

# Summary of Product Characteristics

## 1 NAME OF THE MEDICINAL PRODUCT

GEODON 10 mg/ml oral suspension.

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ml contains ziprasidone hydrochloride monohydrate equivalent to 10 mg of ziprasidone.

Excipient(s) with known effects:

Each ml contains 1.36 mg methyl parahydroxybenzoate.

Each ml contains 0.17 mg propyl parahydroxybenzoate.

Each ml contains 4.65 mg sodium.

For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Oral suspension

White to slightly yellow opaque cherry flavoured aqueous suspension.

## 4 CLINICAL PARTICULARS

### 4.1 Therapeutic Indications

Ziprasidone is indicated for the treatment of schizophrenia in adults.

Ziprasidone is indicated for the treatment of manic or mixed episodes of moderate severity in bipolar disorder in adults, children and adolescents aged 10-17 years (prevention of episodes of bipolar disorder has not been established - see section 5.1).

### 4.2 Posology and method of administration

Posology

#### *Adults*

The recommended dose, in acute treatment of schizophrenia and bipolar mania, is 40 mg twice daily taken with food. Daily dosage may subsequently be adjusted on the basis of individual clinical status up to a maximum of 80 mg twice daily. If indicated, the maximum recommended dose may be reached as early as day 3 of treatment.

It is of particular importance not to exceed the maximum dose as the safety profile above 160mg/day has not been confirmed and ziprasidone is associated with dose-related prolongation of the QT interval (see sections 4.3 and 4.4).

In maintenance treatment of schizophrenia patients, ziprasidone should be administered at the lowest effective dose; in many cases, a dose of 20 mg twice daily may be sufficient.

Measure the dose as closely as possible using the oral syringe provided. Ziprasidone oral suspension should be administered directly into the mouth. Although the suspension should be taken with food, it should not be diluted or mixed into any food or beverages prior to administration.

#### *Elderly*

A lower starting dose is not routinely indicated but should be considered for those 65 and over when clinical factors warrant.

*Patients with renal impairment*

No dose adjustment is required in patients with impaired renal function (see section 5.2).

*Patients with hepatic impairment*

In patients with hepatic insufficiency, lower doses should be considered. (See sections 4.4 and 5.2).

*Paediatric Population**Bipolar Mania*

The recommended dose, in acute treatment of bipolar mania, in paediatric patients (age 10 to 17 years) is a single dose of 20 mg on day 1, with food. Ziprasidone should subsequently be administered with food in two daily divided doses, and should be titrated over 1-2 weeks to a target range of 120-160 mg/day for patients weighing  $\geq 45$  kg, or to a target range of 60-80 mg/day for patients weighing  $< 45$  kg. Subsequent dosing should be adjusted on the basis of individual clinical status within the range of 80-160 mg/day for patients weighing  $\geq 45$  kg, or 40-80 mg/day for patients weighing  $< 45$  kg. Asymmetric dosing, with morning doses 20 mg or 40 mg less than evening doses, was permitted in the clinical trial (see sections 4.4, 5.1 and 5.2).

It is of particular importance not to exceed the weight-based maximum dose as the safety profile above the maximum dose (160 mg/day for children  $\geq 45$  kg and 80 mg/day for children  $< 45$  kg) has not been confirmed and ziprasidone is associated with dose-related prolongation of the QT interval (see sections 4.3 and 4.4)

*Schizophrenia*

The safety and efficacy of ziprasidone in paediatric patients with schizophrenia have not been established (see sections 4.4).

**4.3 Contraindications**

Hypersensitivity to the active substance or any of the excipients listed in section 6.1. Known QT-interval prolongation. Congenital long QT syndrome. Recent acute myocardial infarction. Uncompensated heart failure. Arrhythmias treated with class IA and III antiarrhythmic medicinal products.

Concomitant treatment with medicinal products that prolong the QT interval, such as Class IA and III antiarrhythmics, arsenic trioxide, halofantrine, levomethadyl acetate, mesoridazine, thioridazine, pimozide, sparfloxacin, gatifloxacin, moxifloxacin, dolasetron mesilate, mefloquine, sertindole or cisapride (see sections 4.4 and 4.5)

**4.4 Special warnings and precautions for use**

A medical history, including assessment of family history, and physical examination should be undertaken to identify patients for whom ziprasidone treatment is not recommended (see section 4.3).

*QT interval*

Ziprasidone causes a mild to moderate dose-related prolongation of the QT-interval (see section 4.8 and 5.1).

