

# ZIPRASIDONE HYDROCHLORIDE- ziprasidone hydrochloride capsule

## TYA Pharmaceuticals

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### HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use ziprasidone hydrochloride capsules safely and effectively. See full prescribing information for ziprasidone hydrochloride capsules. Ziprasidone Hydrochloride Capsules Initial U.S. Approval: 2001

#### WARNING INCREASED MORTALITY IN ELDERLY PATIENTS WITH DEMENTIA-RELATED PSYCHOSIS

*See full prescribing information for complete boxed warning*

- Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death compared to placebo treatment ( ) 5.1
- Ziprasidone hydrochloride capsules are not approved for elderly patients with dementia-related psychosis ( ) 5.1

#### RECENT MAJOR CHANGES

Warnings and Precautions: Hyperprolactinemia ( ) 5.11

Metabolic Changes ( ) 07/2013 5.5

#### INDICATIONS AND USAGE

Ziprasidone hydrochloride capsules are an atypical antipsychotic. In choosing among treatments, prescribers should be aware of the capacity of ziprasidone hydrochloride capsules to prolong the QT interval and may consider the use of other drugs first ( ) Ziprasidone hydrochloride capsules are indicated as an oral formulation for the: Treatment of schizophrenia. ( ) 5.2

1.1

- Adults: Efficacy was established in four 4 to 6 week trials and one maintenance trial in adult patients with schizophrenia ( ) 14.1

#### DOSAGE AND ADMINISTRATION

Give oral doses with food.

- Schizophrenia: Initiate at 20 mg twice daily. Daily dosage may be adjusted up to 80 mg twice daily. Dose adjustments should occur at intervals of not less than 2 days. Safety and efficacy has been demonstrated in doses up to 100 mg twice daily. The lowest effective dose should be used. ( ) 2.1

#### DOSAGE FORMS AND STRENGTHS

- Capsules: 20 mg, 40 mg, 60 mg, and 80 mg ( ) 3

#### CONTRAINDICATIONS

- Do not use in patients with a known history of QT prolongation ( ) 4.1
- Do not use in patients with recent acute myocardial infarction ( ) 4.1
- Do not use in patients with uncompensated heart failure ( ) 4.1
- Do not use in combination with other drugs that have demonstrated QT prolongation ( ) 4.1
- Do not use in patients with known hypersensitivity to ziprasidone ( ) 4.2

#### WARNINGS AND PRECAUTIONS

- : ziprasidone hydrochloride use should be avoided in patients with bradycardia, hypokalemia or hypomagnesemia, congenital prolongation of the QT interval, or in combination with other drugs that have demonstrated QT

prolongation ( ) *QT Interval Prolongation*:5.2

- Potentially fatal symptom complex has been reported with antipsychotic drugs. Manage with immediate discontinuation of drug and close monitoring. ( ) *Neuroleptic Malignant Syndrome (NMS)*:5.3
- May develop acutely or chronically ( ) *Tardive Dyskinesia*:5.4
- : Atypical antipsychotic drugs have been associated with metabolic changes that may increase cardiovascular/cerebrovascular risk. These metabolic changes include hyperglycemia, dyslipidemia, and weight gain. ( ) *Metabolic Changes*:5.5
- Monitor all patients for symptoms of hyperglycemia including polydipsia, polyuria, polyphagia, and weakness. Patients with DM risk factors should undergo blood glucose testing before and during treatment. ( ) *Hyperglycemia and Diabetes Mellitus (DM)*:5.5
- Undesirable alterations have been observed in patients treated with atypical antipsychotics. ( ) *Dyslipidemia*:5.5
- Weight gain has been reported. Monitor weight gain. ( ) *Weight Gain*:5.5
- Discontinue in patients who develop a rash without an identified cause ( ) *Rash*:5.6
- Use with caution in patients with known cardiovascular or cerebrovascular disease ( ) *Orthostatic Hypotension*:5.7
- has been reported with antipsychotics. Patients with a pre-existing low white blood cell count (WBC) or a history of leukopenia/neutropenia should have their complete blood count (CBC) monitored frequently during the first few months of therapy and should discontinue ziprasidone hydrochloride at the first sign of a decline in WBC in the absence of other causative factors ( ). *Leukopenia, Neutropenia, and Agranulocytosis*:5.8
- Use cautiously in patients with a history of seizures or with conditions that lower seizure threshold ( ) *Seizures*:5.9
- Patients should use caution when operating machinery ( ) *Potential for Cognitive and Motor impairment*:5.10
- Closely supervise high-risk patients ( ) *Suicide*:5.11

#### ----- ADVERSE REACTIONS -----

Commonly observed adverse reactions (incidence  $\geq$ 5% and at least twice the incidence for placebo) were:

- Somnolence, respiratory tract infection ( ) *Schizophrenia*:6.1

**To report SUSPECTED ADVERSE REACTIONS, contact Sandoz Inc. at 1-800-525-8747 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch)**

#### ----- DRUG INTERACTIONS -----

- Ziprasidone should not be used in combination with other drugs that have demonstrated QT prolongation ( , ) 4.17.3
- The absorption of ziprasidone is increased up to two-fold in the presence of food ( ) 7.9
- The full prescribing information contains additional drug interactions ( ). 7

#### ----- USE IN SPECIFIC POPULATIONS -----

- Ziprasidone should be used during pregnancy only if the potential benefit justifies the potential risk ( ). *Pregnancy*:8.1
- Breast feeding is not recommended ( ) *Nursing Mothers*:8.3
- Safety and effectiveness for pediatric patients has not been established ( ) *Pediatric Use*:8.4

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 9/2014

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## FULL PRESCRIBING INFORMATION

### **WARNING - INCREASED MORTALITY IN ELDERLY PATIENTS WITH DEMENTIA-RELATED PSYCHOSIS**

**Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death. Analyses of seventeen placebo-controlled trials (modal duration of 10 weeks), largely in patients taking atypical antipsychotic drugs, revealed a risk of death in drug-treated patients of between 1.6 to 1.7 times the risk of death in placebo-treated patients. Over the course of a typical 10-week controlled trial, the rate of death in drug-treated patients was about 4.5%, compared to a rate of about 2.6% in the placebo group. 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. Observational studies suggest that, similar to atypical antipsychotic drugs, treatment with conventional antipsychotic drugs may increase mortality. The extent to which the findings of increased mortality in observational studies may be attributed to the antipsychotic drug as opposed to some characteristic(s) of the patients is not clear. Ziprasidone hydrochloride capsules are not approved for the treatment of patients with Dementia-Related Psychosis [see WARNINGS AND PRECAUTIONS (5.1)].**

## 1 INDICATIONS AND USAGE

Ziprasidone hydrochloride capsules are indicated for the treatment of schizophrenia. When deciding among the alternative treatments available for the condition needing treatment, the prescriber should consider the finding of ziprasidone's greater capacity to prolong the QT/QTc interval compared to several other antipsychotic drugs [see ]. Prolongation of the QTc interval is associated in some other drugs with the ability to cause torsade de pointes-type arrhythmia, a potentially fatal polymorphic ventricular tachycardia, and sudden death. In many cases this would lead to the conclusion that other drugs should be tried first. Whether ziprasidone will cause torsade de pointes or increase the rate of sudden death is not yet known [see ] **WARNINGS AND PRECAUTIONS ( ) 5.2****WARNINGS AND PRECAUTIONS ( ) 5.2**

### 1.1 Schizophrenia

Ziprasidone hydrochloride capsules are indicated for the treatment of schizophrenia. The efficacy of oral ziprasidone was established in four short-term (4- and 6-week) controlled trials of adult schizophrenic inpatients and in one maintenance trial of stable adult schizophrenic inpatients [see ]. **CLINICAL STUDIES ( ) 14.1**

## 2 DOSAGE AND ADMINISTRATION

### 2.1 Schizophrenia

#### *Dose Selection*

Ziprasidone hydrochloride capsules should be administered at an initial daily dose of 20 mg twice daily with food. In some patients, daily dosage may subsequently be adjusted on the basis of individual clinical status up to 80 mg twice daily. Dosage adjustments, if indicated, should generally occur at intervals of not less than 2 days, as steady-state is achieved within 1 to 3 days. In order to ensure use of the lowest effective dose, patients should ordinarily be observed for improvement for several weeks before upward dosage adjustment.

Efficacy in schizophrenia was demonstrated in a dose range of 20 mg to 100 mg twice daily in short-term, placebo-controlled clinical trials. There were trends toward dose response within the range of 20 mg to 80 mg twice daily, but results were not consistent. An increase to a dose greater than 80 mg twice daily is not generally recommended. The safety of doses above 100 mg twice daily has not been systematically evaluated in clinical trials [see ]. **CLINICAL STUDIES ( ) 14.1**

#### *Maintenance Treatment*

While there is no body of evidence available to answer the question of how long a patient treated with ziprasidone should remain on it, a maintenance study in patients who had been symptomatically stable and then randomized to continue ziprasidone or switch to placebo demonstrated a delay in time to relapse for patients receiving ziprasidone hydrochloride capsules. [see ]. No additional benefit was demonstrated for doses above 20 mg twice daily. Patients should be periodically reassessed to determine the need for maintenance treatment. **CLINICAL STUDIES ( ) 14.1**

### 2.4 Dosing in Special Populations

Dosage adjustments are generally not required on the basis of age, gender, race, or renal or hepatic impairment. Ziprasidone hydrochloride capsules are not approved for use in children or adolescents.  
*Oral:*

## 3 DOSAGE FORMS AND STRENGTHS

Ziprasidone hydrochloride capsules 20 mg, are hard gelatin capsules with opaque yellow cap and opaque blue body, cap imprinted 'SZ 656' with black ink.

Ziprasidone hydrochloride capsules 40 mg, are hard gelatin capsules with opaque blue cap and opaque blue body, cap imprinted 'SZ 657' with white ink.

Ziprasidone hydrochloride capsules 60 mg, are hard gelatin capsules with opaque yellow cap and opaque yellow body, cap imprinted 'SZ 658' with black ink.

Ziprasidone hydrochloride capsules 80 mg, are hard gelatin capsules with opaque yellow cap and opaque blue body, cap imprinted 'SZ 659' with black ink.

