (median exposure duration of 5 weeks) were found to have high glucose levels (≥ 6.99 mmol/L) during olanzapine treatment versus 11.5% (N = 96) of those treated with placebo.

The proportion of patients who had a change in glucose level from normal or borderline at baseline to high increased over time. Treatment-emergent clinically significant changes in fasting glucose were observed in patients with or without evidence of glucose dysregulation at baseline. Glycosuria was commonly reported in olanzapine-treated patients during clinical trials.

Prolactin:
In controlled clinical trials (up to 12 weeks), elevations in prolactin were observed in 30% of olanzapine-treated patients as compared to 10.5% of placebo-treated patients. In the majority of these patients, the elevations were mild. In patients with schizophrenia, menstrual-related adverse events potentially associated with prolactin elevations[1] were common (< 10% to ≥ 1%), whereas sexual function-related and breast-related adverse events were infrequent (< 1% to ≥ 0.1%). In patients treated for other mental illnesses[2], sexual function-related adverse events (erectile dysfunction, libido decreased, loss of libido, orgasm abnormal) potentially associated with prolactin elevations were common (< 10% to ≥ 1%), whereas breast-related and menstrual-related adverse events were infrequent (< 1% to ≥ 0.1%).

Vital Sign Changes:

[1]*TEAEs analysis up to 52 weeks of treatment*
[2]*Bipolar Depression, Psychotic Depression, Borderline Personality Disorder and Bipolar Mania*

Bradycardia was uncommonly observed in clinical trials.

Photosensitivity Reactions:
Photosensitivity reactions were uncommonly observed in clinical trials.
Table 3 summarizes core adverse drug reaction terms and their frequencies identified from an integrated database of 42 completed olanzapine clinical studies in adults, consisting of 7787 patients exposed to olanzapine in placebo- or comparator-controlled clinical studies.

Table 3: Core Adverse Drug Reactions from Clinical Trials of Olanzapine

Body System/Adverse Reaction Term	**Frequency**				
	≥ 10%	**< 10% and ≥ 1%**	**< 1% and ≥ 0.1%**	**< 0.1% and ≥ 0.01%**	**< 0.01%**
Body as a Whole					
Pyrexia		X			
Cardiovascular					
[1]Orthostatic Hypotension	X				
Digestive System					
Abdominal Distension			X		
Musculoskeletal System					
Arthralgia		X			
Nervous System					
Amnesia			X		
Respiratory, Thoracic and Mediastinal Disorders					
Epistaxis			X		
Laboratory Analytes					
Clinical Chemistry					
[1]Alkaline phosphatase –Increased		X			
[1]Gamma Glutamyltransferase (GGT) (U/L) - High		X			
[1]Uric Acid (µmol/L) – High		X			
Hematology					
[1]Leukopenia, including Neutropenia		X			

[1] As assessed by measured values within the clinical trial database.

Dose-Dependent Adverse Events:
Dose-relatedness of adverse events was assessed using data from a clinical trial with a fixed dosage range. Table 4 enumerates the treatment-emergent adverse events in which there was a statistically significantly increasing dose response in this clinical trial.

Table 4: Schizophrenia Trials: Dose-Dependent Adverse Events in a Fixed Dosage Range, Placebo-Controlled Clinical Trial[1] of Olanzapine

	Percentage of Patients Reporting Event			
Body System/ Adverse Event	**Placebo (N = 68)**	**Olanzapine 5 ± 2.5 mg/day (N = 65)**	**Olanzapine 10 ± 2.5 mg/day (N = 64)**	**Olanzapine 15 ± 2.5 mg/day (N = 69)**
Digestive System				
Constipation	0%	6.2%	9.4%	14.5%
Nervous System				
Abnormal dreams	0%	0%	1.6%	4.3%

	Percentage of Patients Reporting Event			
Body System/ Adverse Event	**Placebo (N = 68)**	**Olanzapine 5 ± 2.5 mg/day (N = 65)**	**Olanzapine 10 ± 2.5 mg/day (N = 64)**	**Olanzapine 15 ± 2.5 mg/day (N = 69)**
Dizziness	2.9%	7.7%	9.4%	17.4%
Somnolence	16.2%	20.0%	29.7%	39.1%
Respiratory System				
Pharyngitis	1.5%	3.1%	1.6%	10.1%

[1] Fungal dermatitis was also reported with a statistically significantly increasing dose response, but is not included as a drug cause was remote.

Table 5 enumerates the treatment-emergent adverse events from one 8-week, randomized, double-blind, fixed-dose trial comparing 10 (N = 199), 20 (N = 200) and 40 (N = 200) mg/day of olanzapine in patients with schizophrenia or schizoaffective disorder. Statistically significant differences among the 3 dose groups were observed for the following safety outcomes: fatigue, dizziness, prolactin elevation, and weight gain (mean change).

Table 5: Schizophrenia Trial: Dose-Dependent Adverse Events in a Fixed Dose, Placebo-Controlled Clinical Trial of Olanzapine[1]

Adverse Event	**Olanzapine 10 mg/day (N = 195)**	**Olanzapine 20 mg/day (N = 191)**	**Olanzapine 40 mg/day (N = 197)**
Fatigue[2,3] (% reporting event)	1.5%	2.1%	6.6%
Dizziness[3] (% reporting event)	2.6%	1.6%	6.6%
Prolactin Elevation[2,3,4] (% reporting event)	31.2%	42.7%	61.1%
Prolactin Elevation[2,3] (mean change from baseline to endpoint)	-10.5 ng/mL	-1.7 ng/mL	4.9 ng/mL
Weight Gain ≥ 7% at any time (% reporting event)	14.0%	18.4%	20.5%
Weight Gain[2] (mean change from baseline to endpoint)	1.9 kg	2.3 kg	3.0 kg

[1] Study HGLF: 8-week, Phase IV, parallel, randomized, double-blind, fixed-dose study in patients with schizophrenia and schizoaffective disorder evaluating the dose-response efficacy and safety of olanzapine 10, 20, and 40 mg/day. Patients were titrated up to their randomized dose over 2 weeks.

[2] significant difference between 10 vs. 40 mg/day

[3] significant difference between 20 vs. 40 mg/day

[4] > 24.2 ng/mL (female) or > 18.77 ng/mL (male) at any time during the trial

Incidence of Treatment-Emergent Extrapyramidal Symptoms:

Table 6 enumerates the percentage of patients with treatment-emergent extrapyramidal symptoms as assessed by categorical analyses of formal rating scales during acute therapy in a controlled clinical trial comparing olanzapine at 3 fixed dosage ranges with placebo in the treatment of schizophrenia.

Table 6: Schizophrenia Trials: Treatment-Emergent Extrapyramidal Symptoms Assessed By Rating Scales Incidence In A Fixed Dosage Range, Placebo-Controlled Clinical Trial -- Acute Phase[1]

	Percentage of Patients			
	Placebo	**Olanzapine 5 ± 2.5 mg/day**	**Olanzapine 10 ± 2.5 mg/day**	**Olanzapine 15 ± 2.5 mg/day**
Parkinsonism[2]	15%	14%	12%	14%
Akathisia[3]	23%	16%	19%	27%

1 No statistically significant differences.
2 Percentage of patients with a Simpson-Angus Scale total score ≥ 3.
3 Percentage of patients with a Barnes Akathisia Scale global score ≥ 2.

Table 7 enumerates the percentage of patients with treatment-emergent extrapyramidal symptoms as assessed by spontaneously reported adverse events during acute therapy in the same controlled clinical trial comparing oral olanzapine at 3 fixed dosage ranges with placebo in the treatment of schizophrenia. Similar results were found during the long-term (up to 1-year) double-blind monotherapy extension bipolar maintenance trial comparing olanzapine with placebo; there was a higher statistical incidence of akathisia for combined doses of olanzapine versus placebo.

Table 7: Schizophrenia Trials: Treatment-Emergent Extrapyramidal Symptoms Assessed by Adverse Events Incidence in an Oral Fixed Dosage Range, Placebo-Controlled Clinical Trial --Acute Phase[1]

	Percentage of Patients Reporting Event			
Extrapyramidal Symptoms	**Placebo (N=68)**	**Olanzapine 5 ± 2.5 mg/day (N=65)**	**Olanzapine 10 ± 2.5 mg/day (N=64)**	**Olanzapine 15 ± 2.5 mg/day (N = 69)**
Dystonic events[2]	1%	3%	2%	3%
Parkinsonism events[3]	10%	8%	14%	20%
Akathisia events[4]	1%	5%	11%[1]	10%[1]
Dyskinetic events[5]	4%	0%	2%	1%
Residual events[6]	1%	2%	5%	1%
Any extrapyramidal event	16%	15%	25%	32%[1]

1 Statistically significantly different from placebo.
2 Patients with the following COSTART terms were counted in this category: dystonia, generalized spasm, neck rigidity, oculogyric crisis, opisthotonos, torticollis.
3 Patients with the following COSTART terms were counted in this category: akinesia, cogwheel rigidity, extrapyramidal syndrome, hypertonia, hypokinesia, masked facies, tremor.
4 Patients with the following COSTART terms were counted in this category: akathisia, hyperkinesia.
5 Patients with the following COSTART terms were counted in this category: buccoglossal syndrome, choreoathetosis, dyskinesia, tardive dyskinesia.
6 Patients with the following COSTART terms were counted in this category: movement disorder, myoclonus, twitching.

Other Investigational Trials

Cerebrovascular Adverse Events (CVAEs) in Elderly Patients with Dementia

Data from 5 placebo-controlled trials in elderly patients with dementia-related psychosis (Alzheimer's, vascular, and mixed; olanzapine n = 1178 and placebo n = 478) suggest that there

was a higher incidence of CVAE in patients treated with olanzapine compared to patients treated with placebo (1.3% vs. 0.4%, respectively). Although the incidence of CVAE was not significantly different when analyzed with Fisher's Exact Test (p = 0.177), the difference was found to be significant when simultaneously controlling for age, gender, and type of dementia using Poisson Regression (p = 0.0428). Four patients died in the olanzapine group versus 1 patient in the placebo group. In open-label safety trials studied for up to 59 weeks in dementia patients (N = 231), 7 cases of CVAEs, including 2 fatalities, were reported (see WARNINGS AND PRECAUTIONS section).

Data from these trials suggest that patients with a dementia diagnosis of vascular or mixed type had a 5-fold higher likelihood of experiencing CVAEs than patients with a diagnosis of Alzheimer's. There is insufficient information to determine whether CVAEs in elderly patients with dementia are associated specifically with olanzapine or all antipsychotic agents.

AURO-OLANZAPINE ODT is not approved for use in elderly patients with dementia.

Overall Mortality
Elderly patients with dementia treated with atypical antipsychotic drugs showed increased mortality compared to placebo in a meta-analysis of 13 controlled trials of various atypical antipsychotic drugs. In five placebo-controlled trials with olanzapine in this population, the incidence of mortality was 3.5 % for olanzapine-treated patients compared to 1.5% for placebo-treated patients. AURO-OLANZAPINE ODT is not indicated in elderly patients with dementia.

Laboratory Changes
In 5 double-blind, placebo controlled clinical trials of elderly patients with dementia-related psychosis (n = 1184 total in the olanzapine arms and n = 478 total in placebo arms), olanzapine-treated patients showed significantly greater incidence rates compared to placebo-treated patients of low albumin (10.4% vs 5.5%, respectively), low hemoglobin (4.2% vs 1.8%) and low hematocrit (4.6% vs 2.4%). Of patients who had low albumin values, 3.6% in the olanzapine-treated group vs 1.4% in the placebo-treated also experienced a treatment-emergent respiratory infection. A causal relationship between the two adverse events has not been determined.

Adverse Events in Adolescent Patients (ages 13-17 years)
The types of adverse events observed in adolescent patients treated with olanzapine were similar to those seen in adult patients. Although no clinical trials designed to compare adolescents to adults were conducted, the data from the adolescent trials were compared to those of the adult trials.

Mean increase in weight in adolescents (4.6 kg over 3 weeks median duration of exposure) was greater than in adults (2.6 kg over 7 weeks median duration of exposure).

In long-term studies (at least 24 weeks), both the magnitude of weight gain and the proportion of adolescent olanzapine-treated patients who had clinically significant weight gain were greater than in short-term studies and were greater than in adult patients with comparable exposures. With long-term exposure, approximately half of adolescent patients gained ≥ 15% and almost a

third gained ≥ 25% of their baseline body weight. Among adolescent patients, mean weight gain was greatest in patients who were overweight or obese at baseline.

Mean increases in fasting glucose were similar in adolescents and adults treated with olanzapine; however, the difference between olanzapine and placebo groups was greater in adolescents compared to adults.

In long-term studies (at least 24 weeks), changes in glucose from normal at baseline to high were uncommon (< 1% and ≥ 0.1%).

Mean increases in fasting total cholesterol, LDL cholesterol, and triglycerides were generally greater in adolescents than in adults treated with olanzapine. However, in short term trials, the differences between olanzapine and placebo were similar for adolescents and adults. The proportion of treatment-emergent clinically significant changes in normal-to-high or borderline-to-high fasting total cholesterol, LDL cholesterol and triglycerides was greater in adolescents compared to adults, and the differences between olanzapine and placebo in these categories of laboratory values were also generally greater in adolescents. In long-term studies, treatment-emergent clinically significant changes in total cholesterol, LDL cholesterol, and triglycerides were observed in adolescents with or without evidence of dyslipidemia at baseline.

Compared with adults, adolescents treated with olanzapine experienced a significantly higher incidence of elevated prolactin levels (47% in olanzapine-treated adolescents vs 30% in olanzapine-treated adults) and significantly higher mean increases in prolactin levels.

Hepatic aminotransferase elevations (3 times the Upper limit of Normal) are more common in adolescents (12.1%) as compared to adults (5.4%).

Sedation-related events are more common in adolescents (44%) as compared to adults (29%).

Table 8 summarizes core adverse drug reaction terms for olanzapine compared to placebo and their frequencies identified only during clinical trials in adolescent patients (ages 13 to 17 years). AURO-OLANZAPINE ODT is not indicated in adolescent patients (ages 13-17 years).