Ziprasidone should not be given together with medicinal products that are known to prolong the QT-interval (see sections 4.3 and 4.5). Caution is advised in patients with significant bradycardia. Electrolyte disturbances such as hypokalaemia and hypomagnesaemia increase the risk for malignant arrhythmias and should be corrected before treatment with ziprasidone is started. If patients with stable cardiac disease are treated, an ECG review should be considered before treatment is started.

If cardiac symptoms, such as palpitations, vertigo, syncope or seizures occur, then the possibility of a malignant cardiac arrhythmia should be considered and a cardiac evaluation including an ECG should be performed. If the QTc interval is  $> 500$  msec, then it is recommended that the treatment should be stopped (see section 4.3).

There have been rare post-marketing reports of torsade de pointes in patients with multiple confounding risk factors taking ziprasidone.

*Paediatric Population*

Safety and efficacy of ziprasidone in the treatment of schizophrenia in children and adolescents has not been evaluated.

*Neuroleptic malignant syndrome (NMS)*

NMS is a rare but potentially fatal complex that has been reported in association with antipsychotic medicinal products, including ziprasidone. The management of NMS should include immediate discontinuation of all antipsychotic medicinal products.

*Severe Cutaneous Adverse Reactions*

Drug reaction with eosinophilia and systemic symptoms (DRESS) has been reported with ziprasidone exposure.

DRESS consists of a combination of three or more of the following:

cutaneous reaction (such as rash or exfoliative dermatitis), eosinophilia, fever, lymphadenopathy and one or more systemic complications such as hepatitis, nephritis, pneumonitis, myocarditis, and pericarditis.

Other severe cutaneous adverse reactions, such as Stevens-Johnson syndrome, have been reported with ziprasidone exposure.

Severe cutaneous adverse reactions are sometimes fatal. Discontinue ziprasidone if severe cutaneous reactions occur.

*Tardive dyskinesia*

There is a potential for ziprasidone to cause tardive dyskinesia and other tardive extrapyramidal syndromes after long-term treatment. Patients with bipolar disorder are known to be particularly vulnerable to this category of symptoms.

This is more frequent with increased duration of treatment and increasing age. If signs and symptoms of tardive dyskinesia appear, dose reduction or discontinuation of ziprasidone should be considered.

*Seizures*

Caution is recommended when treating patients with a history of seizures.

*Hepatic Impairment*

There is a lack of experience in patients with severe hepatic insufficiency and ziprasidone should be used with caution in this group (see sections 4.2 and 5.2).

*Increased risk of cerebrovascular accidents in the dementia population*

An approximately 3-fold increased risk of cerebrovascular adverse events has been seen in randomised placebo-controlled clinical trials in the dementia population with some atypical antipsychotics. The mechanism for this increased risk is not known. An increased risk cannot be excluded for other antipsychotics or other patient populations. Geodon should be used with caution in patients with risk factors for stroke.

Geodon 10 mg/ml oral suspension contains methyl parahydroxybenzoate and propyl parahydroxybenzoate which may cause allergic reactions (possibly delayed).

Geodon 10 mg/ml oral suspension contains 4.65 mg sodium per ml dosed and this should be taken into account in patients on a controlled sodium diet. The usual dose is between 4 and 8 ml twice a day so the total daily sodium load is between 37.2 and 74.4 mg.

*Increased Mortality in Elderly people with Dementia* Data from two large observational studies showed that elderly people with dementia who are treated with antipsychotics are at a small increased risk of death compared with those who are not treated. There are insufficient data to give a firm estimate of the precise magnitude of the risk and the cause of the increased risk is not known.

Geodon is not licensed for the treatment of dementia-related behavioural disturbances.

*Venous Thromboembolism*

Cases of venous thromboembolism (VTE) have been reported with antipsychotic drugs. Since patients treated with antipsychotics often present with acquired risk factors for VTE, all possible risk factors for VTE should be identified before and during treatment with ziprasidone and preventive measures undertaken.

*Priapism*

Cases of priapism have been reported with antipsychotic use, including ziprasidone. This adverse reaction, as with other psychotropic drugs, did not appear to be dose-dependent and did not correlate with the duration of treatment.

*Hyperprolactinemia*

As with other drugs that antagonize dopamine D2 receptors, ziprasidone may elevate prolactin levels. Disturbances such as galactorrhea, amenorrhea, gynecomastia, and impotence have been reported with prolactin-elevating compounds. Long-standing hyperprolactinemia when associated with hypogonadism may lead to decreased bone density.

