

BRISDELLE- paroxetine capsule

Legacy Pharma USA, Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use BRISDELLE safely and effectively. See full prescribing information for BRISDELLE.

BRISDELLE® (paroxetine) capsules, for oral use
Initial U.S. Approval: 1992

WARNING: SUICIDAL THOUGHTS AND BEHAVIORS

See full prescribing information for complete boxed warning.

Antidepressants, including selective serotonin reuptake inhibitors (SSRIs), increased the risk of suicidal thoughts and behavior in pediatric and young adult patients with major depressive disorder and other psychiatric disorder. Because BRISDELLE is an SSRI, closely monitor patients for clinical worsening and for emergence of suicidal thoughts and behaviors. BRISDELLE is not approved for use in pediatric and young adult patients (5.1).

INDICATIONS AND USAGE

BRISDELLE is a selective serotonin reuptake inhibitor (SSRI) indicated for the treatment of moderate to severe vasomotor symptoms associated with menopause (VMS) (1)

Limitations of Use: BRISDELLE is not indicated for the treatment of any psychiatric condition (1)

DOSAGE AND ADMINISTRATION

- The recommended dosage of BRISDELLE is 7.5 mg once daily, at bedtime (2.1)

DOSAGE FORMS AND STRENGTHS

Capsules: 7.5 mg (3)

CONTRAINDICATIONS

- Concurrent use with monoamine oxidase inhibitors (MAOI) or use within 14 days of MAOI use (2.2, 4.1, 5.2, 7.3)
- Use with thioridazine (4.2, 7.1)
- Use with pimozide (4.3, 7.1)
- Hypersensitivity to any ingredient in BRISDELLE (4.4)
- Pregnancy (4.5, 8.1)

WARNINGS AND PRECAUTIONS

- *Suicidality:* Monitor for suicidality or unusual changes in behavior (5.1)
- *Serotonin Syndrome:* BRISDELLE can cause serotonin syndrome with increased risk when co-administered with other serotonergic agents, but also when taken alone. If it occurs, discontinue BRISDELLE and serotonergic agents and initiate supportive measures (5.2, 7.3).
- *Tamoxifen:* Efficacy of tamoxifen may be reduced when administered concomitantly with BRISDELLE (5.3, 7.1)
- *Abnormal Bleeding:* Caution patients about the risk of bleeding associated with the concomitant use of BRISDELLE and non-steroidal anti-inflammatory drugs (NSAIDs), aspirin, or other drugs that affect coagulation (5.4, 7.1)
- *Angle-Closure Glaucoma:* Angle closure glaucoma has occurred in patients who have untreated anatomically narrow angles and who are treated with antidepressants. (5.5)
- *Hyponatremia:* Can occur in association with syndrome of inappropriate antidiuretic hormone secretion (SIADH) (5.6)
- *Bone Fracture:* Epidemiological studies have reported an association between SSRI treatment and fractures (5.7)
- *Activation of Mania/Hypomania:* Screen for bipolar disorder and monitor for mania/ hypomania (5.8)
- *Seizures:* Use cautiously in patients with a history of seizures or with conditions that potentially lower the seizure threshold (5.9)
- *Sexual Dysfunction:* BRISDELLE use may cause symptoms of sexual dysfunction. (5.12)

ADVERSE REACTIONS

The most common adverse reactions ($\geq 2\%$) reported in clinical trials were: headache, fatigue, and nausea/vomiting (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Legacy Pharma Inc. at 1-800-727-7151 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

DRUG INTERACTIONS

Paroxetine is a strong CYP2D6 inhibitor. Co-administration of BRISDELLE can alter concentrations of other drugs that are metabolized by CYP2D6. Consider potential drug interactions prior to and during therapy (5.3, 7.1, 7.3). See Full Prescribing Information for a list of clinically significant drug interactions (7.1, 7.2, 7.3)

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.

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FULL PRESCRIBING INFORMATION

WARNING: SUICIDAL THOUGHTS AND BEHAVIORS

Selective serotonin reuptake inhibitors (SSRIs) increased the risk of suicidal thoughts and behavior in pediatric and young adult patients in short-term trials for the treatment of major depressive disorder and other psychiatric disorders. Because BRISDELLE is an SSRI, closely monitor BRISDELLE-treated patients closely for emergence of suicidal thoughts and behaviors [see Warnings and Precautions (5.1)].

BRISDELLE is not approved for use in any psychiatric condition or in pediatric and young adult patients [see Indications and Usage (1) and Use in Specific Populations (8.4)].

1 INDICATIONS AND USAGE

BRISDELLE is indicated for the treatment of moderate to severe vasomotor symptoms (VMS) associated with menopause.

Limitations of Use:

BRISDELLE is not indicated for the treatment of any psychiatric condition. BRISDELLE has a lower recommended paroxetine dosage than that used to treat major depressive disorder, obsessive compulsive disorder, panic disorder, generalized anxiety disorder, social anxiety disorder, and post-traumatic stress disorder. The safety and effectiveness of the lower BRISDELLE dosage has not been established for any psychiatric condition. Patients who require paroxetine for treatment of a psychiatric condition should discontinue BRISDELLE and initiate a paroxetine-containing product that is indicated for such use.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage

The recommended oral dosage of BRISDELLE for the treatment of moderate to severe VMS associated with menopause is 7.5 mg once daily, at bedtime, with or without food [see *Clinical Pharmacology (12.3)*].

2.2 Use of BRISDELLE Before or After a Monoamine Oxidase Inhibitor

Wait at least 14 days after discontinuation of a monoamine oxidase inhibitor (MAOI) before initiating therapy with BRISDELLE. Conversely, allow at least 14 days after stopping BRISDELLE before starting an MAOI [see *Contraindications (4.1)*, *Warnings and Precautions (5.2)* and *Drug Interactions (7.3)*].