Table 8: Core Adverse Drug Reactions and Frequencies from Clinical Trials in Adolescent Patients (ages 13-17 years)

Body System/Adverse Reaction Term	Olanzapine		Placebo	
	Frequency	**N**	**Frequency**	**N**
Body as a Whole				
Weight gain ≥ 7% of baseline body weight (kg)[7]	40.6%	197	9.8%	112
Weight gain ≥ 15% of baseline body weight (kg)[8]	7.1%	197	2.7%	112
Digestive System				

Body System/Adverse Reaction Term	Olanzapine		Placebo	
	Frequency	**N**	**Frequency**	**N**
Dry Mouth	6.15%	179	0%	89
Increased Appetite	24%	179	6%	89
Nervous System				
Sedation[1]	44.1%	179	9%	89
Clinical Chemistry				
ALT/SGPT > 3X ULN all randomized patients with ALT baseline <= 3X ULN[2]	12.1%	174	2.3%	87
AST/SGOT - Increased[3]	27.6%	163	3.8%	79
Total bilirubin –Decreased[4]	22.1%	154	6.7%	75
GGT - Increased[5]	10.1%	169	1.2%	83
Prolactin - Increased[6]	47.4%	116	6.8%	59
Cholesterol – total, fasting normal to high (< 4.40 mmol/L to ≥ 5.18 mmol/L)[9]	6.9%	87	2.3%	43
Cholesterol – total, fasting borderline to high (≥ 4.40 mmol/L and < 5.18 mmol/L to ≥ 5.18 mmol/L)[9]	38.9 %	36	7.7%	13
LDL cholesterol: fasting normal to high (< 2.85 mmol/L to ≥ 3.37 mmol/L)	5.1%	98	4.5%	44
LDL cholesterol: fasting borderline to high (≥ 2.85 mmol/L and < 3.37 mmol/L to ≥ 3.37 mmol/L)	48.3%	29	0%	9
Triglycerides, fasting normal to high (< 1.02 mmol/L to > 1.47 mmol/L)[9]	26.9%	67	10.7%	28
Triglycerides, fasting borderline to high (≥ 1.02 mmol/L and ≤ 1.47 mmol/L to > 1.47 mmol/L)[9]	59.5%	37	35.3%	17
Glucose, fasting normal to high (< 5.55 mmol/L to ≥ 6.99 mmol/L)[9]	0%	124	1.9%	53
Glucose, fasting borderline to high (≥ 5.55 mmol/L and < 6.99 mmol/L to ≥ 6.99 mmol/L)[10]	14.3%	14	0%	13

[1] Represented cluster of MedDRA terms including: hypersomnia, lethargy, sedation, somnolence.

[2] Covance reference ranges:

		Low	High
(U/L)	Female 13 - ≤ 17.999	6	34
(U/L)	Male 13 - ≤ 17.999	6	43

[3] Covance reference ranges:

		Low	High
(U/L)	Female 13 - ≤ 17.999	10	40
(U/L)	Male 13 - ≤ 17.999	10	40

[4] Covance reference ranges:

		Low	High
(mmol/L)	Female 13 -≤ 17.999	3	21
(mmol/L)	Male 13 -≤ 17.999	3	21

[5] Covance reference ranges

		Low	High

(U/L)	Female 13 - ≤ 17.999	0	33
(U/L)	Male 13 - ≤ 17.999	0	51

[6] Covance reference ranges for prolactin as published by Wiedemann and Jonetz-Mentzel (1993)

Female:	12 to 14 years:	2.52 – 16.90 ng/mL
	14 to 19 years:	4.20 – 39.00 ng/mL
Male:	12 to 14 years:	2.84 – 24.00 ng/mL
	14 to 19 years:	2.76 – 16.10 ng/mL

[7] Median duration of exposure to event = 4 weeks
[8] Median duration of exposure to event = 19 weeks
[9] Median duration of exposure was 3 weeks
[10] Median duration of exposure was 5 weeks

Post-Market Adverse Drug Reactions

Table 9 summarizes core adverse drug reaction terms and their frequencies identified from global post-marketing surveillance in addition to what was reported in clinical trials (*see* preceding section ADVERSE REACTIONS, Clinical Trial Adverse Drug Reactions). A causal relationship between olanzapine and the emergence of these events has not been established.

Table 9: Core Adverse Drug Reactions Seen with Olanzapine Formulations[1]

Body System/Adverse Reaction Term	**Frequency**				
	≥ 10%	**<10% and ≥ 1%**	**< 1% and ≥ 0.1%**	**< 0.1% and ≥ 0.01%**	**< 0.01%**
Body as a Whole					
Allergic reaction [2]					X
Discontinuation reaction [3]					X
Cardiovascular					
Venous Thromboembolism, including Pulmonary Embolism and Deep Vein Thrombosis					X
Digestive System					
Pancreatitis					X
Salivary Hypersecretion[8]			X		
Hematologic					
Thrombocytopenia [4]					X
Hepatobiliary disorders					
Hepatitis				X	
Jaundice					X
Hepatic failure					X
Metabolic					
Diabetic Coma					X
Diabetic Ketoacidosis [5]					X
Hypercholesterolemia [7]					X
Hyperglycaemia				X	
Hypertriglyceridemia[6, 7]					X

Body System/Adverse Reaction Term	Frequency				
	≥ 10%	**<10% and ≥ 1%**	**< 1% and ≥ 0.1%**	**< 0.1% and ≥ 0.01%**	**< 0.01%**
Exacerbation of pre-existing diabetes				X	
Musculoskeletal System					
Rhabdomyolysis					X
Nervous System					
Restless Legs Syndrome (RLS)[8]			X		
Seizures				X	
Stuttering[1, 9]			X		
Skin and Appendages					
Alopecia					X
Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS)					X
Rash				X	
Urogenital System					
Priapism					X
Urinary Incontinence					X
Urinary Retention					X
Laboratory Analytes					
Clinical Chemistry					
Total bilirubin –Increased					X

1 Adverse event identified from spontaneous post-marketing surveillance.
2 e.g., maculopapular rash, anaphylactoid reaction, angioedema, pruritus, or urticaria.
3 i.e., diaphoresis, nausea or vomiting.
4 Including a case of thrombocytopenic purpura.
5 COSTART term is diabetic acidosis.
6 COSTART term is hyperlipemia.
7 Random cholesterol levels of ≥ 6.22 mmol/L and random triglyceride levels of ≥ 11.30 mmol/L have been very rarely reported.
8 Adverse event identified from spontaneous post-marketing reporting with frequency determined using the olanzapine clinical trial database.
9 Stuttering was only studied in oral formulations and the review did not include details about the rapid IM formulation.

As with other atypical anti-psychotics, there have been isolated post-market reports with olanzapine of serious cardiovascular-related adverse events, including fatalities (see WARNINGS AND PRECAUTIONS, Cardiovascular).

Neutropenia, granulocytopenia and agranulocytosis have been reported during antipsychotic use. Therefore, it is recommended that patients have their complete blood count (CBC) tested prior to starting AURO-OLANZAPINE ODT and then periodically throughout treatment.

Venous Thromboembolism (VTE), including fatal pulmonary embolism, has been reported with antipsychotic drugs, including olanzapine. However, since patients who require treatment with

antipsychotics often present with acquired risk factors for VTE all possible risk factors of VTE e.g. immobilization, should be identified and preventative measures undertaken.

Patients should be advised of the risk of severe constipation during AURO-OLANZAPINE ODT treatment, and that they should tell their doctor if constipation occurs or worsens, as they may need laxatives.

Atypical antipsychotic drugs, including olanzapine, have been associated with cases of sleep apnea, with or without concomitant weight gain. In patients who have a history of or are at risk for sleep apnea, AURO-OLANZAPINE ODT should be prescribed with caution.

Risks of somnambulism (sleep walking) and sleep-related eating disorder have been associated with the use of atypical antipsychotics including olanzapine.

DRUG INTERACTIONS

Drug-Drug Interactions

Alcohol: Given the primary CNS effects of AURO-OLANZAPINE ODT, caution should be used when it is taken in combination with other centrally acting drugs and alcohol since additive pharmacological effects such as increased sedation may occur.
Levodopa and Dopamine Agonists: As it exhibits *in vitro* dopamine antagonism, olanzapine may antagonize the effects of levodopa and dopamine agonists.

Antihypertensive Agents: Because of its potential for inducing hypotension, olanzapine may enhance the effects of certain antihypertensive agents. Caution should be exercised in patients who receive medicinal products that can induce hypotension, bradycardia, or respiratory depression.

Potential for Other Drugs to Affect AURO-OLANZAPINE ODT:
Carbamazepine: Concomitant carbamazepine therapy may induce the metabolism of olanzapine.

Activated Charcoal: The concomitant administration of activated charcoal reduced the oral bioavailability of olanzapine by 50% to 60%.

Antacids: Single doses of antacid (aluminium, magnesium) or cimetidine did not affect the oral bioavailability of olanzapine.

Valproate: Studies *in vitro* using human liver microsomes showed that olanzapine has little potential to inhibit the glucuronidation of valproate, which is the major metabolic pathway. Furthermore, valproate was found to have little effect on the metabolism of olanzapine *in vitro*. Daily concomitant *in vivo* administration of 10 mg olanzapine for 2 weeks did not affect steady state plasma concentrations of valproate. Therefore, concomitant olanzapine administration does not require dosage adjustment of valproate.

Fluoxetine: Fluoxetine (60 mg single dose or 60 mg daily for 8 days) causes a mean 16% increase in the maximum concentration of olanzapine and a mean 16% decrease in olanzapine clearance. The magnitude of the impact of this factor is small in comparison to the overall variability between individuals, and therefore dose modification is not routinely recommended.

CYP1A2 Inducers: Agents that induce CYP1A2 such as omeprazole may increase clearance of olanzapine.

CYP1A2 Inhibitors: Fluvoxamine, a specific CYP1A2 inhibitor, has been shown to significantly inhibit the metabolism of olanzapine. The mean increase in olanzapine C_{max} following fluvoxamine was 54% in female nonsmokers and 77% in male smokers. The mean increase in olanzapine AUC was 52% and 108%, respectively. A lower starting dose of olanzapine should be considered in patients who are using fluvoxamine or any other CYP1A2 inhibitors, such as ciprofloxacin or ketoconazole. A decrease in the dose of olanzapine should be considered if treatment with an inhibitor of CYP1A2 is initiated.

Potential for AURO-OLANZAPINE ODT to Affect Other Drugs:
Theophylline: The pharmacokinetics of theophylline, a drug principally metabolized by CYP1A2, were not altered by olanzapine in a clinical trial with single doses of IV theophylline.

Imipramine/Desipramine: In clinical trials with single doses of olanzapine, no inhibition of the metabolism of imipramine/desipramine (P450-CYP2D6) was evident.

Warfarin: In clinical trials with single doses of olanzapine, no inhibition of the metabolism of warfarin (P450 CYP2C9) was evident.

Diazepam: In clinical trials with single doses of olanzapine, no inhibition of the metabolism of diazepam (P450 CYP3A4) was evident.

Lithium or Biperiden: Olanzapine showed no interaction when coadministered with lithium or biperiden.

Drugs Metabolized via P450-CYP1A2, -CYP2C9, -CYP2C19, -CYP2D6, and -CYP3A: In *in vitro* studies with human microsomes, olanzapine showed little potential to inhibit cytochromes P450-CYP1A2, -CYP2C9, -CYP2C19, -CYP2D6, and -CYP3A (see PART II: DETAILED PHARMACOLOGY). Olanzapine is thus unlikely to cause clinically important drug-drug interactions mediated through the metabolic routes outlined above. However, the possibility that olanzapine may alter the metabolism of other drugs, or that other drugs may alter the metabolism of olanzapine, should be considered when prescribing AURO-OLANZAPINE ODT.

Phenylketonurics: AURO-OLANZAPINE ODT contains phenylalanine (0.28, 0.56, 0.84, or 1.12 mg per 5, 10, 15, or 20 mg orally disintegrating tablet, respectively).

Drug-Food Interactions
Absorption of olanzapine is not affected by food.

Drug-Herb Interactions
Interactions with herbal products have not been identified.

Drug-Laboratory Interactions
Interactions with laboratory tests have not been identified.

Drug-Lifestyle Interactions
Smoking: Concomitant smoking may induce the metabolism of olanzapine.

DOSAGE AND ADMINISTRATION

Schizophrenia and Related Disorders
Adults: AURO-OLANZAPINE ODT (olanzapine) should be administered on a once-a-day schedule without regard to meals, generally beginning with 5 to 10 mg, with a target dose of 10 mg/day within several days. Further dosage adjustments, if indicated, should generally occur at intervals of not less than 1 week, since steady state for AURO-OLANZAPINE ODT would not be achieved for approximately 1 week in the typical patient. When dosage adjustments are necessary, dose increments/ decrements of 5 mg per day are recommended. An increase to a dose greater than target dose of 10 mg/day (i.e., to a dose of 15 mg/day or greater) is normally recommended only after clinical assessment.

In clinical trials a dose range of 5-20 mg/day was studied (see Part II: CLINICAL TRIALS).

Doses above 20 mg/day have been evaluated from a safety perspective (see Table 6 in Adverse Events, Dose-Dependent Adverse Events subsection); however, efficacy at doses above 20 mg/day has not been systematically evaluated.

Maintenance Therapy in Schizophrenia:
It is recommended that responding patients with schizophrenia be continued on AURO-OLANZAPINE ODT at the lowest dose needed to maintain remission. Patients should be reassessed periodically to determine the need for maintenance treatment. While there is no body of evidence available to answer the question of how long the patient should be treated with AURO-OLANZAPINE ODT, the effectiveness of maintenance treatment is well established for many other antipsychotic drugs.

Bipolar Disorder
Bipolar Mania
Adults: The recommended starting dose for olanzapine is 15 mg administered once a day in monotherapy and 10 mg daily in combination therapy.

It may be given without regard to meals as its absorption is not affected by food. The dosage range of olanzapine is from 5 mg to 20 mg per day. Daily dosage should be adjusted in response to clinical assessment.