**4.5 Interaction with other medicinal products and other forms of interaction**

Pharmacokinetic and pharmacodynamic studies between ziprasidone and other medicinal products that prolong the QT interval have not been performed. An additive effect of ziprasidone and these medicinal products cannot be excluded, therefore ziprasidone should not be given with medicinal products that prolong the QT interval, such as Class IA and III antiarrhythmics, arsenic trioxide, halofantrine, levomethadyl acetate, mesoridazine, thioridazine, pimozide, sparfloxacin, gatifloxacin, moxifloxacin, dolasetron mesilate, mefloquine, sertindole or cisapride (see section 4.3)

No studies on the interaction of ziprasidone with other medicinal products have been performed in children

*CNS medicinal products /alcohol*

Given the primary effects of ziprasidone, caution should be used when it is taken in combination with other centrally acting medicinal products and alcohol.

*Effect of ziprasidone on other medicinal products*

An *in vivo* study with dextromethorphan showed no marked inhibition of CYP2D6 at plasma concentrations 50% lower than those obtained after 40 mg ziprasidone twice daily. *In vitro* data indicated that ziprasidone may be a modest inhibitor of CYP2D6 and CYP3A4. However, it is unlikely that ziprasidone will affect the pharmacokinetics of medicinal products metabolised by these cytochrome P450 isoforms to a clinically relevant extent.

Oral contraceptives – Ziprasidone administration resulted in no significant change to the pharmacokinetics of oestrogen (ethinyl oestradiol, a CYP3A4 substrate) or progesterone components.

Lithium - Co-administration of ziprasidone had no effect on the pharmacokinetics of lithium.

As ziprasidone and lithium are associated with cardiac conduction changes, the combination may pose a risk for pharmacodynamic interactions including arrhythmias, however, in controlled clinical trials, the combination of ziprasidone plus lithium has not demonstrated an increased clinical risk, compared to lithium alone.

There are limited data on co-medication with the mood stabilisers carbamazepine

A pharmacokinetic interaction of ziprasidone with valproate is unlikely due to the lack of common metabolic pathways for the two drugs. In a study in patients, the co-administration of ziprasidone and valproate showed that the mean concentrations of valproate were within the therapeutic range as compared to valproate administered with placebo.

*Effects of other medicinal products on ziprasidone*

The CYP3A4 inhibitor ketoconazole (400 mg/day) increased the serum concentrations of ziprasidone by <40%. The serum concentrations of S-methyl-dihydroziprasidone and ziprasidone sulphoxide, at the expected T<sub>max</sub> of ziprasidone, were increased by 55% and 8% respectively. No additional QTc prolongation was observed. Changes in pharmacokinetics due to co-administration of potent CYP3A4 inhibitors are unlikely to be of clinical importance, therefore no dosage adjustment is required. *In vitro* data indicates that ziprasidone is a P-glycoprotein (p-gp) substrate. The *in vivo* relevance is unknown, however co-administration with known p-gp inhibitors such as verapamil, macrolide antibiotics, quinidine, itraconazole and ritonavir could cause increased plasma concentrations of ziprasidone. Co-administration with p-gp inducers such as rifampin and St John's Wort could cause decreased concentrations of ziprasidone. This should be taken into account if co-administration is considered.

Carbamazepine therapy, 200 mg b.i.d for 21 days, resulted in a decrease of approximately 35% in the exposure to

ziprasidone.

Antacid - multiple doses of aluminium and magnesium containing antacid or cimetidine have no clinically significant effect on the pharmacokinetics of ziprasidone under fed conditions.

#### *Serotonergic medicinal products*

In isolated cases, there have been reports of serotonin syndrome temporally associated with the therapeutic use of ziprasidone in combination with other serotonergic medicinal products such as SSRIs (see section 4.8). The features of serotonin syndrome can include confusion, agitation, fever, sweating, ataxia, hyperreflexia, myoclonus and diarrhoea.

#### *Protein binding*

Ziprasidone extensively binds to plasma proteins. The in vitro 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 unlikely.

## **4.6 Fertility, pregnancy and lactation**

Reproductive toxicity studies have shown undesirable effects on the reproductive process, at doses associated with maternal toxicity and/or sedation. There was no evidence of teratogenicity (see section 5.3).

#### *Pregnancy*

No studies have been conducted in pregnant women. Women of child bearing potential receiving ziprasidone should therefore be advised to use an appropriate method of contraception. As human experience is limited, administration of ziprasidone is not recommended during pregnancy unless the expected benefit to the mother outweighs the potential risk to the foetus.