3 DOSAGE FORMS AND STRENGTHS

BRISDELLE is available as 7.5 mg pink capsules printed with black edible ink with “BRISDELLE” and “7.5 mg” on the capsule. Each capsule contains 9.69 mg of paroxetine mesylate equivalent to 7.5 mg paroxetine base.

4 CONTRAINDICATIONS

BRISDELLE is contraindicated in patients:

- Taking, or within 14 days of stopping, MAOIs (including the MAOIs linezolid and intravenous methylene blue) because of an increased risk of serotonin syndrome [see *Warnings and Precautions (5.2)*, *Drug Interaction (7)*].
- Taking thioridazine because of risk of QT prolongation [see *Warnings and Precautions (5.3)*, *Drug Interaction (7)*].
- Taking pimozide because of risk of QT prolongation [see *Warnings and Precautions (5.3)*, *Drug Interaction (7)*].
- With known hypersensitivity (e.g., anaphylaxis, angioedema, Stevens-Johnson syndrome) to paroxetine or to any of the inactive ingredients in BRISDELLE [see *Adverse Reactions (6.2)*]
- Who are or become pregnant because menopausal VMS does not occur during pregnancy and BRISDELLE may cause fetal harm [see *Use in Specific Populations (8.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Suicidal Thoughts and Behaviors in Adolescents and Young Adults

SSRIs increased the risk of suicidal thoughts and behavior in pediatric and young adult patients in short-term trials for the treatment of major depressive disorder (MDD) and other psychiatric disorders - BRISDELLE is not approved for use in any psychiatric condition or in pediatric and young adult patients [see *Indications and Usage (1)* and *Use in Specific Populations (8.4)*]. There is limited information regarding suicidal thoughts and behaviors in females who use BRISDELLE for treatment of moderate to severe VMS associated with menopause. The BRISDELLE trials excluded females with a presence or history of previous psychiatric disorders.

Monitor all BRISDELLE-treated patients for any emergence of suicidal thoughts and behaviors, especially during the initial few months of BRISDELLE therapy. Counsel family

members to monitor for changes in behavior and to alert the health care provider if such changes occur. Consider discontinuing BRISDELLE in patients who experience emergent suicidal thoughts or behaviors or symptoms that might be precursors to suicidal thoughts or behaviors, especially if these symptoms are severe, abrupt in onset, or were not part of the patient's presenting symptoms.

5.2 Serotonin Syndrome

BRISDELLE can precipitate serotonin syndrome, a potentially life-threatening condition. The risk is increased with concomitant use of other serotonergic agents (including triptans, tricyclic antidepressants, fentanyl, tramadol, meperidine, methadone, lithium, tryptophan, buspirone, amphetamines, and St. John's Wort) and with drugs that impair metabolism of serotonin, i.e., monoamine oxidase inhibitors (MAOIs) [see *Contraindications (4), Drug Interactions (7.3)*]. Serotonin syndrome can also occur when BRISDELLE is used alone.

Serotonin syndrome symptoms may include mental status changes (e.g., agitation, hallucinations, delirium, and coma), autonomic instability (e.g., tachycardia, labile blood pressure, dizziness, diaphoresis, flushing, hyperthermia), neuromuscular symptoms (e.g., tremor, rigidity, myoclonus, hyperreflexia, incoordination), and/or gastrointestinal symptoms (e.g., nausea, vomiting, diarrhea).

The concomitant use of BRISDELLE with MAOIs is contraindicated. Do not start BRISDELLE in a patient who is being treated with MAOIs such as linezolid or intravenous methylene blue. All reports with methylene blue that provided information on the route of administration involved intravenous administration in the dosage range of 1 mg/kg to 8 mg/kg. No reports involved the administration of methylene blue by other routes (such as oral or local tissue injection) or at lower dosages. If it is necessary to initiate treatment with an MAOI such as linezolid or intravenous methylene blue in a patient taking BRISDELLE, the patient should be taken off BRISDELLE before initiating treatment with the MAOI [see *Contraindications (4.1)*]. If concomitant use of BRISDELLE with other serotonergic drugs (besides MAOIs) is clinically warranted, consider the increased risk of serotonin syndrome and carefully observe the patient, particularly during BRISDELLE initiation [see *Contraindications (4.1) Drug Interactions (7.3)*].

Monitor all patients taking BRISDELLE for the emergence of serotonin syndrome. Discontinue BRISDELLE and any concomitant serotonergic agents immediately if the above events occur and initiate supportive symptomatic treatment.

5.3 Potential Impact on Tamoxifen Efficacy

Some studies have shown that the efficacy of tamoxifen, as measured by the risk of breast cancer relapse/mortality, may be reduced when concomitantly administered with paroxetine as a result of paroxetine's irreversible inhibition of CYP2D6 and lower tamoxifen blood levels [see *Drug Interactions (7.1)*]. However, other studies have failed to demonstrate such a risk.

When tamoxifen is used for the treatment or prevention of breast cancer, weigh the likely benefit of BRISDELLE for treating moderate to severe VMS associated with menopause vs. the risk of possible decreased tamoxifen effectiveness, and consider avoiding the concomitant use of BRISDELLE.

5.4 Increased Risk of Bleeding

SSRIs, including BRISDELLE, increased the risk of bleeding events. Concomitant use of aspirin, nonsteroidal anti-inflammatory drugs (NSAIDs), other antiplatelet drugs, warfarin, and other anticoagulants may add to this risk [see *Drug Interactions (7.3)*]. Case reports and epidemiological studies (case-control and cohort design) have demonstrated an association between use of drugs that interfere with serotonin reuptake and the occurrence of gastrointestinal bleeding. Bleeding events related to SSRIs have ranged from ecchymosis, hematoma, epistaxis, and petechiae to life-threatening hemorrhages.