## 4 CONTRAINDICATIONS

### 4.1 QT Prolongation

Because of ziprasidone's dose-related prolongation of the QT interval and the known association of fatal arrhythmias with QT prolongation by some other drugs, ziprasidone is contraindicated:

- in patients with a known history of QT prolongation (including congenital long QT syndrome)
- in patients with recent acute myocardial infarction
- in patients with uncompensated heart failure

Pharmacokinetic/pharmacodynamic studies between ziprasidone and other drugs that prolong the QT interval have not been performed. An additive effect of ziprasidone and other drugs that prolong the QT interval cannot be excluded. Therefore, ziprasidone should not be given with:

- dofetilide, sotalol, quinidine, other Class Ia and III anti-arrhythmics, mesoridazine, thioridazine, chlorpromazine, droperidol, pimozide, sparfloxacin, gatifloxacin, moxifloxacin, halofantrine, mefloquine, pentamidine, arsenic trioxide, levomethadyl acetate, dolasetron mesylate, probucol or tacrolimus.
- other drugs that have demonstrated QT prolongation as one of their pharmacodynamic effects and have this effect described in the full prescribing information as a contraindication or a boxed or bolded warning [see ]. **WARNINGS AND PRECAUTIONS ( ) 5.2**

## 4.2 Hypersensitivity

Ziprasidone is contraindicated in individuals with a known hypersensitivity to the product.

## 5 WARNINGS AND PRECAUTIONS

### 5.1 Increased Mortality in Elderly Patients with Dementia-Related Psychosis

**] Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death. Ziprasidone hydrochloride is not approved for the treatment of dementia-related psychosis. [see BOXED WARNING**

### 5.2 QT Prolongation and Risk of Sudden Death

Ziprasidone use should be avoided in combination with other drugs that are known to prolong the QTc interval [see ] Additionally, clinicians should be alert to the identification of other drugs that have been consistently observed to prolong the QTc interval. Such drugs should not be prescribed with ziprasidone. Ziprasidone should also be avoided in patients with congenital long QT syndrome and in patients with a history of cardiac arrhythmias [see ] **CONTRAINDICATIONS ( ), DRUG INTERACTIONS ( ) 4.17.4.CONTRAINDICATIONS ( ) 4.**

A study directly comparing the QT/QTc prolonging effect of oral ziprasidone with several other drugs effective in the treatment of schizophrenia was conducted in patient volunteers. In the first phase of the trial, ECGs were obtained at the time of maximum plasma concentration when the drug was administered alone. In the second phase of the trial, ECGs were obtained at the time of maximum plasma concentration while the drug was co-administered with an inhibitor of the CYP4503A4 metabolism of the drug.

In the first phase of the study, the mean change in QTc from baseline was calculated for each drug, using a sample-based correction that removes the effect of heart rate on the QT interval. The mean increase in QTc from baseline for ziprasidone ranged from approximately 9 to 14 msec greater than for four of the comparator drugs (risperidone, olanzapine, quetiapine, and haloperidol), but was approximately 14 msec less than the prolongation observed for thioridazine.

In the second phase of the study, the effect of ziprasidone on QTc length was not augmented by the presence of a metabolic inhibitor (ketoconazole 200 mg twice daily).

In placebo-controlled trials, oral ziprasidone increased the QTc interval compared to placebo by approximately 10 msec at the highest recommended daily dose of 160 mg. In clinical trials with oral ziprasidone, the electrocardiograms of 2/2,988 (0.06%) patients who received ziprasidone hydrochloride and 1/440 (0.23%) patients who received placebo revealed QTc intervals exceeding the potentially clinically relevant threshold of 500 msec. In the ziprasidone-treated patients, neither case suggested a role of ziprasidone. One patient had a history of prolonged QTc and a screening measurement of 489 msec; QTc was 503 msec during ziprasidone treatment. The other patient had a QTc of 391 msec at the end of treatment with ziprasidone and upon switching to thioridazine

experienced QTc measurements of 518 and 593 msec.

Some drugs that prolong the QT/QTc interval have been associated with the occurrence of torsade de pointes and with sudden unexplained death. The relationship of QT prolongation to torsade de pointes is clearest for larger increases (20 msec and greater) but it is possible that smaller QT/QTc prolongations may also increase risk, or increase it in susceptible individuals. Although torsade de pointes has not been observed in association with the use of ziprasidone in premarketing studies and experience is too limited to rule out an increased risk, there have been rare post-marketing reports (in the presence of multiple confounding factors) [see ] **ADVERSE REACTIONS ( ) 6.2.**

As with other antipsychotic drugs and placebo, sudden unexplained deaths have been reported in patients taking ziprasidone at recommended doses. The premarketing experience for ziprasidone did not reveal an excess risk of mortality for ziprasidone compared to other antipsychotic drugs or placebo, but the extent of exposure was limited, especially for the drugs used as active controls and placebo. Nevertheless, ziprasidone's larger prolongation of QTc length compared to several other antipsychotic drugs raises the possibility that the risk of sudden death may be greater for ziprasidone than for other available drugs for treating schizophrenia. This possibility needs to be considered in deciding among alternative drug products [see ]. **INDICATIONS AND USAGE ( ) 1**

Certain circumstances may increase the risk of the occurrence of torsade de pointes and/or sudden death in association with the use of drugs that prolong the QTc interval, including (1) bradycardia; (2) hypokalemia or hypomagnesemia; (3) concomitant use of other drugs that prolong the QTc interval; and (4) presence of congenital prolongation of the QT interval.

It is recommended that patients being considered for ziprasidone treatment who are at risk for significant electrolyte disturbances, hypokalemia in particular, have baseline serum potassium and magnesium measurements. Hypokalemia (and/or hypomagnesemia) may increase the risk of QT prolongation and arrhythmia. Hypokalemia may result from diuretic therapy, diarrhea, and other causes. Patients with low serum potassium and/or magnesium should be repleted with those electrolytes before proceeding with treatment. It is essential to periodically monitor serum electrolytes in patients for whom diuretic therapy is introduced during ziprasidone treatment. Persistently prolonged QTc intervals may also increase the risk of further prolongation and arrhythmia, but it is not clear that routine screening ECG measures are effective in detecting such patients. Rather, ziprasidone should be avoided in patients with histories of significant cardiovascular illness, e.g., QT prolongation, recent acute myocardial infarction, uncompensated heart failure, or cardiac arrhythmia. Ziprasidone should be discontinued in patients who are found to have persistent QTc measurements >500 msec.

For patients taking ziprasidone who experience symptoms that could indicate the occurrence of torsade de pointes, e.g., dizziness, palpitations, or syncope, the prescriber should initiate further evaluation, e.g., Holter monitoring may be useful.

### **5.3 Neuroleptic Malignant Syndrome (NMS)**

A potentially fatal symptom complex sometimes referred to as Neuroleptic Malignant Syndrome (NMS) has been reported in association with administration of antipsychotic drugs. 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 creatinine phosphokinase, myoglobinuria (rhabdomyolysis), and acute renal failure.

The diagnostic evaluation of patients with this syndrome is complicated. In arriving at a diagnosis, it is important to exclude 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 (CNS) pathology.

The management of NMS should include: (1) immediate discontinuation of antipsychotic drugs and other

drugs not essential to concurrent 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 regimens for NMS.

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

#### **5.4 Tardive Dyskinesia**

A syndrome of potentially irreversible, involuntary, dyskinetic movements may develop in patients undergoing treatment with antipsychotic drugs. Although the prevalence of the syndrome appears to be highest among the elderly, especially elderly women, it is impossible to rely upon prevalence estimates to predict, at the inception of antipsychotic treatment, which patients are likely to develop the syndrome. Whether antipsychotic drug products differ in their potential to cause tardive dyskinesia is unknown.

The risk of developing tardive dyskinesia and the likelihood that it will become irreversible are believed to increase as the duration of treatment and the total cumulative dose of antipsychotic drugs administered to the patient increase. However, the syndrome can develop, although much less commonly, after relatively brief treatment periods at low doses.

There is no known treatment for established cases of tardive dyskinesia, although the syndrome may remit, partially or completely, if antipsychotic treatment is withdrawn. Antipsychotic treatment itself, however, may suppress (or partially suppress) the signs and symptoms of the syndrome, and thereby may possibly mask the underlying process. The effect that symptomatic suppression has upon the long-term course of the syndrome is unknown.

Given these considerations, ziprasidone should be prescribed in a manner that is most likely to minimize the occurrence of tardive dyskinesia. Chronic antipsychotic treatment should generally be reserved for patients who suffer from a chronic illness that (1) is known to respond to antipsychotic drugs, and (2) for whom alternative, equally effective, but potentially less harmful treatments are not available or appropriate. In patients who do require chronic treatment, the smallest dose and the shortest duration of treatment producing a satisfactory clinical response should be sought. The need for continued treatment should be reassessed periodically.

If signs and symptoms of tardive dyskinesia appear in a patient on ziprasidone, drug discontinuation should be considered. However, some patients may require treatment with ziprasidone despite the presence of the syndrome.

#### **5.5 Metabolic Changes**

Atypical antipsychotic drugs have been associated with metabolic changes that may increase cardiovascular/cerebrovascular risk. These metabolic changes include hyperglycemia, dyslipidemia, and body weight gain. While all of the drugs in the class have been shown to produce some metabolic changes, each drug has its own specific risk profile.

##### **Hyperglycemia and Diabetes Mellitus**

Hyperglycemia, and diabetes mellitus, in some cases extreme and associated with ketoacidosis or hyperosmolar coma or death, have been reported in patients treated with atypical antipsychotics. There have been few reports of hyperglycemia or diabetes in patients treated with ziprasidone hydrochloride. Although fewer patients have been treated with ziprasidone hydrochloride, it is not known if this more limited experience is the sole reason for the paucity of such reports. 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 reactions is not completely understood. Precise risk estimates for hyperglycemia-related adverse reactions in patients treated with



atypical antipsychotics are not available.

Patients with an established diagnosis of diabetes mellitus who are started on atypical antipsychotics should be monitored regularly for worsening of glucose control. 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. 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 antidiabetic treatment despite discontinuation of the suspect drug.