#### *Antipsychotic class labeling*

Neonates exposed to antipsychotics (including ziprasidone) during the third trimester of pregnancy are at risk of adverse reactions including extrapyramidal and/or withdrawal symptoms that may vary in severity and duration following delivery. There have been reports of agitation, hypertonia, hypotonia, tremor, somnolence, respiratory distress, or feeding disorder. Consequently, newborns should be monitored carefully. Geodon should not be used during pregnancy unless clearly necessary. If discontinuation during pregnancy is necessary, it should not be done abruptly.

#### *Breast-feeding*

It is not known whether ziprasidone is excreted in breast milk. Patients should not breast feed an infant if they are taking ziprasidone. If treatment is necessary, breast-feeding should be discontinued.

## **4.7 Effects on ability to drive and use machines**

Ziprasidone may cause somnolence and may influence on the ability to drive and use machines. Patients likely to drive or operate machines should be cautioned appropriately.

## **4.8 Undesirable effects**

Oral ziprasidone has been administered in clinical trials (see section 5.1) to approximately 6500 adult subjects. The most common adverse reactions in schizophrenia clinical trials were sedation and akathisia. In bipolar mania clinical trials, the most common adverse reactions were sedation, akathisia, extrapyramidal disorder and dizziness.

The table below contains adverse events based on combined short term (4-6 week), fixed dose, schizophrenia studies and short term (3 week), flexible dose, bipolar mania studies, with a probable or possible relationship to treatment with ziprasidone and which occur at an incidence greater than placebo. Additional reactions reported from the post-marketing experience are included as Frequency 'Not known' in italics in the list below.

All adverse reactions are listed by class and frequency: *very common* ( $\geq 1/10$ ); *common* ( $\geq 1/100$  to  $< 1/10$ ); *uncommon* ( $\geq 1/1,000$  to  $< 1/100$ ); *rare* ( $\geq 1/10,000$  to  $< 1/1,000$ ); *very rare* ( $< 1/10,000$ ), *not known* (*cannot be estimated from the available data*).

The adverse reactions listed below may also be associated with the underlying disease and/or concomitant medications.

| <b>System Organ Class<br/>Frequency</b>                | <b>Adverse drug reactions</b>  |
|--|--|
| <b>Infections and Infestations</b>                     |  |
| Rare   | Rhinitis   |
| <b>Metabolism and nutrition disorders</b>              |  |
| Uncommon   | Increased appetite   |
| Rare   | Hypocalcaemia  |
| <b>Psychiatric disorders</b>                           |  |
| Common   | Restlessness   |
| Uncommon   | Agitation, anxiety, throat tightness, nightmare  |
| Rare   | Panic attack, depressive symptom, bradyphrenia, flat affect, anorgasmia  |
| Not known  | Insomnia; mania/hypomania  |
| <b>Nervous system disorders</b>                        |  |
| Common   | Dystonia, akathisia, extrapyramidal disorder, parkinsonism (including cogwheel rigidity, bradykinesia, hypokinesia), tremor, dizziness, sedation, somnolence, headache                           |
| Uncommon   | Generalised tonic clonic seizures, tardive dyskinesia, dyskinesia, drooling, ataxia, dysarthria, oculogyric crisis, disturbance in attention, hypersomnia, hypoaesthesia, paraesthesia, lethargy |
| Rare   | Torticollis, paresis, akinesia, hypertonia, restless legs syndrome   |
| Not known  | Neuroleptic malignant syndrome; serotonin syndrome (see section 4.5); facial droop   |
| <b>Blood and lymphatic system disorders</b>            |  |
| Rare   | Lymphopenia, eosinophil count increased  |
| <b>Cardiac disorders</b>                               |  |
| Uncommon   | Palpitations, tachycardia  |
| Rare   | Electrocardiogram QT corrected interval prolonged  |
| Not known  | Torsade de pointes (see section 4.4)   |
| <b>Eye disorders</b>                                   |  |
| Common   | Vision blurred   |
| Uncommon   | Photophobia  |
| Rare   | Amblyopia, visual disturbance, eye pruritis, dry eyes  |
| <b>Ear and labyrinth disorders</b>                     |  |
| Uncommon   | Vertigo, tinnitus  |
| Rare   | Ear pain   |
| <b>Vascular disorders</b>                              |  |
| Uncommon   | Hypertensive crisis, hypertension, orthostatic hypotension, hypotension  |
| Rare   | Systolic hypertension, diastolic hypertension, labile blood pressure   |
| Not known  | Syncope, venous thromboembolism  |
| <b>Respiratory, thoracic and mediastinal disorders</b> |  |
| Uncommon   | Dyspnoea, sore throat  |
| Rare   | Hiccups  |
| <b>Gastrointestinal disorders</b>                      |  |
| Common   | Nausea, vomiting, constipation, dyspepsia, dry mouth, salivary hypersecretion  |
| Uncommon   | Diarrhoea, dysphagia, gastritis, gastrointestinal discomfort, swollen tongue, tongue thick, flatulence   |
| Rare   | Gastro-oesophageal reflux, loose stools  |
| <b>Skin and subcutaneous tissue disorders</b>          |  |
| Uncommon   | Urticaria, rash, rash maculo-papular, acne   |
| Rare   | Psoriasis, dermatitis allergic, alopecia, swelling face, erythema, rash papular, skin irritation   |