Maintenance Therapy in Bipolar Disorder:
Patients who have been receiving and responding to AURO-OLANZAPINE ODT for the treatment of acute manic or mixed episodes of bipolar disorder should initially continue maintenance therapy at the same dose (see Part II: CLINICAL TRIALS). Subsequent daily dosage should be adjusted on the basis of clinical status within a range of 5-20 mg per day.

Patients should be periodically reassessed to determine the need for maintenance treatment and the appropriate dose for such treatment.

General Considerations for Oral Dosing in Special Populations
The Elderly or Debilitated Patient:
In clinical trials, 44 patients with schizophrenia or related disorders who were 65 years of age or over were treated with olanzapine (5-20 mg daily) (see WARNINGS AND PRECAUTIONS, Special Populations). Given the limited experience with olanzapine in the elderly, and the higher incidence of concomitant illness and concomitant medication in this population, AURO-OLANZAPINE ODT should be used with caution.

The recommended starting dose is 5 mg in patients who are elderly, debilitated, who have a predisposition to hypotensive reactions, who otherwise exhibit a combination of factors that may result in slower metabolism of AURO-OLANZAPINE ODT (e.g., nonsmoking female patients), or who may be pharmacodynamically more sensitive to AURO-OLANZAPINE ODT. When indicated, dose escalation should be performed with caution in these patients.

Patients with Hepatic and/or Renal Impairment:
As clinical experience is lacking in these patients, the lower initial starting dose and slower titration to initial target dose should be considered. Further dose escalation, when indicated, should be conservative (see WARNINGS AND PRECAUTIONS, Special Populations).

Missed Dose
If a patient misses a dose by a few hours, advise patient to take as soon as he/she remembers. If most of the day has passed, advise patient to wait until the next scheduled dose. Advise patients to not take 2 doses of AURO-OLANZAPINE ODT at once.

Administration of AURO-OLANZAPINE ODT
AURO-OLANZAPINE ODT orally disintegrating tablet is intended for oral administration only. It begins disintegrating in the mouth within seconds, allowing its contents to be subsequently swallowed with or without liquid.

The orally disintegrating tablet breaks easily and should be handled carefully, with dry hands. Direct contact with hands should be avoided if possible. When ready for administration, the orally disintegrating tablet should be carefully pushed out from the bottom of the blister cell and placed directly in the mouth. The orally disintegrating tablet may also be stirred into 125 mL (4 ounces) of water, milk, coffee, orange juice or apple juice and the contents promptly consumed.

OVERDOSAGE

Signs and Symptoms
Very common symptoms reported in olanzapine overdose (≥ 10% incidence) include tachycardia, agitation/aggressiveness, dysarthria, various extrapyramidal symptoms, and reduced level of consciousness ranging from sedation to coma.

Other medically significant sequelae of olanzapine overdose include delirium, convulsion, possible neuroleptic malignant syndrome, respiratory depression, aspiration, hypertension or hypotension, cardiac arrhythmias (< 2% of overdose cases) and cardiopulmonary arrest. Fatal outcomes have been reported for acute overdoses as low as 450 mg of oral olanzapine but survival has also been reported following acute overdose of approximately 2,000 mg of oral olanzapine.

Management of Overdose
There is no specific antidote for olanzapine. Induction of emesis is not recommended. Standard procedures for management of overdose may be indicated (i.e., gastric lavage, administration of activated charcoal). The concomitant administration of activated charcoal was shown to reduce the oral bioavailability of olanzapine by 50 to 60%.

Symptomatic treatment and monitoring of vital organ function should be instituted according to clinical presentation, including treatment of hypotension and circulatory collapse and support of respiratory function. Do not use epinephrine, dopamine, or other sympathomimetic agents with beta-agonist activity since beta stimulation may worsen hypotension.

For management of a suspected drug overdose, contact your regional Poison Control Centre immediately.

ACTION AND CLINICAL PHARMACOLOGY

Pharmacodynamics

Pharmacodynamic Properties:
Olanzapine, a thienobenzodiazepine, is an antipsychotic agent that demonstrates a broad pharmacologic profile across a number of receptor systems. Olanzapine displays high receptor affinity binding *in vitro* at dopamine D_2, D_3, D_4 (Ki = 11-31 nM), 5-$HT_{2A/C}$ (Ki = 4 and 11 nM, respectively), 5-HT_3, 5-HT_6, muscarinic M_1-M_5 (Ki = 1.9-2.5 nM), adrenergic α_1 (Ki = 19 nM), and histamine H_1 (Ki = 7 nM) receptor subtypes, while displaying a lower affinity at dopamine D1 and D5 receptor subtypes (Ki = 51–119 nM). In a behavioural paradigm predictive of antipsychotic activity, olanzapine reduced conditioned avoidance response in rats at doses lower than 4 times those required to produce catalepsy. In a single dose (10 mg) PET study in healthy subjects, olanzapine produced higher 5-HT_{2A} than dopamine D_2 receptor occupancy. The percent of D_2 occupancy was less than the threshold value predictive of extrapyramidal events.

In animals olanzapine has been observed to produce a significant reduction in the firing of A10 dopaminergic cells. The number of spontaneously active A9 neurons either remained constant or was increased. This may explain the low incidence of extrapyramidal side effects with olanzapine usually associated with the typical antipsychotics.

Olanzapine also increases extracellular levels of dopamine in a regionally specific manner in the prefrontal cortex, similar to mood stabilizers, lithium and valproate.

Pharmacokinetics

Absorption: Olanzapine is well absorbed after oral administration, reaching peak plasma concentrations within 5 to 8 hours. The absorption is not affected by food.

Distribution: Plasma concentrations of orally administered olanzapine were linear and dose proportional in trials studying doses from 1 to 20 mg. The maximum plasma concentrations (C_{max}) of olanzapine after single oral doses of 5, 10 and 15 mg averaged 7, 14, and 21 ng/mL, respectively (20 ng/mL = 0.064 μM). In young healthy volunteers, after once-a-day repeated dosing, steady-state C_{max} was approximately twice that achieved after a single dose (e.g. 23 ng/mL versus 12 ng/mL for a 10-mg dose). In the elderly, the steady state plasma concentration was approximately 3-fold higher than that achieved after a single dose (e.g. 16 ng/mL versus 5 ng/mL for a 5-mg dose). In both, young and elderly, steady-state concentrations of olanzapine were obtained after seven days of once daily dosing.

Over time and dosage range, pharmacokinetic parameters within an individual are very consistent. However, plasma concentrations, half-life and clearance of olanzapine may vary between individuals on the basis of smoking status, gender, and age (see Special Populations). Data from pooled, single dose pharmacokinetic studies showed the half-life of olanzapine to range from 21 to 54 hours (5^{th} to 95^{th} percentile), and the apparent plasma clearance to range from 12 to 47 L/hr (5^{th} to 95^{th} percentile).

The plasma protein binding of olanzapine was about 93% over the concentration range of about 7 to about 1000 ng/mL. Olanzapine is bound predominantly to albumin and α_1-acid glycoprotein.

Metabolism: Olanzapine is metabolized in the liver by conjugative and oxidative pathways. The major circulating metabolite is the 10-N-glucuronide, which is pharmacologically inactive and does not pass the blood brain barrier. Cytochrome P450 isoforms CYP1A2 and CYP2D6 contribute to the formation of the N-desmethyl and 2-hydroxymethyl metabolites. Both metabolites exhibited significantly less *in vivo* pharmacological activity than olanzapine in animal studies. The predominant pharmacologic activity is from the parent olanzapine.

In vitro microsomal studies show that olanzapine is a weak inhibitor of CYP1A2 (Ki = 36 μM), CYP2D6 (Ki = 89 μM), and CYP3A4 (Ki = 490 μM). Based upon these Ki values, little inhibition of these cytochrome P-450 enzymes is expected *in vivo* at concentrations below 5 μM (roughly 1500 ng/mL) because the olanzapine concentration will be less than 10% of its Ki value. In clinical studies, observed steady-state plasma concentrations of olanzapine are rarely > 150 ng/mL (approximately 0.5 μM). Olanzapine is thus not likely to cause clinically important

pharmacokinetic drug-drug interactions mediated through the metabolic routes outlined above. (See DRUG INTERACTIONS section).

Elimination: After oral administration to healthy subjects, the mean terminal elimination half-life was 33 hours (21 to 54 hours for 5th to 95th percentile) and the mean olanzapine plasma clearance was 26 L/hr (12 to 47 L/hr for the 5th to 95th percentile). Olanzapine pharmacokinetics varied on the basis of smoking status, gender, and age. Table 10 summarizes these effects:

Table 10: Olanzapine Key Pharmacokinetics

Patient Characteristics	Half-Life (hours)	Plasma Clearance (L/hr)
Nonsmoking	38.6	18.6
Smoking	30.4	27.7
Female	36.7	18.9
Male	32.3	27.3
Elderly (65 and older)	51.8	17.5
Non-elderly	33.8	18.2

Although smoking status, gender, and, to a lesser extent, age may affect olanzapine clearance and half-life, the magnitude of the impact of these single factors is small in comparison to the overall variability between individuals.

Special Populations and Conditions

Geriatrics: In a study involving 24 healthy subjects, the mean elimination half-life of olanzapine was about 1.5 times greater in elderly (> 65 years) than in non-elderly subjects (≤ 65 years). Caution should be used in dosing the elderly, especially if there are other factors that might additively influence drug metabolism and/or pharmacodynamic sensitivity (see DOSAGE AND ADMINISTRATION section).

Gender: Clearance of olanzapine is approximately 30% lower in women than in men. There were, however, no apparent differences between men and women in effectiveness or adverse effects. Dosage modifications based on gender should not be needed.

Race: In a study of Caucasians, Japanese, and Chinese subjects, there were no differences in olanzapine pharmacokinetics among the three populations. Cytochrome P450 isoform CYP2D6 status does not affect the metabolism of olanzapine.

Hepatic Insufficiency: No differences in the single-dose pharmacokinetics of oral olanzapine were noted in subjects with clinically significant cirrhosis (who were mostly smokers) when compared to healthy subjects (all non-smokers). Multiple-dose studies in patients with hepatic impairment, however, have not been performed.

Renal Insufficiency: There was no significant difference in mean elimination half-life or olanzapine plasma clearance between subjects with severely impaired renal function compared to

individuals with normal renal function. Approximately 57% of radio-labelled olanzapine is excreted in urine, principally as metabolites.

STORAGE AND STABILITY

Store in the original package at room temperature (15°C to 30°C). Protect from light and moisture.

DOSAGE FORMS, COMPOSITION AND PACKAGING

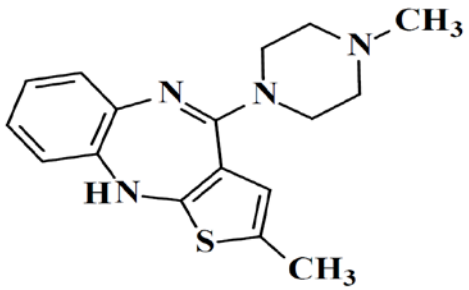
Dosage form	Orally Disintegrating Tablets			
Strength	5 mg	10 mg	15 mg	20 mg
Description	Yellow coloured, circular, flat faced beveled edge tablets debossed with ‘C’ on one side and ‘51’ on the other side.	Yellow coloured, circular, flat faced beveled edge tablets debossed with ‘C’ on one side and ‘52’ on the other side.	Yellow coloured, circular, flat faced beveled edge tablets debossed with ‘C’ on one side and ‘53’ on the other side.	Yellow coloured, circular, flat faced beveled edge tablets debossed with ‘C’ on one side and ‘54’ on the other side.
Composition	5 mg olanzapine	10 mg olanzapine	15 mg olanzapine	20 mg olanzapine
	Non-Medicinal Ingredients: Aspartame, mannitol, polacrilin potassium, crospovidone, colloidal silica anhydrous, microcrystalline cellulose, sodium stearyl fumarate and artificial pineapple flavor.			
Packaging	Blister packs of 30’s	Blister packs of 30’s	Blister packs of 30’s	Blister packs of 30’s

AURO-OLANZAPINE ODT meets USP Disintegration Test 2.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

Proper Name:	Olanzapine
Chemical Name:	2-Methyl-4-(4-methyl-1-piperazinyl)-10*H*-thieno[2,3-*b*] [1,5] benzodiazepine
Molecular Formula and Molecular Weight:	$C_{17}H_{20}N_4S$ 312.4 g/mol
Structural Formula:	
Physicochemical properties:	
Description:	Olanzapine is an antipsychotic agent of the thienobenzodiazepine class. It is a yellow crystalline powder, which is Freely soluble in methylene chloride, slightly soluble in ethanol (96 percent), practically insoluble in water.

CLINICAL TRIALS

<u>Comparative Bio-Availability Study</u>

A double blind, randomized, two-treatment, two-sequence, two-period, crossover, balanced single-dose comparative oral bioavailability study of AURO-OLANZAPINE ODT (Olanzapine) Orally Disintegrating Tablets, 10 mg (Test) of Aurobindo Pharma Limited, India manufactured for Auro Pharma Inc., and ZYPREXA® ZYDIS® (Olanzapine) Orally Disintegrating Tablets, 10 mg (Reference) of Eli Lilly Canada Inc. was conducted in 36 healthy, adult, male subjects under fasting conditions.

Summary Table of the Comparative Bio-Availability Data

Olanzapine (1 X 10 mg) From measured data Geometric Mean Arithmetic Mean (CV %)				
Parameter	**Test***	**Reference †**	**% Ratio of Geometric Means**	**90% Confidence Interval**
$AUC_{0\rightarrow 72}$ (hr.ng/mL)	364.8 379.9 (27.6)	350.9 364.6 (26.8)	104.0	96.9-111.5
$AUC_{0\rightarrow \infty}$ (hr.ng/mL)	453.4 473.4 (28.7)	449.4 463.3 (25.1)	100.9	95.5-106.6
C_{max} (ng/mL)	11.6 12.3 (32.7)	11.8 12.3 (27.5)	98.5	93.0-104.3
T_{max} § (h)	6.0 (2.0-12.0)	6.0 (3.0-12.0)		
$T_{½}$ $ (h)	29.4 (23.1)	29.5 (22.0)		

***** AURO-OLANZAPINE ODT (Olanzapine) Orally Disintegrating Tablets, 10 mg by Auro Pharma Inc.
† ZYPREXA® ZYDIS® (Olanzapine) Orally Disintegrating Tablets, 10 mg manufactured by Eli Lilly Canada Inc., Canada were purchased in Canada.
§ Expressed as median (range) only.
$ Expressed as arithmetic mean (CV%) only.