Inform patients about the risk of bleeding associated with the concomitant use of BRISDELLE and antiplatelet agents or anticoagulants [see *Drug Interactions (7.1)*]. For patients taking warfarin, carefully monitor the international normalized ratio.

5.5 Angle-Closure Glaucoma

The pupillary dilation that occurs following use of SSRIs, including BRISDELLE, may trigger an angle closure attack in a patient with anatomically narrow angles who does not have a patent iridectomy. Cases of angle-closure glaucoma associated with use of paroxetine have been reported. Avoid use of SSRIs, including BRISDELLE, in patients with untreated anatomically narrow angles.

5.6 Hyponatremia

Hyponatremia may occur as a result of treatment with SSRIs, including BRISDELLE. Cases with serum sodium lower than 110 mmol/L have been reported in patients using SSRIs. Geriatric patients, patients taking diuretics, and those who are volume-depleted may be at greater risk of developing hyponatremia with SSRIs [see *Use in Specific Populations (8.5)*]. Signs and symptoms of hyponatremia include headache, difficulty concentrating, memory impairment, confusion, weakness, and unsteadiness, which can lead to falls. Signs and symptoms associated with more severe and/or acute cases have included hallucination, syncope, seizure, coma, respiratory arrest, and death. In many cases, the hyponatremia appears to be the result of the syndrome of inappropriate antidiuretic hormone secretion (SIADH).

In patients with symptomatic hyponatremia, discontinue BRISDELLE and institute appropriate medical intervention.

5.7 Bone Fracture

Epidemiological studies on bone fracture risk following exposure to SSRIs have reported an association between SSRI treatment and fractures. It is unknown to what extent fracture risk is directly attributable to SSRI treatment. If a BRISDELLE-treated patient presents with unexplained bone pain, point tenderness, swelling, or bruising, consider the possibility of a fragility fracture.

5.8 Screening Patients for Bipolar Disorder and Monitoring for Mania/Hypomania

BRISDELLE is only indicated for the treatment of moderate to severe VMS and is not approved for use in treating either depression or bipolar depression. However, prior to initiating treatment with BRISDELLE, all patients should be adequately screened to determine if they are at risk for bipolar disorder; such screening should include a detailed psychiatric history, including a family history of suicide, bipolar disorder, and

depression. It is generally believed (though not established in controlled trials) that use of an antidepressant alone may increase the likelihood of precipitation of a mixed/manic episode in patients at risk for bipolar disorder.

5.9 Seizures

In clinical studies of another paroxetine product, seizures occurred in 0.1% of paroxetine-treated patients.

Use BRISDELLE cautiously in patients with a history of seizures or with conditions that potentially lower the seizure threshold. Discontinue BRISDELLE in any patient who develops seizures.

5.10 Sexual Dysfunction

Use of SSRIs, including BRISDELLE, may cause symptoms of sexual dysfunction. In female patients, SSRI use may result in decreased libido and delayed or absent orgasm.

It is important for prescribers to inquire about sexual function prior to initiation of BRISDELLE and to inquire specifically about changes in sexual function during treatment because sexual function may not be spontaneously reported. When evaluating changes in sexual function, obtaining a detailed history (including timing of symptom onset) is important because sexual symptoms may have other causes. Discuss potential management strategies to support patients in making informed decisions about treatment.

6 ADVERSE REACTIONS

The following serious adverse reactions are discussed elsewhere in labeling:

- Suicidality [see *Warnings and Precautions (5.1)*]
- Serotonin syndrome [see *Warnings and Precautions (5.2)*]
- Abnormal bleeding [see *Warnings and Precautions (5.4)*]
- Angle-Closure Glaucoma [see *Warnings and Precautions (5.5)*]
- Hyponatremia [see *Warnings and Precautions (5.6)*]
- Bone Fracture [see *Warnings and Precautions (5.7)*]
- Mania/Hypomania [see *Warnings and Precautions (5.8)*]
- Seizure [see *Warnings and Precautions (5.9)*]

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 directly be compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The data described below reflect exposure to BRISDELLE in the following randomized, placebo-controlled trials for the treatment of moderate to severe VMS associated with menopause [see *Clinical Studies (14)*]: (1) one 8-week Phase 2 trial, (2) one 12-week Phase 3 trial and (3) one 24-week Phase 3 trial. In these trials, a total of 635 postmenopausal females received BRISDELLE 7.5 mg administered orally once daily and 641 postmenopausal females received placebo. In these trials, 68% were White, 30% were Black or African American (30%), and 2% were other races, with a mean age of 55 years (range 40 to 73 years). Postmenopausal females with a history of suicidal ideation

or suicidal behavior were excluded from these trials.

Serious Adverse Reactions:

In the pooled Phase 2 and Phase 3 trials, three BRISDELLE-treated patients reported a serious adverse reaction of suicidal ideation and one BRISDELLE-treated patient reported a serious adverse reaction of suicide attempt. There were no serious adverse reactions of suicidal ideation or suicide attempt reported among the placebo-treated patients.

Adverse Reactions Leading to Study Discontinuation:

A total of 4.7% of females taking BRISDELLE discontinued from the clinical trials due to an adverse reaction, compared to 3.7% of females on placebo; the most frequent adverse reactions leading to discontinuation among paroxetine-treated females were: abdominal pain (0.3%), attention disturbances (0.3%), headache (0.3%), and suicidal ideation (0.3%).