|   |   |
|---|---|
| Not known   | Hypersensitivity, angioedema, drug reaction with eosinophilia and systemic symptoms (DRESS) |
| <b>Musculoskeletal and connective tissue disorders</b>      |   |
| Common  | Musculoskeletal rigidity  |
| Uncommon  | Musculoskeletal discomfort, muscle cramp, pain in extremity, joint stiffness                |
| Rare  | Trismus   |
| <b>Renal and urinary disorders</b>                          |   |
| Rare  | Urinary incontinence, dysuria   |
| Not known   | Enuresis  |
| <b>Reproductive system and breast disorders</b>             |   |
| Rare  | Erectile dysfunction, erection increased, galactorrhoea, gynaecomastia                      |
| Not known   | Priapism  |
| <b>Immune system disorders</b>                              |   |
| Not known   | Anaphylactic reaction   |
| <b>Hepatobiliary disorders</b>                              |   |
| Uncommon  | Hepatic enzyme increased  |
| Rare  | Liver function test abnormal  |
| <b>General disorders and administration site conditions</b> |   |
| Common  | Asthenia, fatigue   |
| Uncommon  | Chest discomfort, gait abnormal, pain, thirst   |
| Rare  | Pyrexia, feeling hot  |
| <b>Investigations</b>                                       |   |
| Rare  | Blood lactate dehydrogenase increased   |
| <b>Pregnancy, puerperium and perinatal conditions</b>       |   |
| Not known   | Drug withdrawal syndrome neonatal (see section 4.6)   |

In short-term and long-term ziprasidone schizophrenia and bipolar mania clinical trials, the incidence of tonic clonic seizures and hypotension was uncommon, occurring in less than 1% of ziprasidone treated patients.

Ziprasidone causes a mild to moderate dose-related prolongation of the QT interval (see section 5.1). In schizophrenia clinical trials, an increase of 30 to 60 msec was seen in 12.3% (976/7941) of ECG tracings from ziprasidone-treated and 7.5% (73/975) of ECG tracings from placebo-treated patients. A prolongation of >60 msec was seen in 1.6% (128/7941) and 1.2% (12/975) of tracings from ziprasidone and placebo-treated patients, respectively. The incidence of QTc interval prolongation above 500 msec was 3 in a total of 3266 (0.1%) in ziprasidone treated patients and 1 in a total of 538 (0.2%) in placebo treated patients. Comparable findings were observed in bipolar mania clinical trials.

In long term maintenance treatment in schizophrenia clinical trials, prolactin levels in patients treated with ziprasidone were sometimes elevated, but, in most patients, returned to normal ranges without cessation of treatment. In addition, potential clinical manifestations (e.g. gynaecomastia and breast enlargement) were rare.

Paediatric and Adolescent population with Bipolar mania and Adolescents with Schizophrenia.

In a placebo-controlled bipolar disorder trial (ages 10–17 years), the most frequent adverse reactions (reported with a frequency >10%) were sedation, somnolence, headache, fatigue, nausea, and dizziness. In a placebo-controlled schizophrenia trial (ages 13-17 years), the most frequent adverse reactions (reported with a frequency >10%) were somnolence and extrapyramidal disorder. The frequency, type and severity of adverse reactions in these subjects were generally similar to those in adults with bipolar disorder or schizophrenia who are treated with ziprasidone.

Ziprasidone was associated with a similar mild to moderate dose-related prolongation of the QT interval in the paediatric bipolar and schizophrenia clinical trials to those seen in the adult population. Tonic clonic seizures and hypotension were not reported in the placebo-controlled paediatric bipolar clinical trials.

#### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any

suspected adverse reactions via HPRA Pharmacovigilance, Earlsfort Terrace, IRL - Dublin 2; Tel: +353 1 6764971; Fax: +353 1 6762517. Website: [www.hpra.ie](http://www.hpra.ie); E-mail: [medsafety@hpra.ie](mailto:medsafety@hpra.ie)

## 4.9 Overdose

Experience with ziprasidone in overdose is limited. The largest confirmed single ingestion of ziprasidone is 12,800 mg. In this case, extrapyramidal symptoms and a QTc interval of 446 msec (with no cardiac sequelae) were reported. In general, the most commonly reported symptoms following overdose are, extrapyramidal symptoms, somnolence, tremor and anxiety.

The possibility of obtundation, seizures 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. There is no specific antidote to ziprasidone.

## 5 PHARMACOLOGICAL PROPERTIES

### 5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Antipsychotic, indole derivatives, ATC code NO5A E04.

Ziprasidone has a high affinity for dopamine type 2 (D<sub>2</sub>) receptors and substantially higher affinity for serotonin type 2<sub>A</sub> (5HT<sub>2A</sub>) receptors. Receptor blockade, 12 hours after a single dose of 40 mg, was greater than 80% for serotonin type 2<sub>A</sub> and greater than 50% for D<sub>2</sub> using positron emission tomography (PET). Ziprasidone also interacts with serotonin 5HT<sub>2C</sub>, 5HT<sub>1D</sub> and 5HT<sub>1A</sub> receptors where its affinities for these sites are equal to or greater than its affinity for the D<sub>2</sub> receptor. Ziprasidone has moderate affinity for neuronal serotonin and norepinephrine transporters.

Ziprasidone demonstrates moderate affinity for histamine H(1)- and alpha(1)-receptors. Ziprasidone demonstrates negligible affinity for muscarinic M(1)-receptors.

Ziprasidone has been shown to be an antagonist at both serotonin type 2<sub>A</sub> (5HT<sub>2A</sub>) and dopamine type 2 (D<sub>2</sub>) receptors. It is proposed that the therapeutic activity is mediated, in part, through this combination of antagonist activities. Ziprasidone is also a potent antagonist at 5HT<sub>2C</sub> and 5HT<sub>1D</sub> receptors, a potent agonist at the 5HT<sub>1A</sub> receptor and inhibits neuronal reuptake of norepinephrine and serotonin.

#### *Further information on clinical trials*

##### *Schizophrenia*

In a 52 week study, ziprasidone was effective in maintaining the clinical improvement during continuation therapy in patients who showed an initial treatment response: there was no clear evidence for a dose-response relationship amongst the ziprasidone groups. In this study, which included patients with both positive and negative symptoms, ziprasidone's efficacy was demonstrated in both positive and negative symptoms.

The incidence of body weight gain, reported as an adverse event in short term (4-6 week) schizophrenia studies was low and identical in ziprasidone-treated and placebo-treated patients (both 0.4%). In a one-year placebo-controlled study a median weight loss of 1-3 kg was observed in ziprasidone-treated patients compared to a 3 kg median loss in placebo-treated patients.

In a double-blind comparative schizophrenia study, metabolic parameters including weight and fasting levels of insulin, total cholesterol and triglycerides and an insulin resistance (IR) index were measured. In patients receiving ziprasidone no significant changes from baseline were observed in any of these metabolic parameters.

#### *Results of a large post-marketing safety study*

A randomised post-approval study of 18,239 schizophrenic patients with observational follow-up for 1 year was conducted to determine whether ziprasidone's effect on the QTc interval is associated with an increased risk of non-suicide mortality. This study, which was conducted in naturalistic clinical practice settings, showed no difference in the rate of over-all non-suicide mortality between ziprasidone and olanzapine treatments (primary end-point). The study also showed no difference in secondary end-points of all-cause mortality, mortality due to suicide, mortality due to sudden death, however, a non-significant numerically higher incidence of cardiovascular mortality was observed in the ziprasidone group. A statistically significantly higher incidence of all-cause hospitalisation, mainly due to differences in the number of psychiatric hospitalisations, was also observed in the ziprasidone group.