Pooled data from short-term, placebo-controlled studies in schizophrenia are presented in Note that for the flexible dose studies in schizophrenia each subject is categorized as having received either low (20 to 40 mg BID) or high (60 to 80 mg BID) dose based on the subject's modal daily dose. In the tables showing categorical changes, the percentages (% column) are calculated as 100x(n/N). **Tables 1 to 2.**

**Table 1: Glucose\* Mean Change from Baseline in Short-Term (up to 6 weeks), Placebo-Controlled, Fixed-Dose, Oral Ziprasidone, Monotherapy Trials in Adult Patients with Schizophrenia**

Mean Random Glucose Change from Baseline mg/dL (N)						
Ziprasidone						Placebo
5 mg BID	20 mg BID	40 mg BID	60 mg BID	80 mg BID	100 mg BID	
-1.1 (N=45)	+2.4 (N=179)	-0.2 (N=146)	-0.5 (N=119)	-1.7 (N=104)	+4.1 (N=85)	+1.4 (N=260)

“Random” glucose measurements—fasting/non-fasting status unknown \*

**Table 2: Glucose\* Categorical Changes in Short-Term (up to 6 weeks), Placebo-Controlled, Fixed-Dose, Oral Ziprasidone, Monotherapy Trials in Adult Patients with Schizophrenia**

Laboratory Analyte	Category Change (at least once) from Baseline	Treatment Arm	N	n(%)
Random Glucose	Normal to High (<100 mg/dL to ≥126 mg/dL)	Ziprasidone	438	77 (17.6%)
		Placebo	169	26 (15.4%)
	Borderline to High (≥100 mg/dL and <126 mg/dL to ≥126 mg/dL)	Ziprasidone	159	54 (34%)
		Placebo	66	22 (33.3%)

“Random” glucose measurements—fasting/non-fasting status unknown In long-term (at least 1 year), placebo-controlled, flexible-dose studies in schizophrenia, the mean change from baseline in random glucose for ziprasidone 20 to 40 mg BID was -3.4 mg/dL (N=122); for ziprasidone 60 to 80 mg BID was +1.3 mg/dL (N=10); and for placebo was +0.3 mg/dL (N=71). \*

### **Dyslipidemia**

Undesirable alterations in lipids have been observed in patients treated with atypical antipsychotics. Pooled data from short-term, placebo-controlled studies in schizophrenia is presented in **Tables 3 to 4.**

**Table 3: Lipid\* Mean Change from Baseline in Short-Term (up to 6 weeks), Placebo-Controlled,**

## Fixed-Dose, Oral Ziprasidone Monotherapy Trials in Adult Patients with Schizophrenia

Laboratory Analyte	Mean Random Glucose Change from Baseline mg/dL (N)						
	Ziprasidone						Placebo
	5 mg BID	20 mg BID	40 mg BID	60 mg BID	80 mg BID	100 mg BID	
Triglycerides	-12.9 (N=45)	-9.6 (N=181)	-17.3 (N=146)	-0.05 (N=120)	-16 (N=104)	+0.8 (N=85)	-18.6 (N=260)
Total Cholesterol	-3.6 (N=45)	-4.4 (N=181)	-8.2 (N=147)	-3.6 (N=120)	-10 (N=104)	-3.6 (N=85)	-4.7 (N=261)

“Random” lipid measurements, fasting/non-fasting status unknown \*

**Table 4: Lipid\* Categorical Changes in Short-Term (up to 6 weeks), Placebo-Controlled, Fixed-Dose, Oral Ziprasidone Monotherapy Trials in Adult Patients with Schizophrenia**

Laboratory Analyte	Category Change (at least once) from Baseline	Treatment Arm	N	n(%)
Triglycerides	Increase by $\geq 50$ mg/dL	Ziprasidone	681	232 (34.1%)
		Placebo	260	53 (20.4%)
	Normal to High (<150 mg/dL to $\geq 200$ mg/dL)	Ziprasidone	429	63 (14.7%)
		Placebo	152	12 (7.9%)
	Borderline to High ( $\geq 150$ mg/dL and <200 mg/dL to $\geq 200$ mg/dL)	Ziprasidone	92	43 (46.7%)
		Placebo	41	12 (29.3%)
Total Cholesterol	Increase by $\geq 40$ mg/dL	Ziprasidone	682	76 (11.1%)
		Placebo	261	26 (10%)
	Normal to High (<200 mg/dL to $\geq 240$ mg/dL)	Ziprasidone	380	15 (3.9%)
		Placebo	145	0 (0%)
	Borderline to High ( $\geq 200$ mg/dL and <240 mg/dL to $\geq 240$ mg/dL)	Ziprasidone	207	56 (27.1%)
		Placebo	82	22 (26.8%)

“Random” lipid measurements, fasting/non-fasting status unknown \*

In long-term (at least 1 year), placebo-controlled, flexible-dose studies in schizophrenia, the mean change from baseline in random triglycerides for ziprasidone 20 to 40 mg BID was +26.3 mg/dL (N=15); for ziprasidone 60 to 80 mg BID was -39.3 mg/dL (N=10); and for placebo was +12.9 mg/dL (N=9). In long-term (at least 1 year), placebo-controlled, flexible-dose studies in schizophrenia, the mean change from baseline in random total cholesterol for ziprasidone 20 to 40 mg BID was +2.5 mg/dL (N=14); for ziprasidone 60 to 80 mg BID was -19.7 mg/dL (N=10); and for placebo was -28 mg/dL (N=9).

## Weight Gain

Weight gain has been observed with atypical antipsychotic use. Monitoring of weight is recommended. Pooled data from short-term, placebo-controlled studies in schizophrenia are presented in **Tables 5**.

**Table 5: Weight Mean Changes in Short-Term (up to 6 weeks), Placebo-Controlled, Fixed-Dose, Oral Ziprasidone Monotherapy Trials in Adult Patients with Schizophrenia**

Ziprasidone						Placebo
5 mg BID	20 mg BID	40 mg BID	60 mg BID	80 mg BID	100 mg BID	
<b>Mean Weight (kg) Changes from Baseline (N)</b>						
+0.3 (N=40)	+1 (N=167)	+1 (N=135)	+0.7 (N=109)	+1.1 (N=97)	+0.9 (N=74)	-0.4 (227)
<b>Proportion of Patients with <math>\geq 7\%</math> Increase in Weight from Baseline (N)</b>						
0% (N=40)	9% (N=167)	10.4% (N=135)	7.3% (N=109)	15.5% (N=97)	10.8% (N=74)	4% (N=227)

In long-term (at least 1 year), placebo-controlled, flexible-dose studies in schizophrenia, the mean change from baseline weight for ziprasidone 20 to 40 mg BID was -2.3 kg (N=124); for ziprasidone 60 to 80 mg BID was +2.5 kg (N=10); and for placebo was -2.9 kg (N=72). In the same long-term studies, the proportion of subjects with  $\geq 7\%$  increase in weight from baseline for ziprasidone 20 to 40 mg BID was 5.6% (N=124); for ziprasidone 60 to 80 mg BID was 20% (N=10), and for placebo was 5.6% (N=72). In a long-term (at least 1 year), placebo-controlled, fixed-dose study in schizophrenia, the mean change from baseline weight for ziprasidone 20 mg BID was -2.6 kg (N=72); for ziprasidone 40 mg BID was -3.3 kg (N=69); for ziprasidone 80 mg BID was -2.8 kg (N=70) and for placebo was -3.8 kg (N=70). In the same long-term fixed-dose schizophrenia study, the proportion of subjects with  $\geq 7\%$  increase in weight from baseline for ziprasidone 20 mg BID was 5.6% (N=72); for ziprasidone 40 mg BID was 2.9% (N=69); for ziprasidone 80 mg BID was 5.7% (N=70) and for placebo was 2.9% (N=70).

## Schizophrenia

The proportions of patients meeting a weight gain criterion of  $\geq 7\%$  of body weight were compared in a pool of four 4- and 6-week placebo-controlled schizophrenia clinical trials, revealing a statistically significantly greater incidence of weight gain for ziprasidone (10%) compared to placebo (4%). A median weight gain of 0.5 kg was observed in ziprasidone patients compared to no median weight change in placebo patients. In this set of clinical trials, weight gain was reported as an adverse reaction in 0.4% and 0.4% of ziprasidone and placebo patients, respectively. During long-term therapy with ziprasidone, a categorization of patients at baseline on the basis of body mass index (BMI) revealed the greatest mean weight gain and highest incidence of clinically significant weight gain ( $>7\%$  of body weight) in patients with low BMI ( $<23$ ) compared to normal (23 to 27) or overweight patients ( $>27$ ). There was a mean weight gain of 1.4 kg for those patients with a “low” baseline BMI, no mean change for patients with a “normal” BMI, and a 1.3 kg mean weight loss for patients who entered the program with a “high” BMI.

## 5.6 Rash

In premarketing trials with ziprasidone, about 5% of patients developed rash and/or urticaria, with discontinuation of treatment in about one-sixth of these cases. The occurrence of rash was related to dose of ziprasidone, although the finding might also be explained by the longer exposure time in the higher dose patients. Several patients with rash had signs and symptoms of associated systemic illness, e.g., elevated WBCs. Most patients improved promptly with adjunctive treatment with antihistamines or steroids and/or upon discontinuation of ziprasidone, and all patients experiencing these reactions were reported to recover completely. Upon appearance of rash for which an alternative etiology cannot be identified, ziprasidone should be discontinued.

## 5.7 Orthostatic Hypotension

Ziprasidone may induce orthostatic hypotension associated with dizziness, tachycardia, and, in some patients, syncope, especially during the initial dose-titration period, probably reflecting its  $\alpha$ 1-adrenergic antagonist properties. Syncope was reported in 0.6% of the patients treated with ziprasidone.

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

## 5.8 Leukopenia, Neutropenia, and Agranulocytosis

In clinical trial and postmarketing experience, events of leukopenia/neutropenia have been reported temporally related to antipsychotic agents. Agranulocytosis (including fatal cases) has also been reported.

Possible risk factors for leukopenia/neutropenia include pre-existing low white blood cell count (WBC) and history of drug induced leukopenia/neutropenia. Patients with a pre-existing low WBC or a history of drug induced leukopenia/neutropenia should have their complete blood count (CBC) monitored frequently during the first few months of therapy and should discontinue ziprasidone hydrochloride at the first sign of decline in WBC in the absence of other causative factors.

Patients with neutropenia should be carefully monitored for fever or other symptoms or signs of infection and treated promptly if such symptoms or signs occur. Patients with severe neutropenia (absolute neutrophil count  $<1000/\text{mm}^3$ ) should discontinue ziprasidone hydrochloride and have their WBC followed until recovery.<sup>3</sup>

## 5.9 Seizures

During clinical trials, seizures occurred in 0.4% of patients treated with ziprasidone. There were confounding factors that may have contributed to the occurrence of seizures in many of these cases. As with other antipsychotic drugs, ziprasidone should be used cautiously in patients with a history of seizures or with conditions that potentially lower the seizure threshold, e.g., Alzheimer's dementia. Conditions that lower the seizure threshold may be more prevalent in a population of 65 years or older.

## 5.10 Dysphagia

Esophageal dysmotility and aspiration have been associated with antipsychotic drug use. Aspiration pneumonia is a common cause of morbidity and mortality in elderly patients, in particular those with advanced Alzheimer's dementia. Ziprasidone and other antipsychotic drugs should be used cautiously in patients at risk for aspiration pneumonia [see ] **BOXED WARNING**.

## 5.11 Hyperprolactinemia

As with other drugs that antagonize dopamine D receptors, ziprasidone elevates prolactin levels in humans. Increased prolactin levels were also observed in animal studies with this compound, and were associated with an increase in mammary gland neoplasia in mice; a similar effect was not observed in rats [see ( )]. Tissue culture experiments indicate that approximately one-third of human breast cancers are prolactin-dependent, a factor of potential importance if the prescription of these drugs is contemplated in a patient with previously detected breast cancer. Neither clinical studies nor epidemiologic studies conducted to date have shown an association between chronic administration of this class of drugs and tumorigenesis in humans; the available evidence is considered too limited to be conclusive at this time. **NONCLINICAL TOXICOLOGY** 13.1 *in vitro*

Although disturbances such as galactorrhea, amenorrhea, gynecomastia, and impotence have been reported with prolactin-elevating compounds, the clinical significance of elevated serum prolactin levels is unknown for most patients. Long-standing hyperprolactinemia when associated with

hypogonadism may lead to decreased bone density.