Common Adverse Reactions:

Overall, based on investigators' determinations about what events were likely to be drug-related, about 20% of postmenopausal females treated with BRISDELLE reported at least 1 adverse reaction in the three controlled trials. The most common adverse reactions ($\geq 2\%$ and at a higher incidence in BRISDELLE-treated females compared to placebo-treated females) reported in these trials were headache, fatigue/malaise/lethargy, and nausea/vomiting. Of these commonly reported adverse reactions, nausea occurred primarily within the first 4 weeks of BRISDELLE treatment and fatigue occurred primarily within the first week of BRISDELLE treatment, and decreased in frequency with continued therapy.

The adverse reactions that occurred in $\geq 2\%$ of BRISDELLE-treated patients and at a higher incidence in BRISDELLE-treated females compared to placebo-treated females are shown in Table 1 for the pooled Phase 2 and Phase 3 trials.

Table 1: Incidence of Common Adverse Reactions in the Phase 2 and Phase 3 Trials of Postmenopausal Females with Moderate to Severe VMS¹

	Incidence n (%)	
	BRISDELLE (n = 635)	BRISDELLE (n = 641)
Headache	40 (6.3)	31 (4.8)
Fatigue, malaise, lethargy	31 (4.9)	18 (2.8)
Nausea, vomiting	27 (4.3)	15 (2.3)

¹ $\geq 2\%$ BRISDELLE-treated patients and at a higher incidence in BRISDELLE-treated females compared to placebo-treated females.

Adverse Reactions After Discontinuing BRISDELLE

Certain symptoms were seen more frequently in postmenopausal females at the time of discontinuation of BRISDELLE compared to discontinuation of placebo and have also been reported upon discontinuation of other paroxetine products, particularly after abrupt discontinuation. Adverse reactions reported after discontinuation of BRISDELLE

included increased dreaming/nightmares, muscle cramps/spasms/twitching, headache, nervousness/anxiety, fatigue/tiredness, restless feeling in legs, and trouble sleeping/insomnia. While these adverse reactions were generally self-limiting, there have been reports of serious discontinuation symptoms with other paroxetine products.

6.2 Postmarketing Experience

The following adverse reactions have been identified during post-approval use of this and other paroxetine products. Because some of 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.

Blood and Lymphatic System Disorders: Idiopathic thrombocytopenic purpura, Events related to impaired hematopoiesis (including aplastic anemia, pancytopenia, bone marrow aplasia, agranulocytosis).

Cardiac Disorders: Atrial fibrillation, Pulmonary edema, Ventricular fibrillation, Ventricular tachycardia (including torsades de pointes).

Gastrointestinal Disorders: Pancreatitis, Pancreatitis hemorrhagic, Vomiting.

General Disorders and Administration Site Conditions: Death, Drug withdrawal syndrome, Malaise.

Hepatobiliary Disorders: Drug-induced liver injury, Hepatic failure, Jaundice.

Immune System Disorders: Anaphylaxis, Angioedema, Toxic epidermal necrolysis.

Investigations: Elevated liver tests (the most severe cases were deaths due to liver necrosis, and grossly elevated transaminases associated with severe liver dysfunction).

Metabolism and Nutrition Disorders: Diabetes mellitus inadequate control, Type 2 diabetes mellitus.

Nervous System Disorders: Neuroleptic malignant syndrome, Paresthesia, Somnolence, Tremor, Anosmia, Hyposmia

Psychiatric Disorders: Aggression, Agitation, Anxiety, Confusional state, Depression, Disorientation, Homicidal ideation, Insomnia, Restlessness.

Respiratory, Thoracic and Mediastinal Disorders: Pulmonary hypertension.

Skin and Subcutaneous Tissue Disorders: Hyperhidrosis, Stevens-Johnson syndrome, Drug reaction with eosinophilia and systemic symptoms (DRESS).

7 DRUG INTERACTIONS

7.1 Potential for BRISDELLE to Affect Other Drugs

Paroxetine is a strong CYP2D6 inhibitor. Clinical drug interaction studies have been performed with substrates of CYP2D6 and show that paroxetine can inhibit the metabolism of drugs metabolized by CYP2D6 [see *Clinical Pharmacology (12.3)*]. Table 2 contains examples of drugs with a metabolism that may be affected by concomitant use with BRISDELLE.

Table 2 Effects of Paroxetine on Other Drugs

Concomitant Drug Name	Effect of Paroxetine on Other Drugs	Clinical Recommendations
Thioridazine	Increased plasma concentrations of thioridazine Potential QTc prolongation	Concomitant use of thioridazine and BRISDELLE is contraindicated.
Pimozide	Increased plasma concentrations of pimozide. Potential QTc prolongation	Concomitant use of pimozide and BRISDELLE is contraindicated.
Tamoxifen	Reduced plasma concentrations of active tamoxifen metabolite	Consider avoiding concomitant use of tamoxifen and BRISDELLE.
Tricyclic Antidepressants (TCA) (e.g., Desipramine)	Increased plasma concentrations and elimination half-life	Plasma TCA concentrations may need to be monitored and the TCA dosage may need to be reduced if a TCA is used concomitantly with BRISDELLE. Monitor tolerability.
Risperidone	Increased plasma concentrations of risperidone	A lower risperidone dosage may be necessary (see the risperidone Prescribing Information for). Monitor tolerability.
Atomoxetine	Increased exposure of atomoxetine	A lower atomoxetine dosage of may be necessary (see atomoxetine Prescribing Information for). Monitor tolerability.
Drugs Highly Bound to Plasma Protein (e.g., Warfarin)	Increased free plasma concentrations	The warfarin dosage may need to be reduced. Monitor tolerability and the International Normalized Ratio.
Digoxin	Decreased plasma concentrations of digoxin	The digoxin dosage of may need to be increased. Monitor digoxin concentrations and clinical effect.
Theophylline	Increased plasma concentrations of theophylline	The theophylline dosage of may need to be decreased. Monitor theophylline concentrations and tolerability.