Schizophrenia and Related Disorders Trials

The efficacy of olanzapine in the reduction of and maintenance of the reduction of the manifestations of schizophrenia and related psychotic disorders was established in 3 well-controlled clinical trials of psychotic inpatients who, at entry met the DSM-III-R criteria for schizophrenia (most with a course at entry of 'chronic with acute exacerbation') and 1 well-controlled clinical trial of psychotic inpatients and outpatients who, at entry, met the DSM-III-R criteria for schizophrenia, schizophreniform disorder, or schizoaffective disorder.
The results of the trials follow:

(1) A 6-week, placebo-controlled trial (N = 335) compared 3 fixed dosage ranges of olanzapine (5 ± 2.5, 10 ± 2.5, and 15 ± 2.5 mg/day QD), 1 dosage range of haloperidol (15 ± 5 mg/day on a BID schedule), and placebo. The 2 higher dosage ranges of olanzapine were statistically significantly superior to placebo on the Brief Psychiatric Rating Scale (BPRS) total, the Clinical Global Impressions - Severity of Illness (CGI - S) scale, and the BPRS positive psychosis cluster. The highest dosage range of olanzapine was statistically significantly superior to placebo and to haloperidol on the Scale for the Assessment of Negative Symptoms (SANS). Efficacy of olanzapine generally increased with dose. The 5 ± 2.5 mg/day dosage range of olanzapine was numerically, but not statistically, significantly superior to placebo on BPRS total and other assessments of overall psychopathology.

(2) A 6-week, placebo-controlled trial (N = 152) compared 2 fixed doses of olanzapine (1 or 10 mg/day QD) and placebo. Olanzapine, 10 mg/day, was statistically significantly superior to placebo on the BPRS total, the BPRS positive psychosis cluster, the CGI-S scale, the Positive and Negative Syndrome Scale (PANSS) total, the PANSS positive subscale, and the PANSS negative subscale. Olanzapine, 1 mg/day, appeared to be a no-effect dose with no difference, clinically or statistically, from placebo on any assessment of psychopathology.

(3) A 6-week, dose comparison trial (N = 431) compared 3 fixed dosage ranges of olanzapine (5 ± 2.5, 10 ± 2.5 and 15 ± 2.5 mg/day QD), olanzapine (1 mg/day QD), and haloperidol (15 ± 5 mg/day on a BID schedule). There were no statistically significant differences between groups on efficacy measures except for the highest dosage range of olanzapine, which was statistically significantly superior to olanzapine, 1 mg, on the BPRS positive psychosis cluster, PANSS positive subscale, and the CGI-S scale.

(4) A 6-week comparator-controlled trial (N = 1996, 2:1 randomization, olanzapine: haloperidol) compared 1 dosage range of olanzapine (5 to 20 mg/day QD) and 1 dosage range of haloperidol (5 to 20 mg/day QD). The acute mean modal doses (for those patients with at least 3 weeks of treatment) were 13.2 mg/day for olanzapine and 11.8 mg/day for haloperidol. Olanzapine was statistically significantly superior to haloperidol on the BPRS total, the BPRS negative psychosis cluster, the PANSS negative subscale, and the CGI-S scale. Olanzapine was also statistically significantly superior to haloperidol on the Montgomery-Asberg Depression Rating Scale (MADRS). The validity of this scale in patients with schizophrenia, however, is not established.

(5) The effectiveness of olanzapine in long-term therapy, ie., > 6 weeks, was evaluated in 3 double-blind, controlled, extension maintenance trials (of acute trials 1, 3, and 4 above). Patients who showed adequate clinical improvement following double-blind acute therapy were allowed to continue on in a double-blind, long-term extension maintenance phase on their acute dosage regimen. Long-term maintenance of treatment response (as defined by continued reduction in signs and symptoms sufficient to not require hospitalization for psychosis) was compared over time (894 olanzapine-treated patients; median length of treatment was 237 days). The percentage of patients maintaining treatment response over one year was compared. Olanzapine was statistically significantly superior to placebo in the one placebo-controlled trial and was comparable or statistically significantly superior to the active comparator in 3 of 3 active comparator-controlled trials.

Summary of Schizophrenia and Related Disorders Trials

While the efficacy of olanzapine at a dose of 5 mg/day was not statistically superior to placebo (see (1 above)), some individual patients receiving this dose had a good acute response, and were well maintained during a 1-year extension phase.

The above trials (including open-label extension) and an additional trial in geriatric patients with primary degenerative dementia of the Alzheimer's type constitute the primary database (N = 2500 patients treated with olanzapine, corresponding to 1122.2 patient-years; N = 810 patients

treated with haloperidol, corresponding to 193.0 patient-years; N = 236 patients treated with placebo, corresponding to 27.1 patient-years).

Bipolar Disorder Trials

Bipolar Mania:
The efficacy of oral olanzapine in treating acute bipolar mania was demonstrated in 5 controlled studies, including 2 placebo-controlled studies, 2 active comparator studies and 1 cotherapy study. All patients enrolled in these studies had a diagnosis of bipolar I disorder and displayed an acute manic or mixed episode (with or without psychotic features) according to the DSM-IV criteria based on clinical assessment and confirmed by the structured clinical interview for the diagnostic and statistical manual, SCID-P.

- ***Placebo-Controlled Trials:*** The 2 placebo-controlled trials evaluated the efficacy of olanzapine versus placebo in treating bipolar manic or bipolar mixed episodes as measured by the Y-MRS total score LOCF mean change from baseline to endpoint over 3 weeks (n = 70 and n = 69, respectively) and 4 weeks (n = 60 and n = 55, respectively). These trials demonstrated superiority in the efficacy of olanzapine compared with placebo. The key findings were as follows:

    - Olanzapine, at a dose range of 5-20 mg/day, was statistically superior to placebo in improving manic symptoms in each study ($p = 0.019$ and $p < 0.001$, respectively).
    - In each study, a statistically significantly greater percentage of olanzapine-treated patients (48.6% and 64.8%, respectively) compared with placebo-treated patients (24.2% and 42.9%, respectively) responded to treatment (≥ 50% reduction in Y-MRS total score) ($p = 0.004$ and $p = 0.023$, respectively).
    - In each study, the percentage of patients that were in clinical remission (endpoint Y-MRS total score ≤ 12) were significantly greater among olanzapine patients (45.7% and 61.1%, respectively) compared with placebo patients (25.8% and 35.7%, respectively) ($p = 0.020$ and $p = 0.013$, respectively).
    - Olanzapine efficacy did not differ significantly among the main subtypes of bipolar mania, for example patients with a history of rapid cycling, with or without psychotic features, and bipolar mixed or bipolar manic.

(1) ***Active Comparator Trials:*** Two active comparator trials were conducted.

(a) The first active comparator study evaluated the efficacy of olanzapine versus divalproex in treating bipolar manic and bipolar mixed episodes by using the Y-MRS total score LOCF mean change from baseline to endpoint. This study was a 3-week, double-blind study with a double-blind continuation phase of 11 months. The primary objective of the study was to demonstrate non-inferiority in the efficacy of olanzapine compared with divalproex at 3 weeks. Patients were randomized to either olanzapine (5-20 mg/day, n = 125) or divalproex (500-2500 mg/day, n = 126). The key findings were as follows:

- Olanzapine was statistically superior to divalproex in improving manic symptoms as measured by Y-MRS change score at 3 weeks (mean improvements of 13.4 and 10.4 points, respectively, $p = 0.028$).

- The proportion of patients meeting the criteria for response was not statistically significantly different between olanzapine and divalproex groups (54.4% and 42.3%, respectively) (p = 0.059).
- The proportion of patients that was in clinical remission was significantly greater among olanzapine patients (47.2%) compared with divalproex patients (34.1%) (p = 0.039).

(b) The second active comparator study evaluated the efficacy of olanzapine versus haloperidol in treating bipolar manic or mixed episodes by assessing the proportion of patients in protocol-defined remission from manic and depressive symptoms at 6 weeks. Remission was defined as: 1) achieving improvement in clinical symptomatology in manic and depressive symptoms; 2) having achieved specific reductions in Y-MRS and HAMD-21 total scores; and 3) continuing to take study medication at Week 6. This trial consisted of a 6-week double-blind phase followed by a 6-week double-blind maintenance of response phase in the absence of a placebo arm. Patients were randomly assigned to treatment with olanzapine 5-20 mg/day (n = 234) or haloperidol 3-15 mg/day (n = 219). The key findings were as follows:

- Olanzapine and haloperidol were similarly effective in improving manic symptoms.
- A clinical response to treatment was defined as a ≥ 50% improvement in Y-MRS total score from baseline to endpoint. In both treatment groups, a large proportion of patients responded to treatment. At the end of the acute phase 72.3% and 74.2% of olanzapine and haloperidol patients, respectively met the response criteria, and at the end of the continuation phase almost all patients were classified as responders (96.3% of 160 olanzapine patients and 94.1% of 136 haloperidol patients).
- The proportion of patients in symptomatic remission at the end of the acute phase (6 weeks) was similar for olanzapine and haloperidol patients (52.1% versus 46.1%, respectively (p = 0.152)). Among patients who entered the continuation phase and were not in symptomatic remission at 6-weeks, significantly more olanzapine patients (68.3%) than haloperidol patients (41.0%) were in remission by the end of the continuation period (p = 0.014).
- Manic symptoms continued to improve among olanzapine patients to a statistically significant extent.
- Olanzapine was statistically significantly more efficacious than haloperidol in patients without psychotic features (acute phase remission rates were 56.7% in 104 olanzapine patients and 41.6% in 89 haloperidol patients, respectively) (p = 0.043).

(2) ***Cotherapy Trial:*** This trial evaluated the efficacy of olanzapine plus either valproate or lithium (cotherapy, n = 229) versus valproate or lithium alone (monotherapy, n = 115) in treating bipolar manic or mixed episodes as measured by the Y-MRS total score LOCF mean change from baseline to endpoint. This study was a 6-week, double-blind study with a re-randomized double-blind phase of 18 months. The key findings were as follows:

- Olanzapine in combination with either valproate or lithium was significantly more efficacious than monotherapy (valproate or lithium) in improving manic symptoms (mean improvements of 13.1 and 9.1 points, respectively) ($p = 0.003$).
- The proportion of patients that clinically responded to treatment was statistically significantly greater among patients receiving olanzapine cotherapy (67.7%) than lithium or valproate monotherapy (44.7%, $p < 0.001$).
- The percentage of patients that were in clinical remission was significantly greater in the olanzapine cotherapy group (78.6%) compared with the lithium or valproate monotherapy group (65.8%, $p = 0.012$).
- The difference in time to remission was also statistically significantly different ($p = 0.002$). The median estimated remission time was 14 days for olanzapine cotherapy-treated patients and 22 days for monotherapy-treated patients.

Bipolar Maintenance:
The efficacy of oral olanzapine as monotherapy for maintenance treatment of bipolar disorder in patients who responded to acute treatment with olanzapine for a manic or mixed episode was demonstrated in two 1-year 'time to event' controlled trials: one placebo-controlled and one active comparator trial against lithium monotherapy.

All patients enrolled in these studies had a diagnosis of bipolar I disorder and displayed an acute manic or mixed episode (with or without psychotic features) according to the DSM-IV criteria.

For both studies: Patients had to meet study-defined response criteria (YMRS total score of ≤ 12 and a HAMD-21 total score ≤ 8) during open-label treatment with olanzapine (or olanzapine plus lithium in the active comparator study) in order to be randomized into the double-blind maintenance period for observation of study-defined relapse. Dosing was flexible (5 - 20 mg/day for olanzapine; serum levels 0.6 - 1.2 mEq/L for lithium).

The exit criteria was symptomatic relapse of bipolar disorder, either mania or depression. Symptomatic relapse of mania was defined as reaching a YMRS total score ≥ 15, and symptomatic relapse of depression as reaching a HAMD-21 total score ≥ 15; for the placebo-controlled study only, the definitions also included being hospitalized for mania or depression. Thus, the primary efficacy variable was time to, and incidence of, the exit symptomatic relapse of bipolar disorder, based on analysis of Kaplan-Meier time-to-relapse curves.

1) Placebo-controlled trial:
The study evaluated the efficacy of olanzapine vs placebo in maintenance treatment of manic or mixed bipolar episodes by using survival curve analysis to assess the time to, and incidence of, relapse of bipolar disorder. In this trial, 361 patients who had demonstrated response criteria for an average of 16 days were randomized to either continuation of olanzapine at their same dose (n = 225) or to placebo (n = 136), for observation of relapse for up to one year.

The key findings were as follows:
- Figure 1 shows the 1-year time-to-relapse curves for total discontinuations from the study in each arm over time, whether exiting due to relapse, or withdrawal due to adverse events or

other reasons. The percentage of patients remaining in the study (i.e., relapse-free, and have not withdrawn for any reason) can be seen at each of the 3, 6, 9 and 12 month points; at study-end, this is 24% (n = 53) for olanzapine and 10% (n = 13) for placebo. The time-point at which 50% of the patients in a specific arm had withdrawn for any reason was Day 59 for the olanzapine group compared to Day 23 for the placebo group.

- Figure 2 shows the 1-year time-to-relapse curves for specifically the exit criterion of bipolar relapse (i.e., patients who withdrew for other reasons were censored and excluded from the calculated numerators and denominators). Olanzapine was superior to placebo, for both incidence of bipolar relapse (46.7% vs 80.1%, respectively), and median time to relapse (174 days vs 22 days, respectively). Note that a high relapse incidence for the placebo arm is not unexpected given the limited time that patients had been demonstrating response criteria prior to randomization.