### **5.12 Potential for Cognitive and Motor Impairment**

Somnolence was a commonly reported adverse reaction in patients treated with ziprasidone. In the 4- and 6-week placebo-controlled trials, somnolence was reported in 14% of patients on ziprasidone compared to 7% of placebo patients. Somnolence led to discontinuation in 0.3% of patients in short-term clinical trials. Since ziprasidone has the potential to impair judgment, thinking, or motor skills, patients should be cautioned about performing activities requiring mental alertness, such as operating a motor vehicle (including automobiles) or operating hazardous machinery until they are reasonably certain that ziprasidone therapy does not affect them adversely.

### **5.13 Priapism**

One case of priapism was reported in the premarketing database. While the relationship of the reaction to ziprasidone use has not been established, other drugs with alpha-adrenergic blocking effects have been reported to induce priapism, and it is possible that ziprasidone may share this capacity. Severe priapism may require surgical intervention.

### **5.14 Body Temperature Regulation**

Although not reported with ziprasidone in premarketing trials, disruption of the body's ability to reduce core body temperature has been attributed to antipsychotic agents. Appropriate care is advised when prescribing ziprasidone for patients who will be experiencing conditions which may contribute to an elevation in core body temperature, e.g., exercising strenuously, exposure to extreme heat, receiving concomitant medication with anticholinergic activity, or being subject to dehydration.

### **5.15 Suicide**

The possibility of a suicide attempt is inherent in psychotic illness and close supervision of high-risk patients should accompany drug therapy. Prescriptions for ziprasidone should be written for the smallest quantity of capsules consistent with good patient management in order to reduce the risk of overdose.

### **5.16 Patients with Concomitant illnesses**

Clinical experience with ziprasidone in patients with certain concomitant systemic illnesses is limited [see ]. **USE IN SPECIFIC POPULATIONS ( ), ( ) 8.68.7**

Ziprasidone has not been evaluated or used to any appreciable extent in patients with a recent history of myocardial infarction or unstable heart disease.

Patients with these diagnoses were excluded from premarketing clinical studies. Because of the risk of QTc prolongation and orthostatic hypotension with ziprasidone, caution should be observed in cardiac patients [see ]. **WARNINGS AND PRECAUTIONS ( ), ( ) 5.25.7**

### **5.17 Laboratory Tests**

Patients being considered for ziprasidone treatment that are at risk of significant electrolyte disturbances should have baseline serum potassium and magnesium measurements. Low serum potassium and magnesium should be replaced before proceeding with treatment. Patients who are started on diuretics during Ziprasidone therapy need periodic monitoring of serum potassium and magnesium. Ziprasidone should be discontinued in patients who are found to have persistent QTc measurements >500 msec. [see ] **WARNINGS AND PRECAUTIONS ( ) 5.2**

## **6 ADVERSE REACTIONS**

### **6.1 Clinical Trials Experience**

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Clinical trials for oral ziprasidone included approximately 5,700 patients and/or normal subjects exposed to one or more doses of ziprasidone. Of these 5700, over 4800 were patients who participated in multiple-dose effectiveness trials, and their experience corresponded to approximately 1,831 patient-years. These patients include: (1) 4,331 patients who participated in multiple-dose trials, predominantly in schizophrenia, representing approximately 1698 patient-years of exposure as of February 5, 2000. The conditions and duration of treatment with ziprasidone included open-label and double-blind studies, inpatient and outpatient studies, and short-term and longer-term exposure.

Adverse reactions during exposure were obtained by collecting voluntarily reported adverse experiences, as well as results of physical examinations, vital signs, weights, laboratory analyses, ECGs, and results of ophthalmologic examinations.

The stated frequencies of adverse reactions represent the proportion of individuals who experienced, at least once, a treatment-emergent adverse reaction of the type listed. A reaction was considered treatment emergent if it occurred for the first time or worsened while receiving therapy following baseline evaluation.

### ***Adverse Findings Observed in Short-Term, Placebo-Controlled Trials with Oral Ziprasidone***

The following findings are based on the short-term placebo-controlled premarketing trials for schizophrenia (a pool of two 6-week, and two 4-week fixed-dose trials) in which ziprasidone was administered in doses ranging from 10 to 200 mg/day.

### ***Commonly Observed Adverse Reactions in Short Term-Placebo-Controlled Trials***

The following adverse reactions were the most commonly observed adverse reactions associated with the use of ziprasidone (incidence of 5% or greater) and not observed at an equivalent incidence among placebo-treated patients (ziprasidone incidence at least twice that for placebo):

Schizophrenia trials (*see Table 6*)

- Somnolence
- Respiratory Tract Infection

## **SCHIZOPHRENIA**

### **Adverse Reactions Associated with Discontinuation of Treatment in Short-Term, Placebo-Controlled Trials of Oral Ziprasidone**

Approximately 4.1% (29/702) of ziprasidone-treated patients in short-term, placebo-controlled studies discontinued treatment due to an adverse reaction, compared with about 2.2% (6/273) on placebo. The most common reaction associated with dropout was rash, including 7 dropouts for rash among ziprasidone patients (1%) compared to no placebo patients [See ]. **WARNINGS AND PRECAUTIONS ( ) 5.6**

### **Adverse Reactions Occurring at an Incidence of 2% or More Among Ziprasidone-Treated Patients in Short-Term, Oral, Placebo-Controlled Trials**

enumerates the incidence, rounded to the nearest percent, of treatment-emergent adverse reactions that occurred during acute therapy (up to 6 weeks) in predominantly patients with schizophrenia, including only those reactions that occurred in 2% or more of patients treated with ziprasidone and for which the incidence in patients treated with ziprasidone was greater than the incidence in placebo-treated patients.

### **Table 6**

### **Table 6 Treatment-Emergent Adverse Reaction Incidence In Short-Term Oral Placebo-**

## Controlled Trials - Schizophrenia

Body System/Adverse Reaction	Percentage of Patients Reporting Reaction	
	Ziprasidone (N=702)	Placebo (N=273)
<b>Body as a Whole</b>		
Asthenia	5	3
Accidental Injury	4	2
Chest Pain	3	2
<b>Cardiovascular</b>		
Tachycardia	2	1
<b>Digestive</b>		
Nausea	10	7
Constipation	9	8
Dyspepsia	8	7
Diarrhea	5	4
Dry Mouth	4	2
Anorexia	2	1
<b>Nervous</b>		
Extrapyramidal Symptoms *	14	8
Somnolence	14	7
Akathisia	8	7
Dizziness †	8	6
<b>Respiratory</b>		
Respiratory Tract Infection	8	3
Rhinitis	4	2
Cough Increased	3	1
<b>Skin and Appendages</b>		
Rash	4	3
Fungal Dermatitis	2	1
<b>Special Senses</b>		
Abnormal Vision	3	2

\* Extrapyramidal Symptoms includes the following adverse reaction terms: extrapyramidal syndrome, hypertonia, dystonia, dyskinesia, hypokinesia, tremor, paralysis and twitching. None of these adverse reactions occurred individually at an incidence greater than 5% in schizophrenia trials.

† Dizziness includes the adverse reaction terms dizziness and lightheadedness.

### Dose Dependency of Adverse Reactions in Short-Term, Fixed-Dose, Placebo-Controlled Trials

An analysis for dose response in the schizophrenia 4-study pool revealed an apparent relation of adverse reaction to dose for the following reactions: asthenia, postural hypotension, anorexia, dry mouth, increased salivation, arthralgia, anxiety, dizziness, dystonia, hypertonia, somnolence, tremor, rhinitis, rash, and abnormal vision.

### Extrapyramidal Symptoms (EPS)

The incidence of reported EPS (which included the adverse reaction terms extrapyramidal syndrome, hypertonia, dystonia, dyskinesia, hypokinesia, tremor, paralysis and twitching) for ziprasidone-treated patients in the short-term, placebo-controlled schizophrenia trials was 14% vs. 8% for placebo. Objectively collected data from those trials on the Simpson-Angus Rating Scale (for EPS) and the Barnes Akathisia Scale (for akathisia) did not generally show a difference between ziprasidone and placebo.

## **Dys tonia**

Symptoms of dystonia, prolonged abnormal contractions of muscle groups, may occur in susceptible individuals during the first few days of treatment. Dystonic symptoms include: spasm of the neck muscles, sometimes progressing to tightness of the throat, swallowing difficulty, difficulty breathing, and/or protrusion of the tongue. While these symptoms can occur at low doses, they occur more frequently and with greater severity with high potency and at higher doses of first generation antipsychotic drugs. An elevated risk of acute dystonia is observed in males and younger age groups.  
*Class Effect:*

## **Vital Sign Changes**

Ziprasidone is associated with orthostatic hypotension [see **WARNINGS AND PRECAUTIONS ( )**].  
**5.7**

## **ECG Changes**

Ziprasidone is associated with an increase in the QTc interval [see ] In the schizophrenia trials, ziprasidone was associated with a mean increase in heart rate of 1.4 beats per minute compared to a 0.2 beats per minute decrease among placebo patients. **WARNINGS AND PRECAUTIONS ( ) 5.2.**

## ***Other Adverse Reactions Observed During the Premarketing Evaluation of Oral Ziprasidone***

Following is a list of COSTART terms that reflect treatment-emergent adverse reactions as defined in the introduction to the section reported by patients treated with ziprasidone in schizophrenia trials at multiple doses >4 mg/day within the database of 3,834 patients. All reported reactions are included except those already listed in or elsewhere in labeling, those reaction terms that were so general as to be uninformative, reactions reported only once and that did not have a substantial probability of being acutely life-threatening, reactions that are part of the illness being treated or are otherwise common as background reactions, and reactions considered unlikely to be drug-related. It is important to emphasize that, although the reactions reported occurred during treatment with ziprasidone, they were not necessarily caused by it. **ADVERSE REACTION Table 6**

Adverse reactions are further categorized by body system and listed in order of decreasing frequency according to the following definitions:

adverse reactions occurring in at least 1/100 patients ( $\geq 1\%$  of patients) (only those not already listed in the tabulated results from placebo-controlled trials appear in this listing); **Frequent -**

adverse reactions occurring in 1/100 to 1/1000 patients (in 0.1 to 1% of patients) **Infrequent-**

adverse reactions occurring in fewer than 1/1000 patients (<0.1% of patients). - **Rare**