Use caution with concomitant use of BRISDELLE with other drugs that are metabolized by CYP2D6, including nortriptyline, amitriptyline, imipramine, desipramine, fluoxetine, phenothiazines, risperidone, and Type 1C antiarrhythmics (e.g., propafenone, flecainide, and encainide).

7.2 Potential for Other Drugs to Affect BRISDELLE

The metabolism and pharmacokinetics of paroxetine may be affected by the induction and inhibition of drug metabolizing enzymes such as CYP2D6. Table 3 contains a list of drugs that may affect the pharmacokinetics of BRISDELLE when administered concomitantly [see *Clinical Pharmacology (12.3)*].

Table 3 Effects of Other Drugs on Paroxetine

Concomitant Drug Name	Effect of Concomitant Drug on Paroxetine	Clinical Recommendations
Phenobarbital	Decreased paroxetine exposure	Monitor clinical effect of BRISDELLE. No BRISDELLE dosage adjustment is needed.
Phenytoin	Decreased paroxetine exposure	
Fosamprenavir/Ritonavir	Decreased plasma concentration of paroxetine	
Cimetidine	Increased plasma concentration of paroxetine	

Use caution if BRISDELLE is used concomitantly with other drugs that inhibit CYP2D6 (e.g., quinidine).

7.3 Other Potentially Significant Drug Interactions

Monoamine Oxidase Inhibitors

Serious adverse reactions such as serotonin syndrome have been reported in patients treated with a SSRI and a concomitant monoamine oxidase inhibitor (MAOI), in patients started on an SSRI who recently received an MAOI and in patients started on an MAOI who recently received an SSRI. Therefore, concomitant use of MAOIs with BRISDELLE or use of BRISDELLE and an MAOI within 14 days of each other is contraindicated [see *Dosage and Administration (2.2)*, *Contraindications (4.1)* and *Warnings and Precautions (5.2)*].

Serotonergic Drugs

If concomitant use of BRISDELLE with other serotonergic drugs (e.g., other SNRIs, SSRIs, triptans, tricyclic antidepressants, opioids, lithium, tryptophan, buspirone, amphetamines, and St. John's Wort) is clinically warranted, consider the increased risk of serotonin syndrome and carefully observe the patient, particularly during treatment initiation [see *Warnings and Precautions (5.2)*].

An interaction between paroxetine and tryptophan may occur when they are co-administered. Adverse reactions, consisting primarily of headache, nausea, sweating, and dizziness, have been reported when tryptophan was administered to patients taking paroxetine. Consequently, concomitant use of BRISDELLE with tryptophan is not recommended.

Drugs that Interfere with Hemostasis (e.g., NSAIDs, Aspirin, and Warfarin)

Altered anticoagulant effects, including increased bleeding, have been reported when SSRIs are concomitantly administered with NSAIDs, aspirin, warfarin, or other drugs that affect coagulation. There may be a pharmacodynamic interaction between paroxetine and warfarin that causes an increased bleeding diathesis despite unaltered prothrombin time. Carefully monitor patients receiving warfarin therapy when BRISDELLE is initiated or discontinued [see *Warnings and Precautions (5.4)*].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

BRISDELLE is contraindicated in pregnant females and not indicated for use in premenopausal females. Based on epidemiologic and animal studies, paroxetine can cause fetal harm. Based on data from published observational studies, exposure to SSRIs, particularly in the month before delivery, has been associated with a less than 2-fold increase in the risk of postpartum hemorrhage [see *Warnings and Precautions (5.4)*].

Paroxetine is associated with a less than 2-fold increase in cardiovascular malformations when administered to a pregnant female during the first trimester. While individual epidemiological studies on the association between paroxetine use and cardiovascular malformations have reported inconsistent findings, some meta-analyses of epidemiological studies have identified an increased risk of cardiovascular malformations (see *Data*). There are risks of persistent pulmonary hypertension of the newborn (PPHN) (see *Data*) and/or poor neonatal adaptation with exposure to selective serotonin reuptake inhibitors (SSRIs), including BRISDELLE, during pregnancy.

No evidence of treatment related malformations was observed in animal reproduction studies, when paroxetine was administered during the period of organogenesis at doses up to 50 mg/kg/day in rats and 6 mg/kg/day in rabbits. These doses are approximately 64 times (rat) and less than 16 times (rabbit) the maximum recommended human dose (MRHD) of BRISDELLE (7.5 mg) on an mg/m² basis. When paroxetine was administered to female rats during the last trimester of gestation and continued through lactation, there was an increase in the number of pup deaths during the first four days of lactation. This effect occurred at a dose of 1 mg/kg/day which is 1.3 times the MRHD on an mg/m² basis (see *Data*).

Data

Human Data: Published epidemiological studies on the association between first trimester paroxetine use and cardiovascular malformations have reported inconsistent results; however, meta-analyses of population-based cohort studies published between 1996-2017 indicate a less than 2-fold increased risk for overall cardiovascular malformations. Specific cardiac malformations identified in two meta-analyses include approximately 2 to 2.5-fold increased risk for right ventricular outflow tract defects. One meta-analysis also identified an increased risk (less than 2-fold) for bulbus cordis anomalies and anomalies of cardiac septal closure, and an increased risk for atrial septal defects (pooled OR 2.38, 95% CI 1.14-4.97). Important limitations of the studies included in these meta-analyses include potential confounding by indication, depression severity, and potential exposure misclassification.

Exposure to SSRIs, particularly later in pregnancy, may have an increased risk for PPHN. PPHN occurs in 1-2 per 1000 live births in the general population and is associated with substantial neonatal morbidity and mortality.