- Figures 2a and 2b show the efficacy time-to-relapse curves for each of manic and depressive relapse, respectively. Olanzapine showed a statistically significant advantage over placebo in terms of each of mania and depression, although a greater advantage was seen in mania.

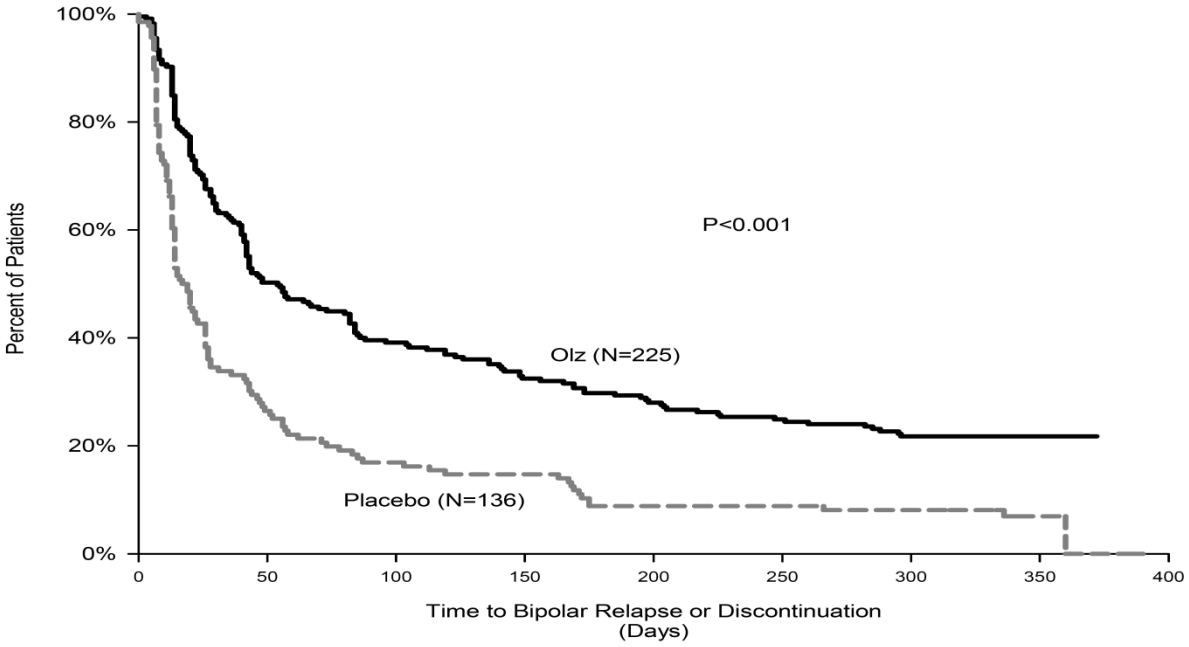


Figure 1: Time to Event (Relapse or Discontinuation)
Study HGHL; Double-Blind Treatment Phase

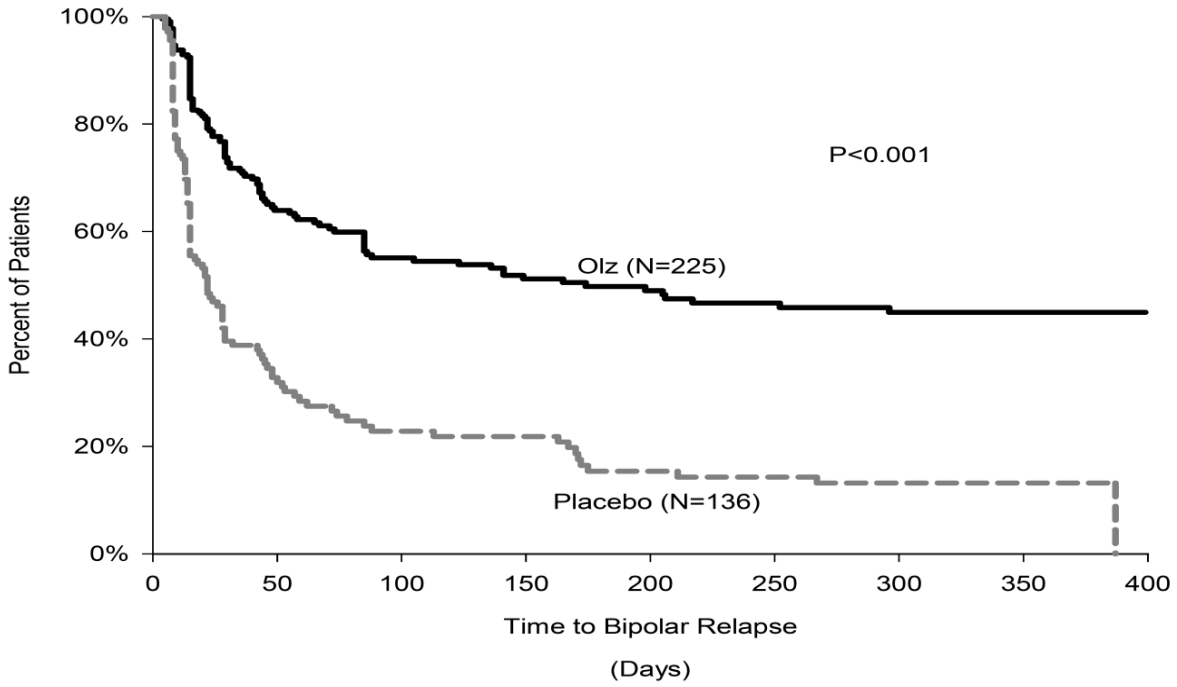


Figure 2: Time to Symptomatic Relapse of Bipolar Disorder, Including Hospitalization Study HGHL; Double-Blind Treatment Phase

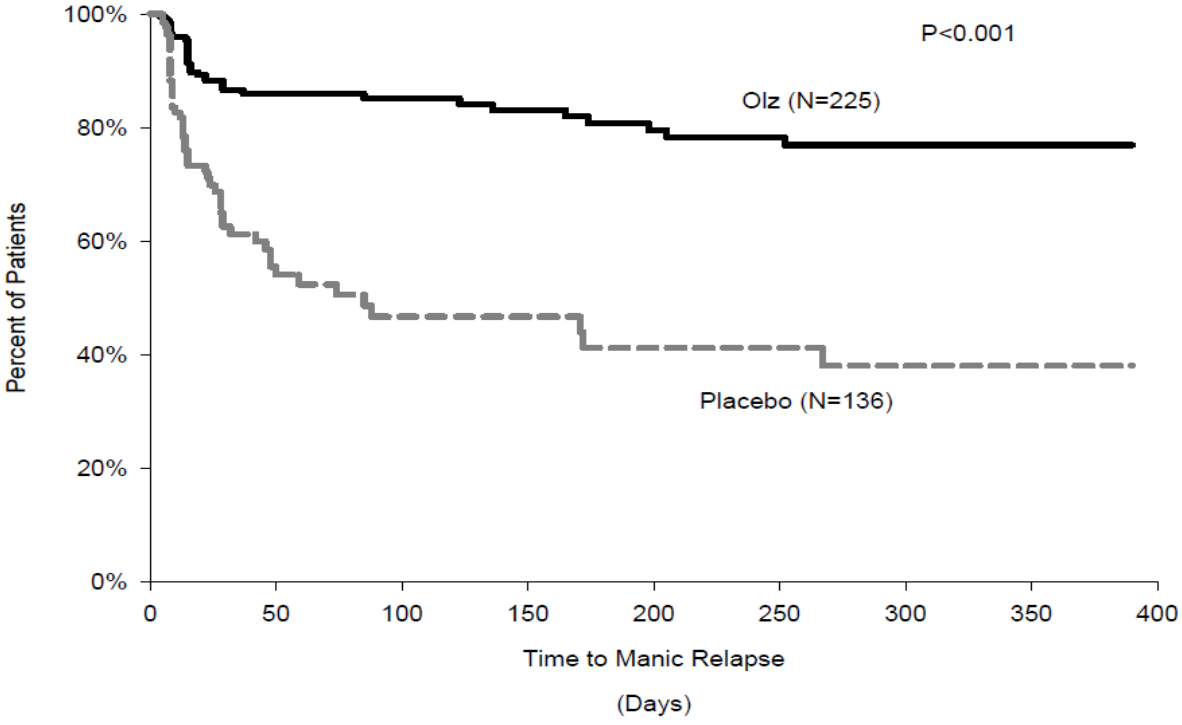


Figure 2a: Time to Symptomatic Relapse of Mania, Including Hospitalization Study HGHL; Double-Blind Treatment Phase

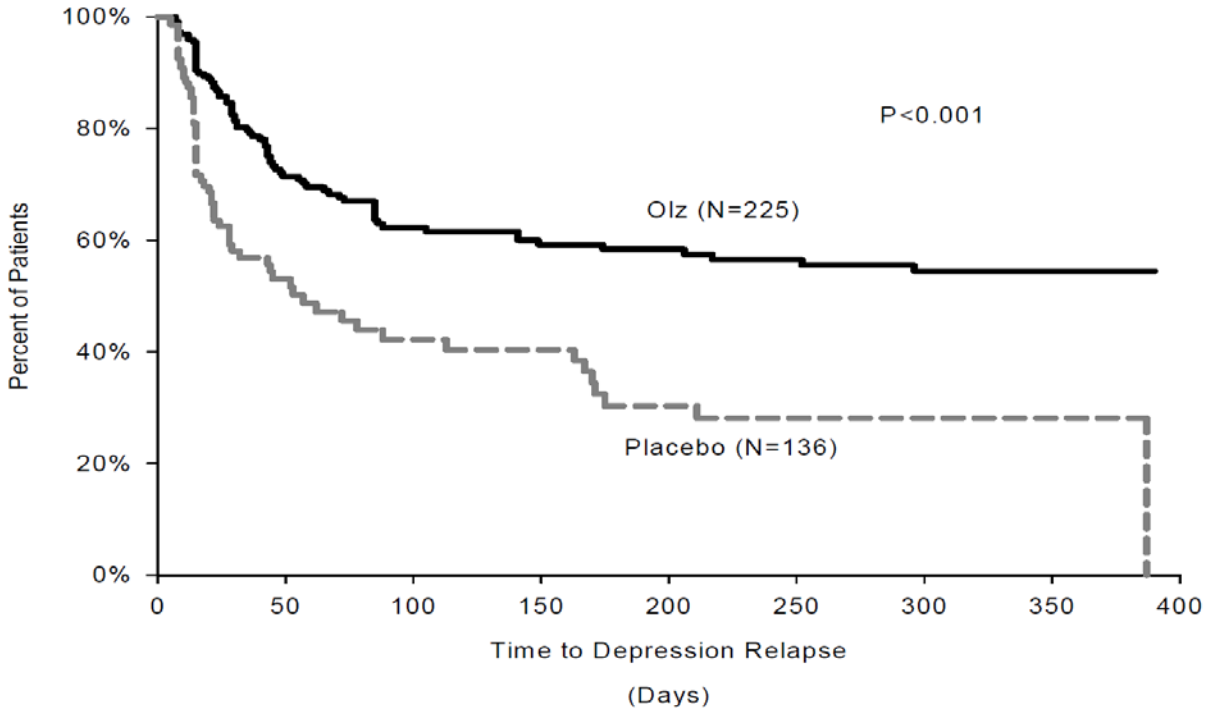


Figure 2b: Time to Symptomatic Relapse of Depression, Including Hospitalization Study HGHL; Double-Blind Treatment Phase

2) Active comparator trial:
The study evaluated the efficacy of olanzapine vs lithium in maintenance treatment of manic or mixed bipolar episodes in a non-inferiority design to assess the incidence of relapse of bipolar disorder and further, by using survival curve analysis to assess time to relapse. In this trial, 543 patients who had demonstrated response criteria for an average of 20 days were randomized to either olanzapine plus placebo (n = 217) or lithium plus placebo (n = 214) for observation of relapse for up to one year. The first month of the double blind period was a taper period to allow for non-abrupt lithium discontinuation. The non-inferiority margin used in this study was: ± 20% of the efficacy seen for the reference population.

The key findings were as follows:

- Figure 3 shows the 1-year time-to-relapse curves for total discontinuations from the study in each arm over time, whether exiting due to relapse, or withdrawal due to adverse events or other reasons. The percentage of patients remaining in the study (i.e., relapse-free, and have not withdrawn for any reason) can be seen at each of the 3, 6, 9 and 12 month points; at study-end, this is 42% (n = 94) for olanzapine and 28% (n = 61) for lithium. The time-point at which 50% of the patients in a specific arm had withdrawn for any reason was Day 255 for the olanzapine group compared to Day 192 for the lithium group.

- Figure 4 shows the 1-year time-to-relapse curves for specifically the exit criterion of bipolar relapse (i.e., patients who withdrew for other reasons were censored and excluded from the calculations of numerators and denominators). Olanzapine was non-inferior to lithium for both incidence of bipolar relapse (30.0% vs 38.8%, respectively), and time to 25% of patients experiencing relapse (122 days vs 143 days, respectively).

- It can be seen from Figure 4 that for approximately the first five months of the 1-year trial, relapse rate was higher in olanzapine-treated patients; thereafter, the rate of relapse for lithium increases, while that for olanzapine flattens out.

- Figures 4a and 4b are the 1-year time-to-relapse curves for each of the exit criterion of manic and depressive relapse respectively. Olanzapine showed a statistically significant advantage over lithium in rate of mania relapse, and was non-inferior for depressive relapse.