## **Body as a Whole**

abdominal pain, flu syndrome, fever, accidental fall, face edema, chills, photosensitivity reaction, flank pain, hypothermia, motor vehicle accident - **Frequent**

## **Cardiovascular System**

tachycardia, hypertension, postural hypotension - **Frequent**

bradycardia, angina pectoris, atrial fibrillation **Infrequent -**

first degree AV block, bundle branch block, phlebitis, pulmonary embolus, cardiomegaly, cerebral infarct, cerebrovascular accident, deep thrombophlebitis, myocarditis, thrombophlebitis **Rare-**

## **Digestive System**

anorexia, vomiting **Frequent -**

rectal hemorrhage, dysphagia, tongue edema **Infrequent -**

gum hemorrhage, jaundice, fecal impaction, gamma glutamyl transpeptidase increased, hematemesis,



cholestatic jaundice, hepatitis, hepatomegaly, leukoplakia of mouth, fatty liver deposit, melena - **Rare**

### **Endocrine**

hypothyroidism, hyperthyroidism, thyroiditis **Rare** -

### **Hemic and Lymphatic System**

anemia, ecchymosis, leukocytosis, leukopenia, eosinophilia, lymphadenopathy **Infrequent** -  
thrombocytopenia, hypochromic anemia, lymphocytosis, monocytosis, basophilia, lymphedema,  
polycythemia, thrombocytopenia - **Rare**

### **Metabolic and Nutritional Disorders**

thirst, transaminase increased, peripheral edema, hyperglycemia, creatine phosphokinase increased,  
alkaline phosphatase increased, hypercholesteremia, dehydration, lactic dehydrogenase increased,  
albuminuria, hypokalemia - **Infrequent**

BUN increased, creatinine increased, hyperlipemia, hypocholesteremia, hyperkalemia, hypochloremia,  
hypoglycemia, hyponatremia, hypoproteinemia, glucose tolerance decreased, gout, hyperchloremia,  
hyperuricemia, hypocalcemia, hypoglycemic reaction, hypomagnesemia, ketosis, respiratory alkalosis -  
**Rare**

### **Musculoskeletal System**

myalgia - **Frequent**

tenosynovitis - **Infrequent**

myopathy **Rare**-

### **Nervous System**

agitation, extrapyramidal syndrome, tremor, dystonia, hypertonia, dyskinesia, hostility, twitching,  
paresthesia, confusion, vertigo, hypokinesia, hyperkinesia, abnormal gait, oculogyric crisis,  
hypesthesia, ataxia, amnesia, cogwheel rigidity, delirium, hypotonia, akinesia, dysarthria, withdrawal  
syndrome, buccoglossal syndrome, choreoathetosis, diplopia, incoordination, neuropathy **Frequent** -  
paralysis **Infrequent** -

myoclonus, nystagmus, torticollis, circumoral paresthesia, opisthotonos, reflexes increased, trismus  
**Rare**-

### **Respiratory System**

dyspnea **Frequent** -

pneumonia, epistaxis **Infrequent** -

hemoptysis, laryngismus **Rare** -

### **Skin and Appendages**

maculopapular rash, urticaria, alopecia, eczema, exfoliative dermatitis, contact dermatitis,  
vesiculobullous rash - **Infrequent**

### **Special Senses**

fungal dermatitis **Frequent**-

conjunctivitis, dry eyes, tinnitus, blepharitis, cataract, photophobia **Infrequent**-

eye hemorrhage, visual field defect, keratitis, keratoconjunctivitis - **Rare**

### **Urogenital System**

impotence, abnormal ejaculation, amenorrhea, hematuria, menorrhagia, female lactation, polyuria,

urinary retention metrorrhagia, male sexual dysfunction, anorgasmia, glycosuria **Infrequent** -  
gynecomastia, vaginal hemorrhage, nocturia, oliguria, female sexual dysfunction, uterine hemorrhage  
**Rare-**

## 6.2 Postmarketing Experience

The following adverse reactions have been identified during post approval use of ziprasidone hydrochloride. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Adverse reaction reports not listed above that have been received since market introduction include rare occurrences of the following : Tachycardia, torsade de pointes (in the presence of multiple confounding factors), [See ] Swollen Tongue; Galactorrhea, priapism; Facial Droop, neuroleptic malignant syndrome, serotonin syndrome (alone or in combination with serotonergic medicinal products), tardive dyskinesia; Insomnia, mania/hypomania; Allergic reaction (such as allergic dermatitis, angioedema, orofacial edema, urticaria), rash; Enuresis, urinary incontinence; Postural hypotension, syncope. **Cardiac Disorders: WARNINGS AND PRECAUTIONS ( ) 5.2; Digestive System Disorders: Reproductive System and Breast Disorders: Nervous System Disorders: Psychiatric Disorders: Skin and subcutaneous Tissue Disorders: Urogenital System Disorders: Vascular Disorders:**

## 7 DRUG INTERACTIONS

Drug-drug interactions can be pharmacodynamic (combined pharmacologic effects) or pharmacokinetic (alteration of plasma levels). The risks of using ziprasidone in combination with other drugs have been evaluated as described below. All interactions studies have been conducted with oral ziprasidone. Based upon the pharmacodynamic and pharmacokinetic profile of ziprasidone, possible interactions could be anticipated:

### 7.1 Metabolic Pathway

Approximately two-thirds of ziprasidone is metabolized via a combination of chemical reduction by glutathione and enzymatic reduction by aldehyde oxidase. There are no known clinically relevant inhibitors or inducers of aldehyde oxidase. Less than one-third of ziprasidone metabolic clearance is mediated by cytochrome P450 catalyzed oxidation.

### 7.2 In Vitro Studies

An enzyme inhibition study utilizing human liver microsomes showed that ziprasidone had little inhibitory effect on CYP1A2, CYP2C9, CYP2C19, CYP2D6 and CYP3A4, and thus would not likely interfere with the metabolism of drugs primarily metabolized by these enzymes. There is little potential for drug interactions with ziprasidone due to displacement [see ] *in vitro* **CLINICAL PHARMACOLOGY ( ) 12.3.**

### 7.3 Pharmacodynamic Interactions

Ziprasidone should not be used with any drug that prolongs the QT interval [see ] **CONTRAINDICATIONS ( ) 4.1.**

Given the primary CNS effects of ziprasidone, caution should be used when it is taken in combination with other centrally acting drugs. Because of its potential for inducing hypotension, ziprasidone may enhance the effects of certain antihypertensive agents.

Ziprasidone may antagonize the effects of levodopa and dopamine agonists.

### 7.4 Pharmacokinetic Interactions

**Carbamazepine**

Carbamazepine is an inducer of CYP3A4; administration of 200 mg twice daily for 21 days resulted in a decrease of approximately 35% in the AUC of ziprasidone. This effect may be greater when higher doses of carbamazepine are administered.

### ***Ketoconazole***

Ketoconazole, a potent inhibitor of CYP3A4, at a dose of 400 mg QD for 5 days, increased the AUC and C<sub>max</sub> of ziprasidone by about 35 to 40%. Other inhibitors of CYP3A4 would be expected to have similar effects.

### ***Cimetidine***

Cimetidine at a dose of 800 mg QD for 2 days did not affect ziprasidone pharmacokinetics.

### ***Antacid***

The co-administration of 30 mL of Maalox with ziprasidone did not affect the pharmacokinetics of ziprasidone.®

## **7.5 Lithium**

Ziprasidone at a dose of 40 mg twice daily administered concomitantly with lithium at a dose of 450 mg twice daily for 7 days did not affect the steady-state level or renal clearance of lithium.

## **7.6 Oral Contraceptives**

Studies have revealed no effect of ziprasidone on the pharmacokinetics of estrogen or progesterone components. Ziprasidone at a dose of 20 mg twice daily did not affect the pharmacokinetics of concomitantly administered oral contraceptives, ethinyl estradiol (0.03 mg) and levonorgestrel (0.15 mg). *In vivo*

## **7.7 Dextromethorphan**

Consistent with results, a study in normal healthy volunteers showed that ziprasidone did not alter the metabolism of dextromethorphan, a CYP2D6 model substrate, to its major metabolite, dextrorphan. There was no statistically significant change in the urinary dextromethorphan/dextrorphan ratio. *in vitro*

## **7.8 Valproate**

A pharmacokinetic interaction of ziprasidone with valproate is unlikely due to the lack of common metabolic pathways for the two drugs.

## **7.9 Other Concomitant Drug Therapy**

Population pharmacokinetic analysis of schizophrenic patients enrolled in controlled clinical trials has not revealed evidence of any clinically significant pharmacokinetic interactions with benztropine, propranolol, or lorazepam.

## **7.10 Food Interaction**

The absolute bioavailability of a 20 mg dose under fed conditions is approximately 60%. The absorption of ziprasidone is increased up to two-fold in the presence of food [see ]. **CLINICAL PHARMACOLOGY ( ) 12.3**

# **8 USE IN SPECIFIC POPULATIONS**

## **8.1 Pregnancy**

### ***Pregnancy Category C***

In animal studies ziprasidone demonstrated developmental toxicity, including possible teratogenic

effects at doses similar to human therapeutic doses. When ziprasidone was administered to pregnant rabbits during the period of organogenesis, an increased incidence of fetal structural abnormalities (ventricular septal defects and other cardiovascular malformations and kidney alterations) was observed at a dose of 30 mg/kg/day (3 times the MRHD of 200 mg/day on a mg/m basis). There was no evidence to suggest that these developmental effects were secondary to maternal toxicity. The developmental no-effect dose was 10 mg/kg/day (equivalent to the MRHD on a mg/m basis). In rats, embryofetal toxicity (decreased fetal weights, delayed skeletal ossification) was observed following administration of 10 to 160 mg/kg/day (0.5 to 8 times the MRHD on a mg/m basis) during organogenesis or throughout gestation, but there was no evidence of teratogenicity. Doses of 40 and 160 mg/kg/day (2 and 8 times the MRHD on a mg/m basis) were associated with maternal toxicity. The developmental no-effect dose was 5 mg/kg/day (0.2 times the MRHD on a mg/m basis).<sup>22222</sup>

There was an increase in the number of pups born dead and a decrease in postnatal survival through the first 4 days of lactation among the offspring of female rats treated during gestation and lactation with doses of 10 mg/kg/day (0.5 times the MRHD on a mg/m basis) or greater. Offspring developmental delays and neurobehavioral functional impairment were observed at doses of 5 mg/kg/day (0.2 times the MRHD on a mg/m basis) or greater. A no-effect level was not established for these effects.<sup>22</sup>

There are no adequate and well-controlled studies in pregnant women. Ziprasidone should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

### ***Non-teratogenic Effects***

Neonates exposed to antipsychotic drugs, 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.

Ziprasidone hydrochloride should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

## **8.2 Labor and Delivery**

The effect of ziprasidone on labor and delivery in humans is unknown.