### *Bipolar mania*

The efficacy of ziprasidone in adults with mania was established in two placebo controlled, double blind, 3 week studies which compared ziprasidone with placebo and one double blind, 12 week study which compared ziprasidone to haloperidol and placebo. These studies included approximately 850 patients meeting DSM-IV criteria for bipolar I disorder with an acute manic or mixed episode, with or without psychotic features. The baseline presence of psychotic features in the studies was 49.7%, 34.7% or 34.9%. Efficacy was assessed using the Mania Rating Scale (MRS). The Clinical Global Impression-Severity (CGI-S) scale was either a co-primary or key secondary efficacy variable in these studies. Ziprasidone treatment (40-80 mg BID, mean daily dose 120 mg) resulted in statistically significantly greater improvement in both MRS and CGI-S scores at Last Visit (3 weeks) compared with placebo. In the 12 week study, haloperidol treatment (mean daily dose 16 mg) produced significantly greater reductions in MRS scores compared with ziprasidone (mean daily dose 121 mg). Ziprasidone demonstrated comparable efficacy to haloperidol in terms of the proportion of patients maintaining a response to treatment from week 3 to week 12.

The efficacy of ziprasidone in the treatment of Bipolar I Disorder in paediatric patients (10 to 17 years of age) was evaluated in one four-week placebo-controlled trial (n=237) of inpatients or outpatients who met DSM-IV criteria for Bipolar I Disorder manic or mixed episodes with or without psychotic features and had a Y-MRS score  $\geq 17$  at baseline. This double-blind, placebo-controlled trial compared flexibly-dosed oral ziprasidone (80-160 mg/day (40-80 mg BID) in two divided doses for patients weighing  $\geq 45$  kg; 40-80 mg/day (20-40 mg BID) for patients weighing  $< 45$  kg) to placebo. Ziprasidone was administered as a single dose of 20 mg on the first day, then titrated over 1-2 weeks, in two daily doses, to a target range of 120-160 mg/day for patients weighing  $\geq 45$  kg, or 60-80 mg/day for patients weighing  $< 45$  kg. Asymmetric dosing, with morning doses 20 mg or 40 mg less than evening doses, was permitted. Ziprasidone was superior to placebo in change from baseline to week 4 on the Y-MRS total score. In this clinical trial, the mean daily doses administered were 119 mg and 69 mg in the patients weighing  $\geq 45$  kg and  $< 45$  kg, respectively.

Ziprasidone has been evaluated for safety in 267 paediatric patients (age 10 to 17 years) who participated in multiple-dose, clinical trials in bipolar mania; a total of 82 paediatric patients with Bipolar I Disorder were dosed with oral ziprasidone for at least 180 days.

In a 4-week trial in paediatric patients (10-17 years) with bipolar mania, there were no differences between ziprasidone and placebo patients in the mean change from baseline in body weight, fasting glucose, total cholesterol, LDL cholesterol, or triglyceride levels.

There are no long-term double-blind clinical studies investigating the efficacy and tolerability of ziprasidone in children and adolescents.

There are no long-term clinical studies investigating the efficacy of ziprasidone in the prevention of recurrence of manic/depressive symptoms.

## **5.2 Pharmacokinetic properties**

### *Absorption*

Following oral administration of multiple doses of ziprasidone with food, peak serum concentrations typically occur 6 to 8 hours post-dose. The absolute bioavailability of a 20 mg dose is 60% in the fed state. Pharmacokinetic studies have demonstrated that the bioavailability of ziprasidone is increased by up to 100% in the presence of food. It is therefore recommended that ziprasidone should be taken with food.

In a multiple dose study ziprasidone oral suspension was shown to be bioequivalent to ziprasidone capsules under steady-state conditions. In a single dose administration study, equivalence was demonstrated with regard to AUC. The mean C<sub>max</sub> of the oral suspension was reduced by approximately 10-15% as compared with capsules. For the multiple dose study, the associated 90% confidence interval around this mean ratio was (79.6–101.0%).

#### *Distribution*

The volume of distribution is approximately 1.1 L/kg. Ziprasidone is more than 99% protein bound in serum.

#### *Biotransformation and elimination*

The mean terminal half-life of ziprasidone after oral administration is 6.6 hours. Steady state is reached within 1-3 days. Mean clearance of ziprasidone administered intravenously is 5 ml/min/kg. Approximately 20% of the dose is excreted in urine, and approximately 66% is eliminated in faeces.

Ziprasidone demonstrates linear kinetics over the therapeutic dose range of 40 to 80 mg twice daily in fed subjects.

Ziprasidone is extensively metabolised after oral administration with only a small amount excreted in urine (<1%) or faeces (<4%) as unchanged ziprasidone. Ziprasidone is primarily cleared via three proposed metabolic routes to yield four major circulating metabolites, benisothiazole piperazine (BITP) sulphoxide, BITP sulphone, ziprasidone sulphoxide and S-methyl-dihydroziprasidone. Unchanged ziprasidone represents about 44% of total drug-related material in serum.