Animal Data: Reproduction studies were performed at doses up to 50 mg/kg/day in rats and 6 mg/kg/day in rabbits administered during organogenesis. These doses are approximately 64 times (rat) and less than 16 times (rabbit) MRHD of BRISDELLE (7.5 mg) on an mg/m² basis. These studies have revealed no evidence of malformations. However, in rats, there was an increase in pup deaths during the first 4 days of lactation when dosing occurred during the last trimester of gestation and continued throughout lactation. This effect occurred at a dose of 1 mg/kg/day which is 1.3 times the MRHD on an mg/m² basis. The no effect dose for rat pup mortality was not determined. The cause of these deaths is not known.

8.4 Pediatric Use

Safety and effectiveness of BRISDELLE in pediatric patients have not been established; BRISDELLE is not indicated in the pediatric population.

8.5 Geriatric Use

Clinical trials of BRISDELLE did not include sufficient numbers of patients aged 65 and over to determine whether they respond differently from younger adult patients. Patients 65 years of age and older may have elevated paroxetine plasma concentrations compared to younger adult patients. However, the recommended BRISDELLE dosage in patients 65 years of age and older is that same as that as younger patients [see *Clinical Pharmacology* (12.3)].

SSRIs have been associated with cases of clinically significant hyponatremia in geriatric patients, who may be at greater risk for this adverse reaction [see *Warnings and Precautions* (5.6)].

8.6 Renal Impairment

The recommended BRISDELLE dosage in patients with renal impairment is the same as those with normal renal function [see *Clinical Pharmacology* (12.3)].

8.7 Hepatic Impairment

The recommended BRISDELLE dosage in patients with hepatic impairment is the same as those with normal hepatic function [see *Clinical Pharmacology* (12.3)].

10 OVERDOSAGE

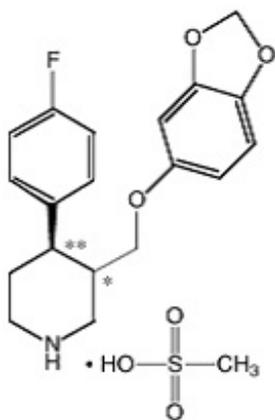
The following have been reported with paroxetine tablets overdose:

- Seizures, which may be delayed, and altered mental status including coma.
- Cardiovascular toxicity, which may be delayed, including QRS and QTc interval prolongation. Hypertension most commonly seen, but rarely can see hypotension alone or with co-ingestants including alcohol.
- Serotonin syndrome (patients with a multiple drug overdose with other pro-serotonergic drugs may have a higher risk).

Consider contacting the Poison Help Line (1-800-221-2222) or a medical toxicologist for overdose management recommendations.

11 DESCRIPTION

BRISDELLE (paroxetine) is an orally administered selective serotonin reuptake inhibitor (SSRI) for the treatment of moderate to severe VMS associated with menopause. It is identified chemically as (-)-*trans*-4R-(4'-fluorophenyl)-3S-[(3',4'-methylenedioxyphenoxy)methyl]piperidine mesylate and has the empirical formula of $C_{19}H_{20}FNO_3 \cdot CH_3SO_3H$. The molecular weight is 425.5 (329.4 as free base). The structural formula is:



The mesylate salt of paroxetine is an odorless, off-white powder, having a melting point range of 147° to 150°C and a solubility of more than 1 g/mL in water.

Each pink capsule contains 9.69 mg of paroxetine mesylate equivalent to 7.5 mg paroxetine base.

Inactive ingredients consist of: dibasic calcium phosphate, sodium starch glycolate, magnesium stearate, gelatin, titanium dioxide, FD&C Yellow #6, FD&C Red #3, FD&C Red #40, shellac, and black iron oxide.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Nonclinical studies have shown that paroxetine is an SSRI; BRISDELLE is not an estrogen. The mechanism of action of BRISDELLE for the treatment of moderate to

severe VMS associated with menopause is unknown.

Studies at clinically relevant BRISDELLE doses in humans have demonstrated that paroxetine blocks the uptake of serotonin into human platelets. Paroxetine is a neuronal serotonin reuptake inhibitor with weak effects on norepinephrine and dopamine neuronal reuptake in vitro. Paroxetine also has low affinity for muscarinic α_1 -, α_2 -, beta-adrenergic, dopamine (D2)-, 5-HT₁-, 5-HT₂-, and histamine (H₁)-receptors.

12.2 Pharmacodynamics

The exposure-response relationship and time course of pharmacodynamic response for the safety and effectiveness of paroxetine for the treatment of moderate to severe VMS associated with menopause have not been fully characterized.

12.3 Pharmacokinetics

Absorption

Paroxetine is completely absorbed after oral dosing.

In a study in which healthy postmenopausal females (n=24) received BRISDELLE 7.5 mg once daily for 14 days, steady-state paroxetine concentrations were achieved by approximately 12 days of dosing for most subjects, although it may take substantially longer in an occasional patient. Peak concentrations were reached at a median of 6 hours (3 to 8 hours range). Steady-state mean values of C_{max} , C_{min} , and AUC_{0-last} were 13.10 ng/mL (CV 91%), 7.17 ng/mL (CV 99%), and 237 hr*ng/mL (CV 94%), respectively.

Steady-state AUC_{0-24} values were about 3 times those of AUC_{0-inf} following a single BRISDELLE dose, indicating non-linear pharmacokinetics. Steady-state C_{max} values were approximately 5 times greater than those attained after a single BRISDELLE dose and steady-state exposure based on AUC_{0-24} was about 10 times greater than AUC_{0-24} after a single dose.

The nonlinear kinetics and excess accumulation are due to the fact that CYP2D6, an enzyme that is in part responsible for paroxetine metabolism, is readily saturable.

Effect of Food: The effects of food on the bioavailability of paroxetine were studied after administration of another paroxetine product with a dosage higher than BRISDELLE. AUC was slightly increased (6%) when paroxetine was administered with food and the C_{max} was 29% greater, while the time to reach peak plasma concentration decreased from 6.4 hours post-dosing to 4.9 hours. These changes are not clinically significant [see *Dosage and Administration (2.1)*].