## **8.3 Nursing Mothers**

It is not known whether ziprasidone or its metabolites are excreted in human milk. It is recommended that women receiving ziprasidone should not breastfeed.

## **8.4 Pediatric Use**

The safety and effectiveness of ziprasidone in pediatric patients have not been established.

## **8.5 Geriatric Use**

Of the total number of subjects in clinical studies of ziprasidone, 2.4 percent were 65 and over. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly and younger patients, but greater sensitivity of some older individuals cannot be ruled out. Nevertheless, the presence of multiple factors that might increase the pharmacodynamic response to ziprasidone, or cause poorer tolerance or orthostasis, should lead to consideration of a lower starting dose, slower titration, and careful monitoring during the initial dosing period for some elderly patients.

## **8.6 Renal Impairment**

Because ziprasidone is highly metabolized, with less than 1% of the drug excreted unchanged, renal impairment alone is unlikely to have a major impact on the pharmacokinetics of ziprasidone. The

pharmacokinetics of ziprasidone following 8 days of 20 mg twice daily dosing were similar among subjects with varying degrees of renal impairment (n=27), and subjects with normal renal function, indicating that dosage adjustment based upon the degree of renal impairment is not required. Ziprasidone is not removed by hemodialysis.

### **8.7 Hepatic Impairment**

As ziprasidone is cleared substantially by the liver, the presence of hepatic impairment would be expected to increase the AUC of ziprasidone; a multiple-dose study at 20 mg twice daily for 5 days in subjects (n=13) with clinically significant (Childs-Pugh Class A and B) cirrhosis revealed an increase in AUC of 13% and 34% in Childs-Pugh Class A and B, respectively, compared to a matched control group (n=14). A half-life of 7.1 hours was observed in subjects with cirrhosis compared to 4.8 hours in the control group. 0-12

### **8.8 Age and Gender Effects**

In a multiple-dose (8 days of treatment) study involving 32 subjects, there was no difference in the pharmacokinetics of ziprasidone between men and women or between elderly (>65 years) and young (18 to 45 years) subjects. Additionally, population pharmacokinetic evaluation of patients in controlled trials has revealed no evidence of clinically significant age or gender-related differences in the pharmacokinetics of ziprasidone. Dosage modifications for age or gender are, therefore, not recommended.

### **8.9 Smoking**

Based on studies utilizing human liver enzymes, ziprasidone is not a substrate for CYP1A2; smoking should therefore not have an effect on the pharmacokinetics of ziprasidone. Consistent with these results, population pharmacokinetic evaluation has not revealed any significant pharmacokinetic differences between smokers and nonsmokers. *in vitro*

## **9 DRUG ABUSE AND DEPENDENCE**

### **9.3 Dependence**

Ziprasidone has not been systematically studied, in animals or humans, for its potential for abuse, tolerance, or physical dependence. While the clinical trials did not reveal any tendency for drug-seeking behavior, these observations were not systematic and it is not possible to predict on the basis of this limited experience the extent to which ziprasidone will be misused, diverted, and/or abused once marketed. Consequently, patients should be evaluated carefully for a history of drug abuse, and such patients should be observed closely for signs of ziprasidone misuse or abuse (e.g., development of tolerance, increases in dose, drug-seeking behavior).

## **10 OVERDOSAGE**

### **10.1 Human Experience**

In premarketing trials involving more than 5,400 patients and/or normal subjects, accidental or intentional overdosage of oral ziprasidone was documented in 10 patients. All of these patients survived without sequelae. In the patient taking the largest confirmed amount, 3,240 mg, the only symptoms reported were minimal sedation, slurring of speech, and transitory hypertension (200/95).

Adverse reactions reported with ziprasidone overdose included extrapyramidal symptoms, somnolence, tremor, and anxiety. [see ] **ADVERSE REACTIONS ( ) 6.2**

### **10.2 Management of Overdosage**

In case of acute overdosage, establish and maintain an airway and ensure adequate oxygenation and

ventilation. Intravenous access should be established, and gastric lavage (after intubation, if patient is unconscious) and administration of activated charcoal together with a laxative should be considered. The possibility of obtundation, seizure, or dystonic reaction of the head and neck following overdose may create a risk of aspiration with induced emesis.

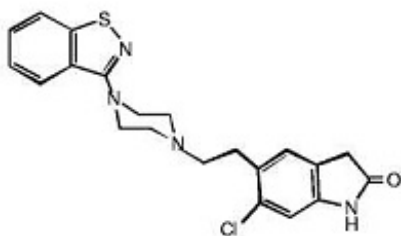
Cardiovascular monitoring should commence immediately and should include continuous electrocardiographic monitoring to detect possible arrhythmias. If antiarrhythmic therapy is administered, disopyramide, procainamide, and quinidine carry a theoretical hazard of additive QT-prolonging effects that might be additive to those of ziprasidone.

Hypotension and circulatory collapse should be treated with appropriate measures such as intravenous fluids. If sympathomimetic agents are used for vascular support, epinephrine and dopamine should not be used, since beta stimulation combined with  $\alpha_1$  antagonism associated with ziprasidone may worsen hypotension. Similarly, it is reasonable to expect that the alpha-adrenergic-blocking properties of bretylium might be additive to those of ziprasidone, resulting in problematic hypotension.

In cases of severe extrapyramidal symptoms, anticholinergic medication should be administered. There is no specific antidote to ziprasidone, and it is not dialyzable. The possibility of multiple drug involvement should be considered. Close medical supervision and monitoring should continue until the patient recovers.

## 11 DESCRIPTION

Ziprasidone hydrochloride capsules are available for oral administration. Ziprasidone is a psychotropic agent that is chemically unrelated to phenothiazine or butyrophenone antipsychotic agents. It has a molecular weight of 412.94 (free base), with the following chemical name: 5-[2-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]ethyl]-6-chloro-1,3-dihydro-2-indol-2-one. The molecular formula of  $C_{17}H_{16}ClN_2OS$  (free base of ziprasidone) represents the following structural formula:  $H_{21214}$



Ziprasidone hydrochloride capsules contain a monohydrochloride, anhydrous salt of ziprasidone. Chemically, ziprasidone hydrochloride anhydrous is 5-[2-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]ethyl]-6-chloro-1,3-dihydro-2-indol-2-one, monohydrochloride, anhydrous. The molecular formula is  $C_{17}H_{16}ClN_2OS \cdot HCl$  and its molecular weight is 449.4. Ziprasidone hydrochloride anhydrous is a light pink to dark pink colored powder.  $H_{21214}$

Ziprasidone hydrochloride capsules are supplied for oral administration in 20 mg (yellow/blue), 40 mg (blue/blue), 60 mg (yellow/yellow), and 80 mg (yellow/blue) capsules. Ziprasidone hydrochloride capsule, intended for oral administration, contains ziprasidone hydrochloride anhydrous equivalent to 20 mg, 40 mg, 60 mg or 80 mg of ziprasidone. In addition, each capsule contains the following inactive ingredients: calcium silicate, citric acid, lactose anhydrous, magnesium stearate and pregelatinized starch.

The capsule shell for the 20 mg and 80 mg strengths consists of carboxymethylcellulose, FD&C blue # 2, gelatin, iron oxide yellow, sodium lauryl sulfate and titanium dioxide. The capsule is printed with edible black pharmaceutical ink.

The capsule shell for the 40 mg strength consists of FD&C blue # 2, gelatin, sodium lauryl sulfate and titanium dioxide. The capsule is printed with edible white pharmaceutical ink.

The capsule shell for the 60 mg strength consists of carboxymethylcellulose, gelatin, iron oxide yellow, sodium lauryl sulfate and titanium dioxide. The capsule is printed with edible black pharmaceutical ink.

The black ink is comprised of iron oxide black, potassium hydroxide, propylene glycol, shellac and ammonia solution.

The white ink is comprised of povidone, propylene glycol, sodium hydroxide, shellac and titanium dioxide.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

The mechanism of action of ziprasidone, as with other drugs having efficacy in schizophrenia, is unknown. However, it has been proposed that this drug's efficacy in schizophrenia is mediated through a combination of dopamine type 2 (D<sub>2</sub>) and serotonin type 2 (5HT<sub>2</sub>) antagonism. 22

### 12.2 Pharmacodynamics

Ziprasidone exhibited high binding affinity for the dopamine D<sub>1</sub> and D<sub>2</sub>, the serotonin 5HT<sub>1A</sub>, 5HT<sub>1B</sub>, 5HT<sub>2A</sub>, 5HT<sub>2B</sub>, and  $\alpha$ -adrenergic receptors (K<sub>s</sub> of 4.8, 7.2, 0.4, 1.3, 3.4, 2, and 10 nM, respectively), and moderate affinity for the histamine H<sub>1</sub> receptor (K<sub>s</sub> =47 nM). Ziprasidone functioned as an antagonist at the D<sub>1</sub>, 5HT<sub>2A</sub>, and 5HT<sub>2B</sub> receptors, and as an agonist at the 5HT<sub>1A</sub> receptor. Ziprasidone inhibited synaptic reuptake of serotonin and norepinephrine. No appreciable affinity was exhibited for other receptor/binding sites tested, including the cholinergic muscarinic receptor (IC<sub>50</sub> >1  $\mu$ M). Antagonism at receptors other than dopamine and 5HT with similar receptor affinities may explain some of the other therapeutic and side effects of ziprasidone. Ziprasidone's antagonism of histamine H<sub>1</sub> receptors may explain the somnolence observed with this drug. Ziprasidone's antagonism of  $\alpha$ -adrenergic receptors may explain the orthostatic hypotension observed with this drug. *in vitro* 232A2C1A1D1i1i2A1D1A50211

### 12.3 Pharmacokinetics

#### Oral Pharmacokinetics

Ziprasidone's activity is primarily due to the parent drug. The multiple-dose pharmacokinetics of ziprasidone are dose-proportional within the proposed clinical dose range, and ziprasidone accumulation is predictable with multiple dosing. Elimination of ziprasidone is mainly via hepatic metabolism with a mean terminal half-life of about 7 hours within the proposed clinical dose range. Steady-state concentrations are achieved within one to three days of dosing. The mean apparent systemic clearance is 7.5 mL/min/kg. Ziprasidone is unlikely to interfere with the metabolism of drugs metabolized by cytochrome P450 enzymes.

#### Absorption

Ziprasidone is well absorbed after oral administration, reaching peak plasma concentrations in 6 to 8 hours. The absolute bioavailability of a 20 mg dose under fed conditions is approximately 60%. The absorption of ziprasidone is increased up to two-fold in the presence of food.