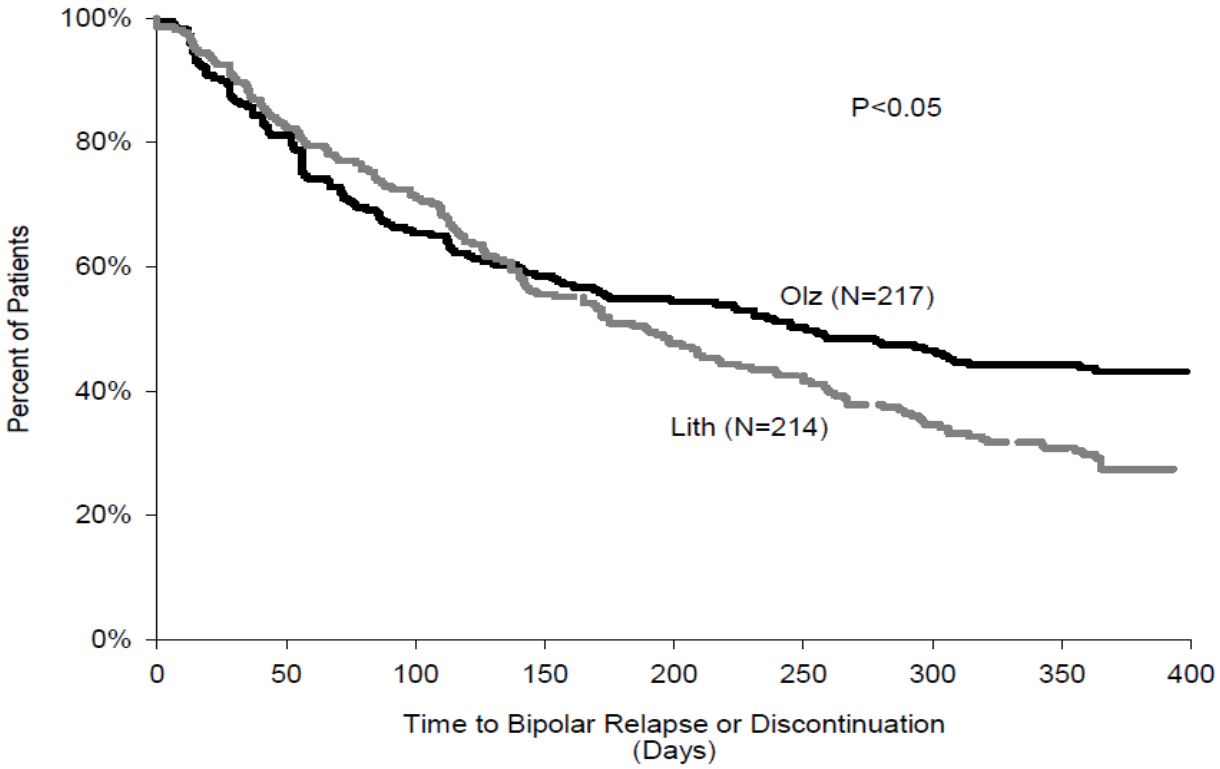


Figure 3: Time to Event (Relapse or Discontinuation)
Study HGHT; Double-Blind Treatment Phase

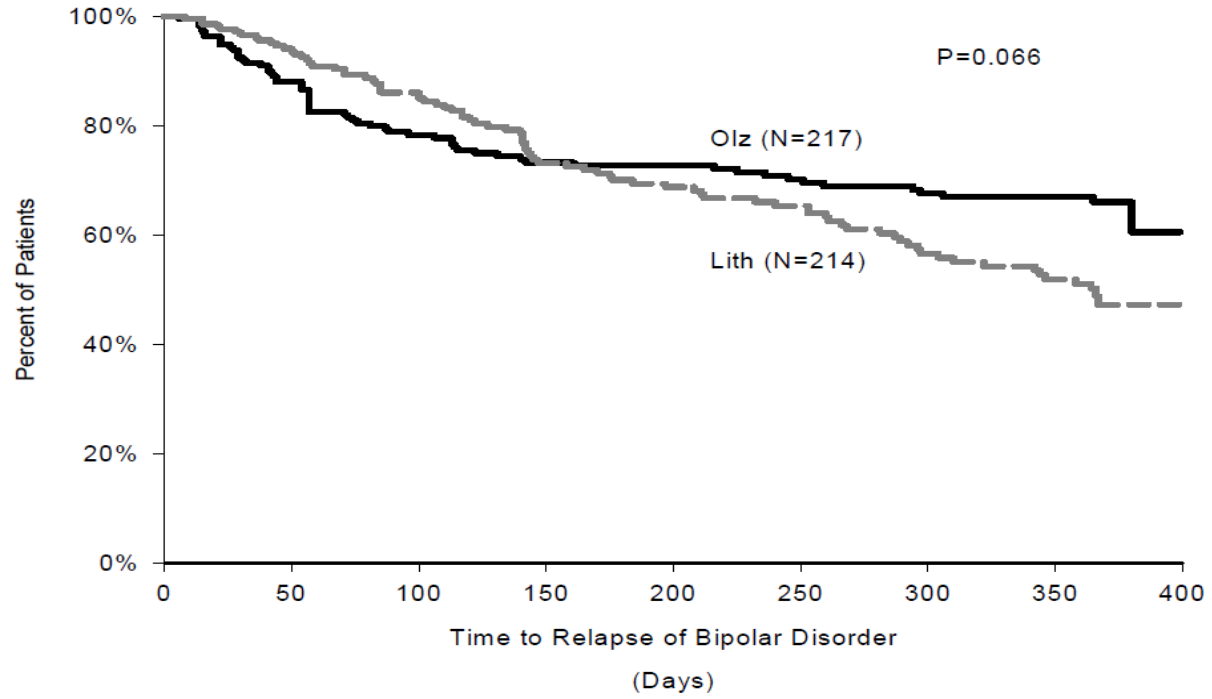


Figure 4: Time to Symptomatic Relapse of Bipolar Disorder
Study HGHT; Double-Blind Treatment Phase

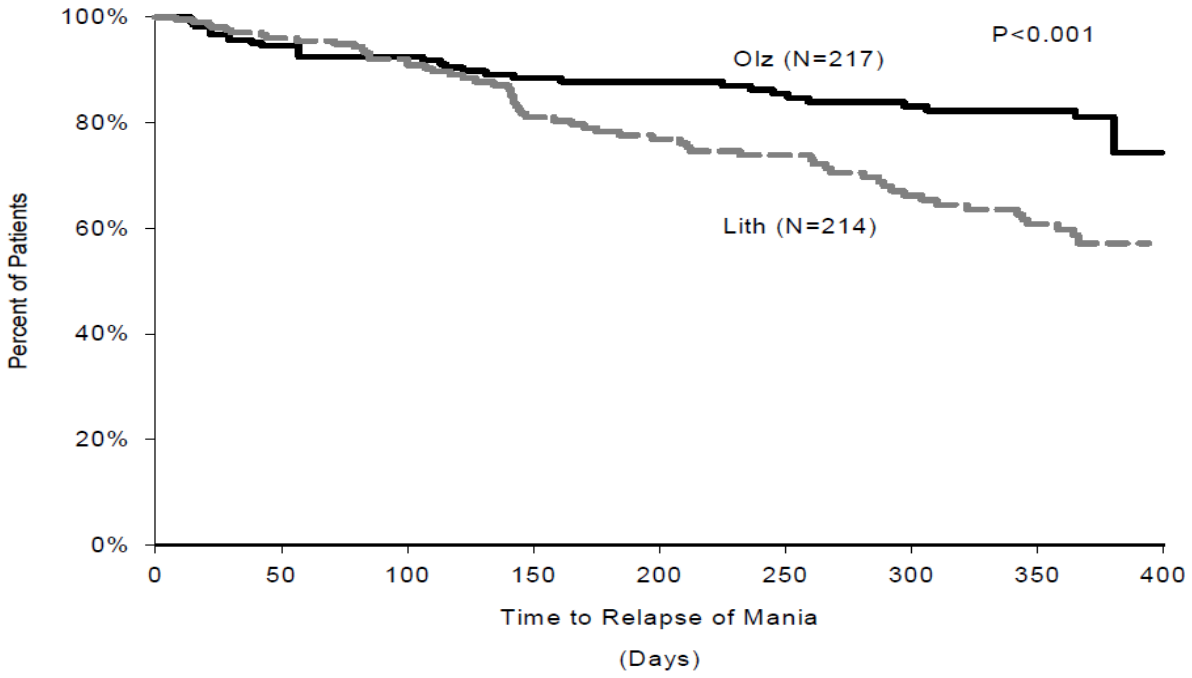


Figure 4a: Time to Symptomatic Relapse of Mania
Study HGHT; Double-Blind Treatment Phase

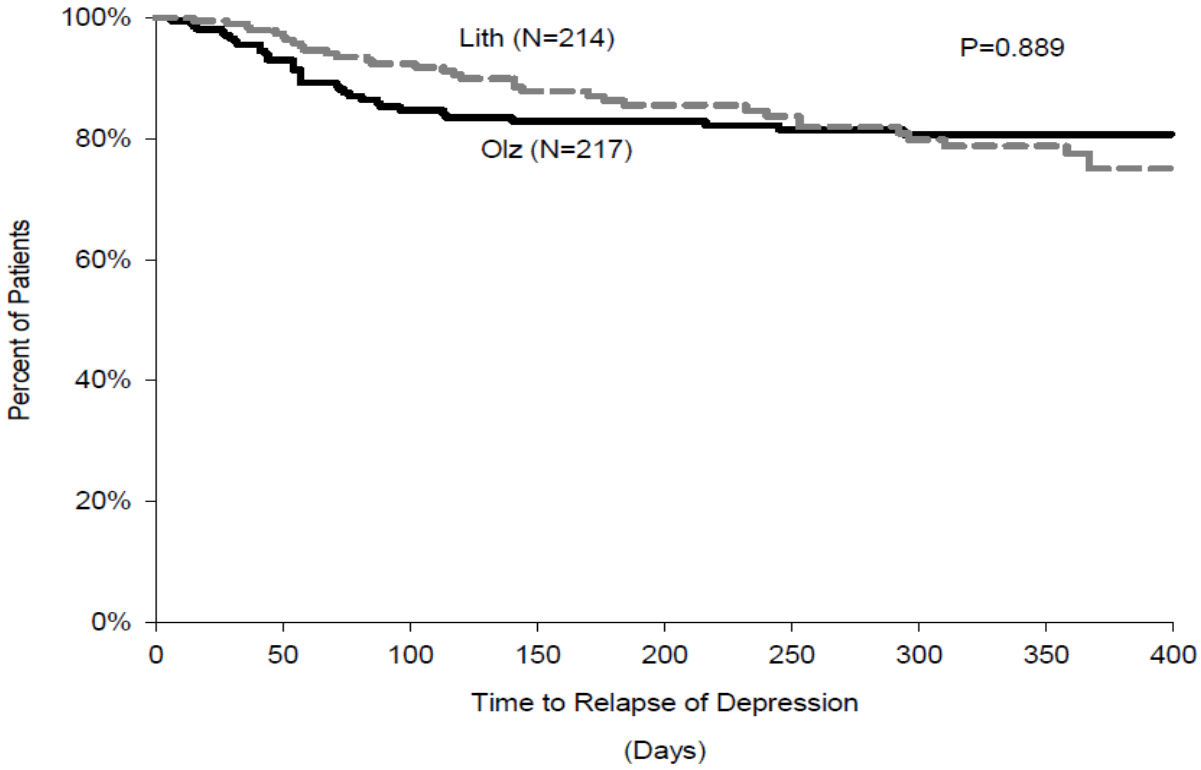


Figure 4b: Time to Symptomatic Relapse of Depression
Study HGHT; Double-Blind Treatment Phase

Summary of Bipolar Disorder Trials
Bipolar Mania: Olanzapine was more efficacious than placebo and divalproex and as effective as haloperidol in improving overall manic symptomatology in patients with acute bipolar I disorder, manic or mixed episode, with or without psychotic symptoms, and with or without a history of rapid cycling. Olanzapine is associated with a faster onset of action (based on median time to remission estimated with Kaplan-Meier analysis) compared to divalproex and similar to that of haloperidol. The addition of olanzapine also improved patients not responding to lithium or valproate. Olanzapine was not associated with inducing or worsening symptoms of depression.

Bipolar Maintenance: Two 1-year controlled studies support the use of olanzapine monotherapy in maintenance treatment of bipolar patients who responded to acute olanzapine treatment for a manic or mixed episode. Based on analysis of one-year Kaplan-Meier survival curves, olanzapine was superior to placebo, and non-inferior to lithium, in both time to, and incidence of, bipolar relapse over one year.

DETAILED PHARMACOLOGY

Pharmacodynamics:
In Vitro Receptor Binding Affinities:
The binding affinities of olanzapine versus clozapine and haloperidol are summarized in Table 11. The binding profile of olanzapine has similarities to that produced by clozapine, although the

affinity of olanzapine is somewhat greater for dopamine D_1 and D_2 receptors and lower at 2 receptors. With respect to 5-HT receptor subtypes, both agents show greatest affinity for 5-HT_{2A} and 5-HT_{2C} receptors. The ratio of activity between 5-HT_{2A} and D_2 receptors is slightly less for olanzapine than for clozapine, although olanzapine is still about twice as active at 5-HT_{2A} receptors compared with D_2 receptors. Both compounds also have a high affinity for muscarinic receptor subtypes, particularly the m1 site. The affinity constants (Ki, nM) for olanzapine, clozapine, and haloperidol are shown below:

Table 11: Affinity constants for olanzapine, clozapine, and haloperidol

Compound	**Dopamine D_1**	**Dopamine D_2**	**α_1**	**α_2**	**Histamine H_1**
Olanzapine	31 ± 0.7	11 ± 2	19 ± 1	230 ± 40	7 ± 0.3
Clozapine	85 ± 0.7	125 ± 20	7 ± 4	8 ± 3	6 ± 2
Haloperidol	25 ± 7	1 ± 0.04	46 ± 6	360 ± 100	3630 ± 85

Compound	**5-HT_{1A}**	**5-HT_{1B}**	**5-HT_{1D}**	**5-HT_{2A}**	**5-HT_{2C}**	**5-HT_3**
Olanzapine	>10,000	1355 ± 380	800 ± 190	4 ± 0.4	11 ± 1	57
Clozapine	770 ± 220	1200 ± 170	980 ± 115	12 ± 3	8 ± 0.8	69
Haloperidol	7930 ± 500	>10,000	6950 ± 950	78 ± 22	3085	>1000

Compound	**m_1**	**m_2**	**m_3**	**m_4**	**m_5**
Olanzapine	1.9 ± 0.1	18 ± 5	25 ± 2	13 ± 2	6 ± 0.8
Clozapine	1.9 ± 0.4	10 ± 1	14 ± 1	18 ± 5	5 ± 1.2
Haloperidol	1475 ± 300	1200 ± 180	1600 ± 305	>10,000	Not Tested

Olanzapine has no significant activity at $GABA_A$, benzodiazepine, or β receptors. Olanzapine also interacts with dopamine D_4 receptors (Ki 27 nM).