Distribution

Paroxetine distributes throughout the body, including the central nervous system, with only 1% remaining in the plasma.

Approximately 95% and 93% of paroxetine is bound to plasma protein at 100 ng/mL and 400 ng/mL, respectively. Paroxetine does not alter the *in vitro* protein binding of phenytoin or warfarin.

Elimination

Metabolism: Paroxetine is extensively metabolized after oral administration. The principal

metabolites are polar and conjugated products of oxidation and methylation, which are readily cleared. Conjugates with glucuronic acid and sulfate predominate, and major metabolites have been isolated and identified. Data indicate that the metabolites have no more than 1/50 the potency of the parent compound at inhibiting serotonin uptake. The metabolism of paroxetine is accomplished in part by cytochrome CYP2D6. The role of this enzyme in paroxetine metabolism also suggests potential drug interactions [see *Drug Interactions (7)*]. At steady state, when the CYP2D6 pathway is essentially saturated, paroxetine clearance is governed by alternative P450 isozymes, which, unlike CYP2D6, show no evidence of saturation.

Excretion: Approximately 64% of a 30 mg oral solution of another paroxetine product (four times the recommended BRISDELLE dosage) was excreted in the urine with 2% as the parent compound and 62% as metabolites over a 10-day post-dosing period. About 36% of the dose was excreted in the feces (probably via the bile), mostly as metabolites and less than 1% as the parent compound over the 10-day post-dosing period.

Specific Populations

Patients with Renal and Hepatic Impairment: Increased plasma concentrations of paroxetine occur in subjects with renal and hepatic impairment. The mean plasma concentration in patients with creatinine clearance below 30 mL/min was approximately 4 times greater than seen in normal volunteers. Patients with creatinine clearance of 30 to 60 mL/min and patients with hepatic impairment had about a 2-fold increase in plasma concentrations (AUC, C_{max}) [see *Use in Specific Populations (8.6, 8.7)*].

Geriatric Patients: In a multiple-dose study in geriatric patients with another paroxetine product at doses of 20, 30, and 40 mg (1.7, 4, and 5.3, times the maximum recommended BRISDELLE dosage, respectively), C_{min} concentrations were about 70% to 80% greater than the respective C_{min} concentrations in younger adult subjects [see *Use in Specific Populations (8.5)*].

Drug Interaction Studies

Potential Effect of BRISDELLE on Other Drugs

- **Drugs Metabolized by CYP3A4:** An *in vivo* drug interaction study involving the concomitant use of paroxetine and terfenadine, a substrate for cytochrome CYP3A4, under steady-state conditions revealed no effect of paroxetine on terfenadine pharmacokinetics. *In vitro* studies have shown ketoconazole, a potent CYP3A4 inhibitor, to be at least 100 times more potent than paroxetine as an inhibitor of the metabolism of several substrates for CYP3A4, including astemizole, triazolam, and cyclosporine. Based on the assumption that the relationship between paroxetine's *in vitro* K_i and its lack of effect on terfenadine's *in vivo* clearance predicts its effect on other CYP3A4 substrates, paroxetine's extent of CYP3A4 inhibition is not likely to be of clinical significance.
- **Drugs Metabolized by CYP2D6:** Many drugs are metabolized by the cytochrome P450 isozyme CYP2D6. Like other agents that are metabolized by CYP2D6, paroxetine may significantly inhibit the activity of this isozyme. In most patients (> 90%), this CYP2D6 isozyme is saturated early during paroxetine dosing. Specific studies investigating the effect of paroxetine on drugs metabolized by CYP2D6 are listed below:
 - **Pimozide:** In a controlled study of healthy volunteers, after another paroxetine product was titrated to 60 mg daily (8 times the maximum recommended BRISDELLE dosage), concomitant use of paroxetine with a single 2 mg dose of pimozide was associated with mean increases in pimozide AUC of 151% and C_{max}

- of 62%, compared to pimozide administered alone [see *Drug Interactions (7.1)*].
- Desipramine: In one study, daily dosing of another paroxetine product (20 mg once daily) (2.7 times the recommended BRISDELLE dosage) under steady-state conditions with a concomitant single dose of desipramine (100 mg) increased desipramine C_{max} , AUC, and $T_{1/2}$ by an average of approximately 2-, 5-, and 3-fold, respectively [see *Drug Interactions (7.1)*].
 - Risperidone: Daily dosing of another paroxetine product at 20 mg in patients stabilized on risperidone (4 to 8 mg/day), a CYP2D6 substrate, increased mean plasma risperidone concentrations approximately 4-fold, decreased 9-hydroxyrisperidone concentrations approximately 10%, and increased concentrations of the active moiety (the sum of risperidone plus 9-hydroxyrisperidone) approximately 1.4-fold [see *Drug Interactions (7.1)*].
 - Atomoxetine: The effect of paroxetine on the pharmacokinetics of atomoxetine has been evaluated when both drugs were at steady state. In healthy volunteers who were extensive metabolizers of CYP2D6, 20 mg daily of another paroxetine product was given in combination with 20 mg atomoxetine every 12 hours resulting in increases in steady-state atomoxetine AUC values that were 6- to 8-fold greater and in atomoxetine C_{max} values that were 3- to 4-fold greater than when atomoxetine was given alone [see *Drug Interactions (7.1)*].
 - Digoxin: Mean digoxin AUC at steady state decreased by 15% in the presence of paroxetine [see *Drug Interactions (7.1)*].
 - Beta Blockers: In a study in which propranolol (80 mg twice daily) was dosed orally for 18 days, the steady-state plasma concentrations of propranolol were unaltered during concomitant use with another paroxetine product (30 mg once daily) (4 times the recommended BRISDELLE dosage) for the final 10 days. The effects of propranolol on paroxetine have not been evaluated.