#### Distribution

Ziprasidone has a mean apparent volume of distribution of 1.5 L/kg. It is greater than 99% bound to plasma proteins, binding primarily to albumin and  $\alpha$ -acid glycoprotein. The plasma protein binding of ziprasidone was not altered by warfarin or propranolol, two highly protein-bound drugs, nor did ziprasidone alter the binding of these drugs in human plasma. Thus, the potential for drug interactions with ziprasidone due to displacement is minimal. *in vitro*

## ***Metabolism and Elimination***

Ziprasidone is extensively metabolized after oral administration with only a small amount excreted in the urine (<1%) or feces (<4%) as unchanged drug. Ziprasidone is primarily cleared via three metabolic routes to yield four major circulating metabolites, benzisothiazole (BITP) sulphoxide, BITP-sulphone, ziprasidone sulphoxide, and S-methyl-dihydroziprasidone. Approximately 20% of the dose is excreted in the urine, with approximately 66% being eliminated in the feces. Unchanged ziprasidone represents about 44% of total drug-related material in serum. Studies using human liver subcellular fractions indicate that S-methyl-dihydroziprasidone is generated in two steps. These studies indicate that the reduction reaction is mediated primarily by chemical reduction by glutathione as well as by enzymatic reduction by aldehyde oxidase and the subsequent methylation is mediated by thiol methyltransferase. Studies using human liver microsomes and recombinant enzymes indicate that CYP3A4 is the major CYP contributing to the oxidative metabolism of ziprasidone. CYP1A2 may contribute to a much lesser extent. Based on abundance of excretory metabolites, less than one-third of ziprasidone metabolic clearance is mediated by cytochrome P450 catalyzed oxidation and approximately two-thirds via reduction. There are no known clinically relevant inhibitors or inducers of aldehyde oxidase. *In vitro*/*In vivo*

## **13 NONCLINICAL TOXICOLOGY**

### **13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility**

#### ***Carcinogenesis***

Lifetime carcinogenicity studies were conducted with ziprasidone in Long Evans rats and CD-1 mice. Ziprasidone was administered for 24 months in the diet at doses of 2, 6, or 12 mg/kg/day to rats, and 50, 100, or 200 mg/kg/day to mice (0.1 to 0.6 and 1 to 5 times the maximum recommended human dose [MRHD] of 200 mg/day on a mg/m basis, respectively). In the rat study, there was no evidence of an increased incidence of tumors compared to controls. In male mice, there was no increase in incidence of tumors relative to controls. In female mice, there were dose-related increases in the incidences of pituitary gland adenoma and carcinoma, and mammary gland adenocarcinoma at all doses tested (50 to 200 mg/kg/day or 1 to 5 times the MRHD on a mg/m basis). Proliferative changes in the pituitary and mammary glands of rodents have been observed following chronic administration of other antipsychotic agents and are considered to be prolactin-mediated. Increases in serum prolactin were observed in a 1-month dietary study in female, but not male, mice at 100 and 200 mg/kg/day (or 2.5 and 5 times the MRHD on a mg/m basis). Ziprasidone had no effect on serum prolactin in rats in a 5-week dietary study at the doses that were used in the carcinogenicity study. The relevance for human risk of the findings of prolactin-mediated endocrine tumors in rodents is unknown [see ].<sup>222</sup>**WARNINGS AND PRECAUTIONS ( ) 5.11**

#### ***Mutagenesis***

Ziprasidone was tested in the Ames bacterial mutation assay, the mammalian cell gene mutation mouse lymphoma assay, the chromosomal aberration assay in human lymphocytes, and the chromosomal aberration assay in mouse bone marrow. There was a reproducible mutagenic response in the Ames assay in one strain of in the absence of metabolic activation. Positive results were obtained in both the mammalian cell gene mutation assay and the chromosomal aberration assay in human lymphocytes. *in vitro*/*in vivo*. *S. typhimurium* *in vitro*/*in vivo*

#### ***Impairment of Fertility***

Ziprasidone was shown to increase time to copulation in Sprague-Dawley rats in two fertility and early embryonic development studies at doses of 10 to 160 mg/kg/day (0.5 to 8 times the MRHD of 200 mg/day on a mg/m basis). Fertility rate was reduced at 160 mg/kg/day (8 times the MRHD on a mg/m basis). There was no effect on fertility at 40 mg/kg/day (2 times the MRHD on a mg/m basis). The effect on fertility appeared to be in the female since fertility was not impaired when males given 160



mg/kg/day (8 times the MRHD on a mg/m basis) were mated with untreated females. In a 6-month study in male rats given 200 mg/kg/day (10 times the MRHD on a mg/m basis) there were no treatment-related findings observed in the testes. <sup>22222</sup>

## 14 CLINICAL STUDIES

### 14.1 Schizophrenia

The efficacy of oral ziprasidone in the treatment of schizophrenia was evaluated in 5 placebo-controlled studies, 4 short-term (4- and 6-week) trials and one maintenance trial. All trials were in adult inpatients, most of whom met DSM III-R criteria for schizophrenia. Each study included 2 to 3 fixed doses of ziprasidone as well as placebo. Four of the 5 trials were able to distinguish ziprasidone from placebo; one short-term study did not. Although a single fixed-dose haloperidol arm was included as a comparative treatment in one of the three short-term trials, this single study was inadequate to provide a reliable and valid comparison of ziprasidone and haloperidol.

Several instruments were used for assessing psychiatric signs and symptoms in these studies. The Brief Psychiatric Rating Scale (BPRS) and the Positive and Negative Syndrome Scale (PANSS) are both multi-item inventories of general psychopathology usually used to evaluate the effects of drug treatment in schizophrenia. The BPRS psychosis cluster (conceptual disorganization, hallucinatory behavior, suspiciousness, and unusual thought content) is considered a particularly useful subset for assessing actively psychotic schizophrenic patients. A second widely used assessment, the Clinical Global Impression (CGI), reflects the impression of a skilled observer, fully familiar with the manifestations of schizophrenia, about the overall clinical state of the patient. In addition, the Scale for Assessing Negative Symptoms (SANS) was employed for assessing negative symptoms in one trial.

#### **The results of the oral ziprasidone trials in schizophrenia follow:**

- In a 4-week, placebo-controlled trial (n=139) comparing 2 fixed doses of ziprasidone (20 and 60 mg twice daily) with placebo, only the 60 mg dose was superior to placebo on the BPRS total score and the CGI severity score. This higher dose group was not superior to placebo on the BPRS psychosis cluster or on the SANS.
- In a 6-week, placebo-controlled trial (n=302) comparing 2 fixed doses of ziprasidone (40 and 80 mg twice daily) with placebo, both dose groups were superior to placebo on the BPRS total score, the BPRS psychosis cluster, the CGI severity score and the PANSS total and negative subscale scores. Although 80 mg twice daily had a numerically greater effect than 40 mg twice daily, the difference was not statistically significant.
- In a 6-week, placebo-controlled trial (n=419) comparing 3 fixed doses of ziprasidone (20, 60, and 100 mg twice daily) with placebo, all three dose groups were superior to placebo on the PANSS total score, the BPRS total score, the BPRS psychosis cluster, and the CGI severity score. Only the 100 mg twice daily dose group was superior to placebo on the PANSS negative subscale score. There was no clear evidence for a dose-response relationship within the 20 mg twice daily to 100 mg twice daily dose range.
- In a 4-week, placebo-controlled trial (n=200) comparing 3 fixed doses of ziprasidone (5, 20, and 40 mg twice daily), none of the dose groups was statistically superior to placebo on any outcome of interest.
- A study was conducted in stable chronic or subchronic (CGI-S  $\leq$  5 at baseline) schizophrenic inpatients (n=294) who had been hospitalized for not less than two months. After a 3-day single-blind placebo run-in, subjects were randomized to one of 3 fixed doses of ziprasidone (20 mg, 40 mg, or 80 mg twice daily) or placebo and observed for relapse. Patients were observed for "impending psychotic relapse," defined as CGI-improvement score of  $\geq$  6 (much worse or very much worse) and/or scores  $\geq$  6 (moderately severe) on the hostility or uncooperativeness items of the PANSS on two consecutive days. Ziprasidone was significantly superior to placebo in time to

relapse, with no significant difference between the different dose groups. There were insufficient data to examine population subsets based on age and race. Examination of population subsets based on gender did not reveal any differential responsiveness.

## **16 HOW SUPPLIED/STORAGE AND HANDLING**

NDC:64725-2164-1 in a CONTAINER of 30 CAPSULES

## **17 PATIENT COUNSELING INFORMATION**

See FDA-Approved Patient Labeling ( ). 17.3

Please refer to the patient package insert. To assure safe and effective use of ziprasidone hydrochloride, the information and instructions provided in the patient information should be discussed with patients.

### **17.1 Administration with Food**

Patients should be instructed to take ziprasidone hydrochloride capsules with food for optimal absorption. The absorption of ziprasidone is increased up to two-fold in the presence of food [see and ]. **DRUG INTERACTIONS ( ) 7.8CLINICAL PHARMACOLOGY ( ) 12.3**

### **17.2 QTc Prolongation**

Patients should be advised to inform their health care providers of the following: History of QT prolongation; recent acute myocardial infarction; uncompensated heart failure; prescription of other drugs that have demonstrated QT prolongation; risk for significant electrolyte abnormalities; and history of cardiac arrhythmia [see and ]. **CONTRAINDICATIONS ( ) 4.1WARNINGS AND PRECAUTIONS ( ) 5.2**

Patients should be instructed to report the onset of any conditions that put them at risk for significant electrolyte disturbances, hypokalemia in particular, including but not limited to the initiation of diuretic therapy or prolonged diarrhea. In addition, patients should be instructed to report symptoms such as dizziness, palpitations, or syncope to the prescriber [see ]. **WARNINGS AND PRECAUTIONS ( ) 5.2**

### **17.3 FDA-Approved Patient Labeling**

## **PATIENT SUMMARY OF INFORMATION ABOUT**

### **Ziprasidone Hydrochloride Capsules**

#### **Information for patients taking ziprasidone hydrochloride capsules or their caregivers**

This summary contains important information about ziprasidone hydrochloride capsules. It is not meant to take the place of your doctor's instructions. Read this information carefully before you take ziprasidone hydrochloride capsules. Ask your doctor or pharmacist if you do not understand any of this information or if you want to know more about ziprasidone hydrochloride capsules.

#### **What are ziprasidone hydrochloride capsules?**

Ziprasidone hydrochloride capsules are a type of prescription medicine called a psychotropic, also known as an atypical antipsychotic. Ziprasidone hydrochloride capsules can be used to treat symptoms of schizophrenia.

#### **Who should take ziprasidone hydrochloride capsules?**

Only your doctor can know if ziprasidone hydrochloride capsules are right for you. Ziprasidone hydrochloride capsules may be prescribed for you if you have schizophrenia.