In vivo biochemical studies were conducted to confirm the binding data and investigate the functional consequences of interacting with these neurotransmitter receptor sites.

In Vivo Neuroendocrine Studies:
It has been shown that corticosterone concentrations in rats can be elevated by 5-HT-mediated or dopamine-mediated mechanisms. Olanzapine antagonizes the 5-HT (quipazine-induced) (ED_{50} 0.57 mg/kg) and D_2 dopamine receptor-mediated (pergolide-induced) (ED_{50} 3 mg/kg) increases in corticosterone. These results show that olanzapine has greater activity at 5-HT compared with D_2 dopamine receptors *in vivo*. These results complement the behavioural studies showing that olanzapine preferentially antagonizes a 5-HT-induced response.

In Vivo Behavioural Pharmacology:
In behavioural studies, olanzapine exhibits a broad pharmacologic profile, as predicted from the biochemical data.

Olanzapine blocks apomorphine-induced climbing behaviour with an ED_{50} of approximately 5 mg/kg. The climbing response has previously been shown to require both D_1 and D_2 receptor activation. These results therefore indicate that olanzapine possesses dopamine antagonist activity *in vivo*.

A second study in mice looked at the ability of olanzapine to block 5-hydroxytryptophan (5-HTP)-induced head twitches, a test probably mediated by $5\text{-}HT_2$ receptors. Olanzapine produced dose-related reductions in the head-twitch response with approximate ED_{50}s of 2 mg/kg. Olanzapine preferentially blocks the head twitch, compared with the climbing response, demonstrating that this agent exhibits greater activity at the 5-HT receptor compared with dopamine receptors *in vivo*. These results agree with those reported in rats, showing that olanzapine preferentially antagonizes 5-HT-mediated rather than dopamine-mediated elevations in corticosterone (Moore et al. 1993).

Olanzapine doses of 2.5 to 10 mg/kg produced a significant reduction in oxotremorine-induced tremor in mice, with an ED_{50} of 3 mg/kg. These results demonstrate that olanzapine possesses anticholinergic activity *in vivo* at doses which also antagonize dopamine-mediated effects.

Inhibition of a conditioned avoidance response has been widely used as a test to predict the antipsychotic potential of a compound, while the induction of catalepsy in rats is associated with the occurrence of extrapyramidal symptoms in the clinic. ED_{50}s for the various compounds in blocking a conditioned avoidance response or inducing catalepsy in rats are given in Table 12.

Table 12: Effect of Olanzapine and Haloperidol on Conditioned Avoidance Responding (CAR) and the Induction of Catalepsy (CAT) in Lister Hooded Rats

Compound	CAR	CAT	Ratio
Olanzapine	5.6 (4.6-6.8)	23 (18.7-29)	4.1
Haloperidol	0.28 (0.24-0.33)	0.74 (0.6-0.9)	2.6

Note: The results are expressed as ED_{50} values (mg/kg p.o.) with 95% confidence intervals stated in parentheses. The ratio is the ED_{50} CAT / ED_{50} CAR.

Although olanzapine induces catalepsy, this only occurs at doses higher than those required to block the conditioned avoidance response.

A number of reports have shown that the "atypical" agent, clozapine, differs from "typical" antipsychotics in its effects on schedule-controlled behaviour. In a rat or pigeon conflict test, olanzapine, clozapine, and chlordiazepoxide produced the characteristic changes in rates of responding associated with anxiolytics, although the effect of olanzapine and clozapine was smaller than that seen with chlordiazepoxide. All three compounds decreased or had no effect on

the high rates of responding produced in the reward component, whereas the rates in time-out and particularly the conflict period were increased. This type of profile was not seen with the "typical" antipsychotic, haloperidol, which only decreased the rates in all the components. These data further emphasize the "atypical" profile of olanzapine.

In Vivo Electrophysiology:
“Typical” antipsychotic agents, such as haloperidol, reduce the spontaneous firing of both A9 and A10 dopaminergic neurons in the CNS following chronic dosing. The A9 (nigrostriatal system) is thought to mediate extrapyramidal motor disturbances, while the A10 (mesolimbic system) has been associated with the antipsychotic activity of compounds. Olanzapine (10 and 20 mg/kg subcutaneously for 21 days) produced a significant reduction in the firing of A10 dopaminergic cells. The number of spontaneously active A9 neurons either remained constant or was increased. These results are very similar to those reported previously for clozapine and further emphasize the “atypical” pharmacologic profile of olanzapine.

Human Versus Animal Metabolism:
In animal species (mice, rats, and dogs) used for toxicologic evaluation, olanzapine was metabolized through aromatic hydroxylation (forming phenolic metabolites and/or their glucuronide conjugates), allylic (alkyl) oxidation, N-dealkylation, and N-oxidation reactions.

Although similarities in the metabolic fate of olanzapine in animals (mice, rats, and dogs) and humans include the 2-alkyl hydroxylation, N-dealkylation, and N-oxidation pathways, two significant differences can be noted. First, direct glucuronidation, producing mainly 10-N-glucuronide and to a lesser extent 4’-N-glucuronide, was a significant metabolic pathway in humans. These N-glucuronides were absent in animal species except for a trace amount of 10-N-glucuronide in dog urine. Second, metabolites resulting from aromatic oxidation were not found in any human biological fluids. The monkey also did not appear to form 10-N-glucuronide, but was similar to humans in apparently not forming metabolites resulting from the oxidative attack of the benzene ring of olanzapine.

TOXICOLOGY

An extensive series of acute, subchronic, chronic, reproduction, and genetic toxicity as well as oncogenicity studies have been conducted to support clinical trials with olanzapine. In most of these studies, olanzapine was given by the oral route to rodents, rabbits, and monkeys in an aqueous suspension with 5% to 10% acacia and to dogs as neat material in capsules.

The predominant effects in laboratory animals given olanzapine were CNS depression and anticholinergic effects related to the pharmacology of the drug. Tolerance to the CNS depression developed in repeated-dose studies. Depressed body weight gain was a consistent finding in mice given 30 mg/kg/day and in rats given 4 mg/kg/day. Effects on hematology parameters were found in each species studied in repeated-dose studies. Rats given 16 mg/kg/day had decreased lymphocyte and neutrophil counts and atrophy of bone marrow consistent with the marked reduction in body weight gain. Mice given 3 mg/kg/day developed leukopenia, due primarily to lymphocytopenia, but also associated with neutropenia. Lymphoid necrosis of thymus and spleen

was seen in mice given ≥ 10 mg/kg/day. Instances of reversible neutropenia, with or without thrombocytopenia, or anemia developed in a low number of individual dogs treated with 8 or 10 mg/kg/day. Bone marrow from some dogs with olanzapine-induced neutropenia responded to olanzapine with lower than expected numbers of maturing granulocytic cells; however, progenitor and proliferating cells were present in adequate numbers. No olanzapine-related hematologic effects were seen in dogs receiving olanzapine at either 2 or 5 mg/kg/day.

Effects observed in rats consistent with increased plasma concentrations of prolactin in rats included decreased weights of ovaries and uterus. Histopathologic tissue alterations in mammary gland morphology and vaginal epithelium and increased prominence of ovarian follicles were also consistent with elevated prolactin concentrations. Prolactin-induced histopathologic tissue alterations found in rats regressed after treatment cessation. No unexpected toxicologically important findings unrelated to pharmacologic activity were found in the 1-year studies in rats given 4 mg/kg/day or in dogs given 5 mg/kg/day.

In a rat oncogenicity study, the only neoplasm with increased incidence related to treatment was malignant mammary gland tumours in females of the 4- and 8-mg/kg/day groups (initial dose levels were increased from 2.5 and 4 mg/kg/day, respectively, on Day 211). The overall incidence of mammary gland tumours was not increased. The shift in expression of mammary gland tumours was not unexpected and was consistent with effects due to elevated prolactin concentrations in rodents. Also consistent with increased prolactin concentrations was an increased total incidence of mammary gland tumours in female mice given 10 or 20 mg/kg/day (the high dose was decreased from 30 mg/kg/day due to excess mortality).

Olanzapine had no mutagenic or teratogenic effects. Mating performance was affected in male rats given 5 mg/kg/day, but the effect was quickly reversed when treatment stopped. Estrous cycles were affected and reproduction parameters were influenced in rats given the higher doses of ≥ 1 mg/kg/day. No adverse effects were observed on numbers of corpora lutea, implantations, fetal viability, or fetal weight, and there were no effects on litter size or on the survival, growth, or development of the offspring from parents given up to 5 mg/kg/day. Transient modest decreases in activity levels of the progeny from females given 0.25 mg/kg/day and skeletal changes indicative of growth retardation in fetuses from females given 5 mg/kg/day were observed. Although the reproductive process in female rats from mating through fertilization was not adversely affected by treatment, this evidence does not exclude a possible interference with maintenance of pregnancy at high doses of olanzapine.

The findings of toxicology studies support the safety of olanzapine for oral use in humans as an antipsychotic agent.

Acute Toxicity Studies:
The acute toxicity of olanzapine was studied in mice, rats, dogs, and monkeys. The estimated median lethal dose for each species is shown below in Table 13:

T able 13: Acute Toxicity Summary

		Estimated Median Lethal Dose (mg/kg/day)	
Species	**Route**	**Males**	**Females**
Mouse	Oral	211	208
Rat	Oral	174	177
Dog	Oral	Both sexes > 100 mg/kg	
Monkey	Nasogastric	Both sexes > 100 mg/kg	
Rat	Intraperitoneal	112	107

Signs of toxicity in rodents included hypoactivity, lethargy, leg weakness, coma, tremors, clonic convulsions, salivation, poor grooming, and depressed body weight gain.

The potential for irritation of an aqueous intramuscular formulation of olanzapine was tested in one *in vitro* and two *in vivo* (dog and rabbit) studies. The intent of these studies was to characterize the effects at the site of injection. Overall, these tests indicated that formulations of olanzapine, at 1.7 to 8.4 mg/mL in a tartaric acid/lactose vehicle, have the potential to cause slight irritation of skeletal muscle. While the *in vitro* model suggested a potential for moderate irritation at the higher concentrations tested, the *in vivo* models indicated either very little or slight potential for irritation.

Subchronic/Chronic/Carcinogenicity and Related Toxicity Studies:

Subchronic Toxicity Studies:

Subchronic administration studies of up to 3 months in duration have been conducted by the oral route in mice, rats, and dogs.

Chronic Toxicity Studies:

Chronic administration studies of up to 1 year were conducted by the oral route in rats and dogs.

Carcinogenicity Studies:

The oncogenic potential of olanzapine was evaluated in studies in rats and mice. Carcinogenicity studies were conducted in CD-1 mice and Fischer 344 rats. Olanzapine was administered orally to mice at doses of 3, 10, or 20 mg/kg for 19 months (males) or 21 months (females) in an initial study, and in a subsequent study at doses of 0.5, 2, or 8 mg/kg for 21 months (males and females). Rats received oral doses of 0.25, 1, 2.5, or 4 mg/kg (males) or 0.25, 1, 2.5, 4, or 8 mg/kg (females) for 24 months. These doses are equivalent to 2 to 70 times the maximum daily human dose (mouse studies) or 0.9 to 28 times the maximum daily human dose (rats). A maximum tolerated dose was achieved in both mouse and rat studies. Increased mortality was seen in mice at doses of 10 and 20 mg/kg and decreases in circulating lymphocytes and neutrophils were seen at doses 0.5 mg/kg. In female mice treated with olanzapine, the incidence of mammary tumours was increased at doses ≥ 2 mg/kg. Female rats treated with 4 or 8 mg/kg had an increase in malignant mammary tumours, but the overall incidence of mammary gland neoplasia was unchanged. Antipsychotic drugs, including olanzapine, have been shown to chronically elevate prolactin concentrations in rodents. An increase in mammary neoplasms has been found in rodents after chronic administration of other antipsychotic drugs and is considered to be prolactin mediated. The role of prolactin in human breast cancer has not been defined

conclusively, and there are presently no epidemiologic data indicating increased risk for breast cancer for humans using antipsychotic drugs.

Reproduction Studies:
Fertility studies in male and female rats and teratology studies in rats and rabbits have been conducted by the oral route. Mating performance was affected by administration of olanzapine due to sedation in male rats given doses greater than 18 times the maximum daily human dose, but the effect was quickly reversed when treatment stopped. Estrous cycles were affected and reproduction parameters were influenced in rats given doses greater than 4 times the maximum daily human dose. No adverse effects were observed on numbers of corpora lutea, implantations, fetal viability, or fetal weight, and there were no effects on litter size or on the survival, growth, or development of the offspring from parents given up to 18 times the maximum daily human dose. Although the reproductive process in female rats from mating through fertilization was not adversely affected by treatment, this evidence does not exclude a possible interference with maintenance of pregnancy at high doses of olanzapine. Reproduction studies, performed in rats and rabbits at doses of olanzapine 3.5 and 7 times the maximum daily human dose (20 mg), respectively, have revealed no evidence of harm to the fetus. Maternal toxicity, developmental toxicity (indicated by fetal growth retardation and slightly delayed ossification at birth), and increased numbers of nonviable offspring occurred at higher doses (in rats at 14 and 63 times the maximum daily human dose and in rabbits at 28 and 105 times the maximum daily human dose). However, fetal malformations were not increased. Transient decreases in offspring activity have occurred at all doses; however, there were no effects on body weight, growth, mating, fertility, or live births in second-generation animals. Placental transfer of olanzapine occurs in rat fetuses. Olanzapine was also detected in the milk of rats at concentrations up to three-fold higher than those in the plasma.