Ziprasidone is primarily metabolised by two pathways: reduction and methylation to generate S-methyldihydroziprasidone which accounts for approximately two-thirds of the metabolism, and oxidative metabolism accounting for the other third. *In vitro* studies using human liver subcellular fractions indicate that S-methyldihydroziprasidone is generated in two steps. These studies indicate that the first step is mediated primarily by chemical reduction by glutathione as well as by enzymatic reduction by aldehyde oxidase. The second step is methylation mediated by thiol methyltransferase. *In vitro* studies indicate that CYP3A4 is the major cytochrome P450 catalysing the oxidative metabolism of ziprasidone with a potential minor contribution of CYP1A2.

Ziprasidone, S-methyl-dihydroziprasidone, and ziprasidone sulphoxide, when tested *in vitro*, share properties which may predict a QTc-prolonging effect. S-methyl-dihydroziprasidone is mainly eliminated in faeces by biliary excretion with a minor contribution by CYP3A4 catalysed metabolism. Ziprasidone sulphoxide is eliminated through renal excretion and by secondary metabolism catalysed by CYP3A4.

#### *Special populations*

Pharmacokinetic screening of patients has not revealed any significant pharmacokinetic differences between smokers and non-smokers.

No clinically significant age- or gender-differences in the pharmacokinetics of ziprasidone has been observed. The pharmacokinetics of ziprasidone in paediatric patients 10 to 17 years of age were similar to those in adults after correcting for the differences in body weights.

Consistent with the fact that renal clearance contributes very little to its overall clearance, no progressive increases in ziprasidone exposure were noted when ziprasidone was administered to subjects with varying degrees of renal function. Exposures in subjects with mild (creatinine clearance 30-60 ml/min), moderate (creatinine clearance 10-29 ml/min) and severe impairment (requiring dialysis) were 146%, 87% and 75% those of healthy subjects (creatinine clearance >70 ml/min) following oral administration of 20 mg BID for seven days. It is unknown whether serum concentrations of the metabolites are increased in these patients.

In mild to moderate impairment of liver function (Child Pugh A or B) caused by cirrhoses, the serum concentrations after oral administration were 30% higher and the terminal half-life was about 2 hours longer than in normal patients. The effect of liver impairment on the serum concentrations of the metabolites is unknown.

### **5.3 Preclinical safety data**

Preclinical safety data reveal no special hazard for humans based on conventional studies of safety pharmacology,

genotoxicity and carcinogenic potential. In reproductive studies in rats and rabbits, ziprasidone has shown no evidence of teratogenicity. Undesirable effects on fertility and decreased pup weights were observed at doses causing maternal toxicity such as decreased body weight gain. Increased perinatal mortality and delayed functional development of offspring occurred at maternal plasma concentrations extrapolated to be similar to the maximal concentrations in human given therapeutic doses.

## **6 PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

Water, purified  
Xylitol, E 967  
Sodium chloride  
Silica, colloidal anhydrous  
Cherry flavour  
Xanthan gum, E 415  
Methyl parahydroxybenzoate, E 218  
Sodium citrate, E 331  
Citric acid, anhydrous, E 330  
Polysorbate 80, E 433  
Propyl parahydroxybenzoate, E 216

### **6.2 Incompatibilities**

Geodon oral suspension should not be diluted or mixed into any food or beverages prior to administration.

### **6.3 Shelf life**

Shelf life of the medicinal product as packaged for sale: 2 years

Shelf life after first opening the bottle: 2 months.

### **6.4 Special precautions for storage**

This medicinal product does not require any special storage conditions.

### **6.5 Nature and contents of container**

HDPE bottles (60 ml or 240 ml) with polypropylene child-resistant closures and low density polyethylene liners.

Polypropylene oral syringe (2 ml or 8 ml) and polyethylene press-in bottle adaptor. The 2 ml oral syringe is calibrated in 0.25 ml (2.5 mg) increments; the 8 ml oral syringe is calibrated in 1.0 ml (10 mg increments).

Not all pack sizes may be marketed.

### **6.6 Special precautions for disposal**

No special requirements.

## **7 MARKETING AUTHORISATION HOLDER**

Pfizer Limited  
Ramsgate Road  
Sandwich  
Kent CT13 9NJ

United Kingdom

**8 MARKETING AUTHORISATION NUMBER**

PA 0019/052/006

**9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorization: 29 October 2004

Date of last renewal: 01 August 2010

**10 DATE OF REVISION OF THE TEXT**

January 2016