Symptoms of schizophrenia may include:

- hearing voices, seeing things, or sensing things that are not there (hallucinations)
- beliefs that are not true (delusions)
- unusual suspiciousness (paranoia)
- becoming withdrawn from family and friends

If you show a response to ziprasidone hydrochloride capsules, your symptoms may improve. If you continue to take ziprasidone hydrochloride capsules there is less chance of your symptoms returning. Do not stop taking the capsules even when you feel better without first discussing it with your doctor.

It is also important to remember that ziprasidone hydrochloride capsules should be taken with food.

### **What is the most important safety information I should know about ziprasidone hydrochloride capsules?**

**Ziprasidone hydrochloride capsules are not approved for the treatment of patients with dementia-related psychosis. Elderly patients with a diagnosis of psychosis related to dementia treated with antipsychotics are at an increased risk of death when compared to patients who are treated with placebo (a sugar pill).**

Ziprasidone hydrochloride capsules are an effective drug to treat the symptoms of schizophrenia. However, one potential side effect is that it may change the way the electrical current in your heart works more than some other drugs. The change is small and it is not known whether this will be harmful, but some other drugs that cause this kind of change have in rare cases caused dangerous heart rhythm abnormalities. Because of this, ziprasidone hydrochloride capsules should be used only after your doctor has considered this risk for ziprasidone hydrochloride capsules against the risks and benefits of other medications available for treating schizophrenia.

**Your risk of dangerous changes in heart rhythm can be increased if you are taking certain other medicines and if you already have certain abnormal heart conditions. Therefore, it is important to tell your doctor about any other medicines that you take, including non-prescription medicines, supplements, and herbal medicines. You must also tell your doctor about any heart problems you have or have had.**

### **Who should NOT take ziprasidone hydrochloride capsules?**

Elderly patients with a diagnosis of psychosis related to dementia. Ziprasidone hydrochloride capsules are not approved for the treatment of these patients.

Anything that can increase the chance of a heart rhythm abnormality should be avoided. Therefore, do not take ziprasidone hydrochloride capsules if:

- You have certain heart diseases, for example, long QT syndrome, a recent heart attack, severe heart failure, or certain irregularities of heart rhythm (discuss the specifics with your doctor)
- You are currently taking medications that should not be taken in combination with ziprasidone, for example, dofetilide, sotalol, quinidine, other Class Ia and III anti-arrhythmics, mesoridazine, thioridazine, chlorpromazine, droperidol, pimozide, sparfloxacin, gatifloxacin, moxifloxacin, halofantrine, mefloquine, pentamidine, arsenic trioxide, levomethadyl acetate, dolasetron mesylate, probucol or tacrolimus.

### **What to tell your doctor before you start ziprasidone hydrochloride capsules**

Only your doctor can decide if ziprasidone hydrochloride capsules are right for you. Before you start ziprasidone hydrochloride capsules, be sure to tell your doctor if you:

- have had any problem with the way your heart beats or any heart related illness or disease
- any family history of heart disease, including recent heart attack
- have had any problem with fainting or dizziness

- are taking or have recently taken any prescription medicines
- are taking any over-the-counter medicines you can buy without a prescription, including natural/herbal remedies
- have had any problems with your liver
- are pregnant, might be pregnant, or plan to get pregnant
- are breast feeding
- are allergic to any medicines
- have ever had an allergic reaction to ziprasidone or any of the other ingredients of ziprasidone hydrochloride capsules. Ask your doctor or pharmacist for a list of these ingredients
- have low levels of potassium or magnesium in your blood

Your doctor may want you to get additional laboratory tests to see if ziprasidone hydrochloride capsules are an appropriate treatment for you.

### **Ziprasidone hydrochloride capsules and other medicines**

There are some medications that may be unsafe to use when taking ziprasidone hydrochloride capsules, and there are some medicines that can affect how well ziprasidone hydrochloride capsules works. While you are on ziprasidone hydrochloride capsules, check with your doctor before starting any new prescription or over-the-counter medications, including natural/herbal remedies.

### **How to take ziprasidone hydrochloride capsules**

- Take ziprasidone hydrochloride capsules only as directed by your doctor.
- Swallow the capsules whole.
- Take ziprasidone hydrochloride capsules with food.
- It is best to take ziprasidone hydrochloride capsules at the same time each day.
- Ziprasidone hydrochloride capsules may take a few weeks to work. It is important to be patient.
- Do not change your dose or stop taking your medicine without your doctor's approval.
- Remember to keep taking your capsules, even when you feel better.

### **Possible side effects**

Because these problems could mean you're having a heart rhythm abnormality, contact your doctor if you: **IMMEDIATELY**

- Faint or lose consciousness
- Feel a change in the way that your heart beats (palpitations)

Common side effects of ziprasidone hydrochloride capsules include the following and should also be discussed with your doctor if they occur:

- Feeling unusually tired or sleepy
- Nausea or upset stomach
- Constipation
- Dizziness
- Restlessness
- Abnormal muscle movements, including tremor, shuffling, and uncontrolled involuntary movements
- Diarrhea
- Rash
- Increased cough / runny nose

If you develop any side effects that concern you, talk with your doctor. It is particularly important to tell your doctor if you have diarrhea, vomiting, or another illness that can cause you to lose fluids. Your doctor may want to check your blood to make sure that you have the right amount of important salts after such illnesses.

For a list of all side effects that have been reported, ask your doctor or pharmacist for the ziprasidone hydrochloride capsules Professional Package Insert.

### **What to do for an overdose**

In case of an overdose, call your doctor or poison control center right away or go to the nearest emergency room.

### **Other important safety information**

A serious condition called neuroleptic malignant syndrome (NMS) can occur with all antipsychotic medications including ziprasidone hydrochloride capsules. Signs of NMS include very high fever, rigid muscles, shaking, confusion, sweating, or increased heart rate and blood pressure. NMS is a rare but serious side effect that could be fatal. Therefore, tell your doctor if you experience any of these signs.

Adverse reactions related to high blood sugar (hyperglycemia), sometimes serious, have been reported in patients treated with atypical antipsychotics. There have been few reports of hyperglycemia or diabetes in patients treated with ziprasidone hydrochloride capsules, and it is not known if ziprasidone hydrochloride capsules are associated with these reactions. Patients treated with an atypical antipsychotic should be monitored for symptoms of hyperglycemia.

Dizziness caused by a drop in your blood pressure may occur with ziprasidone hydrochloride capsules, especially when you first start taking this medication or when the dose is increased. If this happens, be careful not to stand up too quickly, and talk to your doctor about the problem.

Before taking ziprasidone hydrochloride capsules, tell your doctor if you are pregnant or plan on becoming pregnant. It is advised that you don't breast feed an infant if you are taking ziprasidone hydrochloride capsules.

Because ziprasidone hydrochloride capsules can cause sleepiness, be careful when operating machinery or driving a motor vehicle.

Since medications of the same drug class as ziprasidone hydrochloride capsules may interfere with the ability of the body to adjust to heat, it is best to avoid situations involving high temperature or humidity.

It is best to avoid consuming alcoholic beverages while taking ziprasidone hydrochloride capsules. Call your doctor if you take more than the amount of ziprasidone hydrochloride capsules prescribed by your doctor. ***immediately***

Ziprasidone hydrochloride capsules have not been shown to be safe or effective in the treatment of children and teenagers under the age of 18 years old.

**Keep ziprasidone hydrochloride capsules and all medicines out of the reach of children.**

### **How to store ziprasidone hydrochloride capsules**

Store ziprasidone hydrochloride capsules at room temperature 20° to 25°C (68° to 77°F).

### **For more information about ziprasidone hydrochloride capsules**

This sheet is only a summary. Ziprasidone hydrochloride capsules are a prescription medicine and only your doctor can decide if it is right for you. If you have any questions or want more information about ziprasidone hydrochloride capsules, talk with your doctor or pharmacist.

Maalox is a registered trademark of Novartis. ®

Manufactured in India by Sandoz Private Limited for

Sandoz Inc., Princeton, NJ 08540

Rev. August 2013

**ZIPRASIDONE HCL 20MG CAPSULE(30CTCD)**

**RX ONLY**

**0781-2164-60**

**LOT: XXXX**

**EXP: XX-XX-XX**

**MFR: SANDOZ**

**REPACKAGED BY: T.Y.A. PHARMACEUTICALS**

**2930 CRESCENT DR.**

**TALLAHASSEE, FL 32301**

**(850) 385-0228**



**DOSE: SEE PACKAGE INSERT**

**ZIPRASIDONE HCL 80MG CAPSULE(30CTCD)**

**RX ONLY**

**0781-2168-60**

**LOT: XXXX**

**EXP: XX-XX-XX**

**MFR: SANDOZ**

**REPACKAGED BY: T.Y.A. PHARMACEUTICALS**

**2930 CRESCENT DR.**

**TALLAHASSEE, FL 32301**

**(850) 385-0228**



**DOSE: SEE PACKAGE INSERT**

**ZIPRASIDONE HYDROCHLORIDE**

ziprasidone hydrochloride capsule

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:64725-2164(NDC:0781-2164)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZIPRASIDONE HYDROCHLORIDE (UNII: 216X0810RU) (ZIPRASIDONE - UNII:6UKA5VEJ6X)	ZIPRASIDONE	20 mg

### Inactive Ingredients

Ingredient Name	Strength
CALCIUM SILICATE (UNII: S4255P4G5M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6B30)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
GELATIN (UNII: 2G86QN327L)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
AMMONIA (UNII: 5138Q19F1X)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	

### Product Characteristics

<b>Color</b>	YELLOW (opaque yellow cap) , BLUE (opaque blue body)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	14mm
<b>Flavor</b>		<b>Imprint Code</b>	SZ;656
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64725-2164-1	30 in 1 CONTAINER		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077562	02/13/2013	

**ZIPRASIDONE HYDROCHLORIDE**

ziprasidone hydrochloride capsule

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:64725-2168(NDC:0781-2168)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZIPRASIDONE HYDROCHLORIDE (UNII: 216X081ORU) (ZIPRASIDONE - UNII:6UKA5VEJ6X)	ZIPRASIDONE	80 mg

### Inactive Ingredients

Ingredient Name	Strength
CALCIUM SILICATE (UNII: S4255P4G5M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
GELATIN (UNII: 2G86QN327L)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
AMMONIA (UNII: 5138Q19F1X)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	

### Product Characteristics

<b>Color</b>	YELLOW (opaque yellow cap) , BLUE (opaque blue body)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	SZ;659
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64725-2168-1	30 in 1 CONTAINER		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077562	02/13/2013	



**Labeler** - TYA Pharmaceuticals (938389038)

**Registrant** - TYA Pharmaceuticals (938389038)

**Establishment**

Name	Address	ID/FEI	Business Operations
TYA Pharmaceuticals		938389038	RELABEL(64725-2164, 64725-2168) , REPACK(64725-2164)

Revised: 8/2013

TYA Pharmaceuticals