Mutagenicity Studies:
Olanzapine was not mutagenic or clastogenic in a full range of standard tests which included bacterial mutation tests and *in vitro* and *in vivo* mammalian tests. Appropriate positive controls were used in each test to verify the sensitivity of the test systems.

Hematologic Indices:
In animal studies with olanzapine, the principal hematologic findings were reversible peripheral cytopenias in individual dogs given high doses of olanzapine (24 to 30 times the maximum daily human dose), dose-related decreases in lymphocytes and neutrophils in mice, and lymphopenia secondary to compromised nutritional status in rats. A few dogs treated with 24 to 30 times the maximum daily human dose developed reversible neutropenia or reversible hemolytic anemia between 1 and 10 months of treatment. Effects on hematology parameters in each species involved circulating blood cells, and no evidence of bone marrow cytotoxicity was found in any of the species examined.

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PART III: CONSUMER INFORMATION

[Pr]AURO-OLANZAPINE ODT
olanzapine orally disintegrating tablets
5 mg, 10 mg, 15 mg and 20 mg
House Standard

This leaflet is part III of a three-part "Product Monograph" published when AURO-OLANZAPINE ODT was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about AURO-OLANZAPINE ODT. Contact your doctor or pharmacist if you have any questions about the drug.

Keep this information with your medicine in case you need to read it again.

ABOUT THIS MEDICATION

The name of your medicine is AURO-OLANZAPINE ODT and your doctor has prescribed it to help relieve the symptoms that are bothering you. AURO-OLANZAPINE ODT can help to control your symptoms and reduce the risk of relapse. Although AURO-OLANZAPINE ODT cannot cure your symptoms, it can help you keep them under control as you continue your treatment.

What the medication is used for:
AURO-OLANZAPINE ODT is used to treat symptoms of schizophrenia and related psychotic disorders as well as those of bipolar disorder.

Your doctor may have prescribed AURO-OLANZAPINE ODT for another reason. Ask your doctor if you have any questions about why AURO-OLANZAPINE ODT has been prescribed for you.

What it does:
AURO-OLANZAPINE ODT belongs to a group of medicines called antipsychotics. AURO-OLANZAPINE ODT is used to treat symptoms of schizophrenia and related psychotic disorders as well as those of bipolar disorder. Schizophrenia may cause symptoms such as hallucinations (e.g. hearing, seeing, or sensing things which are not there), delusions, unusual suspiciousness, feeling withdrawn, lack of emotions. People with schizophrenia may also feel depressed, anxious or tense. Signs and symptoms of bipolar mania include but are not limited to: feeling invincible or all powerful, inflated self-esteem, racing thoughts, easily lose your train of thought, overreaction to what you see or hear, misinterpretation of events, speeded-up activity, talking very quickly, talking too loudly, or talking more than usual, decreased need for sleep, and poor judgment.

When it should not be used:
Do not take AURO-OLANZAPINE ODT if you have had an allergic reaction to AURO-OLANZAPINE ODT or any of the ingredients listed in the "Nonmedicinal Ingredients" section of this leaflet. Signs of allergic reaction may include a skin rash, itching, shortness of breath or swelling of the face, lips or tongue.

What the medicinal ingredient is:
AURO-OLANZAPINE ODT orally disintegrating tablets contain the active ingredient called olanzapine.

What the nonmedicinal ingredients are:
AURO-OLANZAPINE ODT contains the following inactive ingredients: Aspartame, mannitol, polacrilin potassium, crospovidone, colloidal silica anhydrous, microcrystalline cellulose, sodium stearyl fumarate and artificial pineapple flavor.

What dosage forms it comes in:
AURO-OLANZAPINE ODT orally disintegrating tablets are available in 5 mg, 10 mg, 15 mg, and 20 mg strengths.

WARNINGS AND PRECAUTIONS

Serious Warnings and Precautions
Studies with various medicines of the group to which AURO-OLANZAPINE ODT belongs, including AURO-OLANZAPINE ODT, when used in elderly patients with dementia have been associated with an increased rate of death. AURO-OLANZAPINE ODT is not indicated in elderly patients with dementia.

Before starting AURO-OLANZAPINE ODT and to get the best possible treatment, be sure to tell your doctor if you:

- are pregnant or plan to become pregnant
- are breast feeding or plan on breast feeding
- have had an allergic reaction to any medicine which you have taken previously to treat your current condition
- have diabetes or a family history of diabetes
- have a history of any problems with the way your heart beats or have any heart problems
- have a history of stroke or high blood pressure
- have risk factors for developing blood clots such as: a family history of blood clots, age over 65, smoking, obesity, recent major surgery (such as hip or knee replacement), immobility due to air travel

or other reason, or take oral contraceptives ("The Pill").
- are a smoker
- have ever had blackouts or seizures
- are taking any other medicines (prescriptions or over-the-counter medicines)
- drink alcoholic beverages or use drugs
- exercise vigorously or work in hot or sunny places
- have a history of liver problems, hepatitis, or yellowing of the eyes and skin (jaundice)
- have prostate problems
- have intestinal congestion (paralytic ileus)
- have raised pressure within the eye (glaucoma)
- cannot take phenylalanine because AURO-OLANZAPINE ODT contains aspartame, a source of phenylalanine.

It is important for your doctor to have this information before prescribing your treatment and dosage.

Effects on Newborns:
In some cases, babies born to a mother taking AURO-OLANZAPINE ODT during pregnancy have experienced symptoms that are severe and require the newborn to be hospitalized. Sometimes, the symptoms may resolve on their own. Be prepared to seek immediate emergency medical attention for your newborn if they have difficulty breathing, are overly sleepy, have muscle stiffness, or floppy muscles (like a rag doll), are shaking, or are having difficulty feeding.

INTERACTIONS WITH THE MEDICATIONS

Tell all doctors, dentists and pharmacists who are treating you that you are taking AURO-OLANZAPINE ODT.

Tell your doctor or pharmacist that you are taking AURO-OLANZAPINE ODT before you start taking any new medicines.

A combination of AURO-OLANZAPINE ODT with the following medicines might make you feel drowsy:
- medicines taken for anxiety or to help you sleep
- medicines taken for depression.

The effects of alcohol could be made worse while taking AURO-OLANZAPINE ODT. It is recommended that you DO NOT drink alcohol while taking AURO-OLANZAPINE ODT.

You should tell your doctor if you are taking fluvoxamine (antidepressant), ketoconazole (antifungal), or ciprofloxacin (antibiotic), as these medicines may lead to higher concentrations of olanzapine in your blood.

You should also tell your doctor if you are taking carbamazepine as it may lead to lower concentrations of AURO-OLANZAPINE ODT in your blood, making AURO-OLANZAPINE ODT less effective.

Only take other medicines while you are on AURO-OLANZAPINE ODT if your doctor tells you that you can. DO NOT give AURO-OLANZAPINE ODT to anyone else. Your doctor has prescribed it for you and your condition.

PROPER USE OF THIS MEDICATION

Usual dose:
The most important thing about taking AURO-OLANZAPINE ODT is to take it the way your doctor has prescribed - the right dose, every day. Your doctor has decided on the best dosage for you based on your individual situation and needs. Your doctor may increase or decrease your dose depending on your response.

Although AURO-OLANZAPINE ODT cannot cure your condition, it can help relieve your symptoms. If your symptoms improve or disappear, it is probably because your treatment is working. Studies have shown that, after coming off medication, a relapse of symptoms occurs in about 2 out of 3 patients and is more than double that of patients staying on their medication. That is why it is so important to keep taking AURO-OLANZAPINE ODT, even after your symptoms have improved or disappeared. AURO-OLANZAPINE ODT should be taken for as long as you and your doctor believe it is helping you.

Proper Handling Instructions
AURO-OLANZAPINE ODT should be handled carefully with dry hands.

Following the instructions below:
1 Gently push the tablet out from the bottom of the blister.
2 Avoid touching the tablet with your hands. With dry hands, put the tablet directly into your mouth. It will begin to dissolve in your mouth within a few seconds. You can also place the tablet directly into a full glass of water, milk, coffee, orange juice or apple juice. Stir and drink all of the contents immediately.

Overdose:

If you think you have taken too much AURO-OLANZAPINE ODT, contact your healthcare professional, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

Show the health care practitioner your box of tablets. Do this even if there are no signs of discomfort or poisoning. The most common signs if you have taken too much AURO-OLANZAPINE ODT are drowsiness and slurred speech.

Missed Dose:
Take your prescribed dose at the same time each day. If you miss a dose of AURO-OLANZAPINE ODT by a few hours, take the dose when you remember. If most of the day has passed, wait until your next scheduled dose and try not to miss any more.
Do not take 2 doses at once.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Like other medicines, AURO-OLANZAPINE ODT can cause some side effects. These side effects are most likely to be minor and temporary. However, some may be serious and need medical attention. Many of the side effects are dose related, so it is important not to exceed your prescribed dose. The most common side effects of AURO-OLANZAPINE ODT are:

- drowsiness
- weight gain
- dizziness
- increased appetite
- fluid retention
- constipation
- dry mouth
- a feeling of restlessness (akathisia)
- decreased blood pressure upon rising from a lying or sitting position

Events of stuttering (disruptive speech) and increased salivation (salivary hypersecretion) were uncommonly reported. You should also tell your doctor if you notice any symptoms that worry you, even if you think the problems are not connected with the medicine or are not listed here.

Because some people experience drowsiness, you should avoid driving a car or operating machinery until you know how AURO-OLANZAPINE ODT affects you. Some people may feel dizzy in the early stages of treatment, especially when getting up from a lying or sitting position. This side effect usually passes after taking AURO-OLANZAPINE ODT for a few days.

After prolonged use in women, medicines of this type can cause milk secretion or changes in the regularity of their monthly period. On rare occasions, after prolonged use in men, medicines of this type have been associated with breast enlargement. As well, abnormal liver function tests have been reported on occasion.

Your doctor should check your body weight before starting AURO-OLANZAPINE ODT and continue to monitor it for as long as you are being treated.

Your doctor should take blood tests before starting AURO-OLANZAPINE ODT. They will monitor blood sugar, and the number of infection fighting white blood cells. Your doctor should continue to monitor your blood for as long as you are being treated.

If you have high levels of prolactin (measured with a blood test) and a condition called hypogonadism you may be at increased risk of breaking a bone due to osteoporosis. This occurs in both men and women.

Do not be alarmed by this list of possible side effects. You may not experience any of them. If any of these side effects are experienced, they are usually mild and temporary.

The following table is based on data from placebo-controlled clinical trials and from post-marketing data.

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

Symptom / effect	Talk to your healthcare professional		Stop taking drug and get immediate medical help
	Only if severe	In all cases	
Common			
New or worsening constipation		✓	
Uncommon			
Slow heartbeat[1]		✓	
Rare			
Liver inflammation [symptoms of fever, yellow skin or eyes, dark urine, weakness, abdominal pain, nausea, vomiting, loss of appetite, itching][2]		✓*	

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM			
Symptom / effect	**Talk to your healthcare professional**		**Stop taking drug and get immediate medical help**
	Only if severe	**In all cases**	
Low white blood cell count [symptoms of infection, such as cold, flu-like symptoms, fever, sore throat, as well as weakness or general feeling of unwellness][2]		✓	
Rash[2] (see also Allergic Reaction below)	✓		
Seizure [i.e., loss of consciousness with uncontrollable shaking ("fit")][2]			✓*
Very Rare			
Allergic reaction [symptoms include skin rash, hives, swelling, difficulty breathing][2]			✓*
Bruise easily, excessive bleeding[2]		✓	
High fever, muscle rigidity, rapid heartbeat, profuse sweating, irregular pulse[1,2]			✓*
Increased thirst & hunger, frequent urination[1,2]		✓	
Muscle twitching or abnormal movements of the face or tongue[2]		✓*	
Blood clots: swelling, pain and redness in an arm or leg that can be warm to touch. You may develop sudden chest pain, difficulty breathing and heart palpitations[2]		✓	
Pancreas inflammation [symptoms of severe abdominal pain, fever, nausea, vomiting][2]			✓
Long-lasting (greater than 4 hours in duration) and painful erection of the penis[2]			✓
Sudden weakness or numbness in the face, arms, or legs, and speech or vision problems[3]			✓*
Very dark ("tea coloured") urine, muscle tenderness and/or aching[2]			✓

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM			
Symptom / effect	**Talk to your healthcare professional**		**Stop taking drug and get immediate medical help**
	Only if severe	**In all cases**	
Serious skin reactions: (Drug Reaction with Eosinophilia and Systemic Symptoms [DRESS]): skin rash or redness developing into widespread rash with blisters and peeling skin, swollen lymph nodes and fever.			✓

1 Identified from the clinical trial database.

2 Identified from adverse events reported after release onto market.

3 Identified from data from 5 placebo-controlled trials in elderly patients with dementia-related psychosis.

* If you think you have these side effects, it is important that you seek medical advice from your doctor immediately.

This is not a complete list of side effects. For any unexpected effects while taking AURO-OLANZAPINE ODT, contact your doctor or pharmacist.

HOW TO STORE IT

All medicines should be kept out of the reach and sight of children. AURO-OLANZAPINE ODT should be stored in its original package at room temperature (15°C to 30°C), in a dry place and out of direct light. The expiry date of this medicine is printed on the package label. Do not use the medicine after this date. If your doctor tells you to stop taking AURO-OLANZAPINE ODT or you find that they have passed their expiry date, please return any left over medicine to your pharmacist.

Reporting Side Effects

You can report any suspected side effects associated with the use of health products to Health Canada by:

- Visiting the Web page on Adverse Reaction Reporting (https://www.canada.ca/en/health-canada/services/drugs-health-products/medeffect-canada/adverse-reaction-reporting.html) for information on how to report online, by mail or by fax; or
- Calling toll-free at 1-866-234-2345.

NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

If you want more information about AURO-OLANZAPINE ODT:

- Talk to your healthcare professional
- Find the full product monograph that is prepared for healthcare professionals and includes this consumer information by visiting the Health Canada website (https://health-products.canada.ca/dpd-bdpp/index-eng.jsp); the manufacturer's website www.auropharma.ca, or by calling 1-855-648-6681.

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