Potential Effect of Other Drugs on BRISDELLE

Concomitant use of paroxetine with other drugs that alter CYP enzymes activities including CYP2D6 may affect the plasma concentrations of paroxetine. Specific studies investigating the effect of other drugs on paroxetine are listed below:

- Cimetidine: Cimetidine inhibits many cytochrome P450 enzymes. In a study in which another paroxetine product was dosed orally at 30 mg once daily (4 times the recommended BRISDELLE dosage) for 4 weeks, steady-state plasma concentrations of paroxetine were increased by approximately 50% during concomitant use with oral cimetidine (300 mg three times daily) for the final week [see *Drug Interactions (7.2)*].
- Phenobarbital: Phenobarbital induces many cytochrome P450 enzymes. When a single oral 30 mg dose of another paroxetine product (4 times the recommended BRISDELLE dosage) was administered at phenobarbital steady state (100 mg once daily for 14 days), paroxetine AUC and $T_{1/2}$ were reduced (by an average of 25% and 38%, respectively) compared to paroxetine administered alone. The effect of paroxetine on phenobarbital pharmacokinetics was not studied. Because paroxetine exhibits nonlinear pharmacokinetics, the results of this study may not address the case where the two drugs are both being chronically dosed [see *Drug Interactions (7.2)*].
- Phenytoin: When a single oral 30 mg dose of another paroxetine product (4 times the recommended BRISDELLE dosage) was administered at phenytoin steady state (300 mg once daily for 14 days), paroxetine AUC and $T_{1/2}$ were reduced (by an average of 50% and 35%, respectively) compared to paroxetine administered alone.

In a separate study, when a single oral 300 mg dose of phenytoin was administered with another paroxetine product (30 mg once daily for 14 days) at paroxetine steady state, phenytoin AUC was slightly reduced (12% on average) compared to phenytoin administered alone. Because both drugs exhibit nonlinear pharmacokinetics, the above studies may not address the case where the two drugs are both being chronically dosed [see *Drug Interactions (7.2)*].

- **Digoxin:** A clinical drug interaction study showed that concurrent use of digoxin did not affect paroxetine exposure.
- **Diazepam:** A clinical drug interaction study showed that concurrent use of diazepam did not affect paroxetine exposure.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis

Two-year carcinogenicity studies were conducted in rodents given paroxetine in the diet at 1, 5, and 25 mg/kg/day (mice) and 1, 5, and 20 mg/kg/day (rats). The doses used in these carcinogenicity studies were approximately 16 (mouse) and 26 (rat) times the MHRD for BRISDELLE for the treatment of moderate to severe VMS associated with menopause. There was a significantly greater number of male rats in the high-dose group with reticulum cell sarcomas (1/100, 0/50, 0/50, and 4/50 for control, low-, middle-, and high-dose groups, respectively) and a significantly increased linear trend across groups for the occurrence of lymphoreticular tumors in male rats. Female rats were not affected. Although there was a dose-related increase in the number of tumors in mice, there was no paroxetine-related increase in the number of mice with tumors. The relevance of these findings to humans is unknown.

Mutagenesis

Paroxetine produced no genotoxic effect in a battery of 5 *in vitro* and 2 *in vivo* assays that included the following: bacterial mutation assay, mouse lymphoma mutation assay, unscheduled DNA synthesis assay, and tests for cytogenetic aberrations *in vivo* in mouse bone marrow and *in vitro* in human lymphocytes and in a dominant lethal test in rats.

14 CLINICAL STUDIES

The effectiveness of BRISDELLE as a treatment for moderate to severe vasomotor symptoms (VMS) associated with menopause was established in two Phase 3 randomized, double-blind, placebo-controlled clinical trials in 1,174 postmenopausal females. In these trials, patients must have had a minimum of 7-8 moderate to severe VMS per day at baseline (≥ 50 per week) for 30 days prior to receiving study drug. Patients were randomized to BRISDELLE 7.5 mg orally once daily or daily placebo:

- Trial 1 was a 12-week trial with a total of 606 postmenopausal females (average age 55 years; 65% White, 33% Black or African American, and 2% other races; 10.6% were Hispanic/Latina and 89.4% were not Hispanic/Latina; 18% surgically menopausal and 82% naturally menopausal).
- Trial 2 was a 24-week trial with a total of 568 postmenopausal females (average age

54 years; 76% White, 22% Black or African American, and 2% other races; 6.5% were Hispanic/Latina and 93.5% were not Hispanic/Latina; 20% surgically menopausal and 81% naturally menopausal).

The co-primary efficacy endpoints for both trials were the reduction from baseline in VMS frequency and VMS severity at Weeks 4 and 12.

- Data from Trial 1 showed a statistically significant reduction from baseline in the frequency of moderate to severe VMS at Week 4 and Week 12 and a statistically significant reduction in the severity of moderate to severe VMS at Week 4 in the BRISDELLE group compared to the placebo group (Table 4).
- Data from Trial 2 showed a statistically significant reduction from baseline in the frequency and severity of moderate to severe VMS at Week 4 and Week 12 in the BRISDELLE group compared to placebo group (Table 5).

Table 4: Trial 1: Changes in the Daily Frequency and Daily Severity of VMS at Weeks 4 and 12 in Postmenopausal Females with Moderate to Severe VMS (MITT Population)

	Frequency		Severity	
	BRISDELLE	Placebo	BRISDELLE	Placebo
Baseline				
n	301	305	301	305
Median	10.4	10.4	2.5	2.5
Change from baseline at Week 4				
n	289	293	281	289
Median	-4.3	-3.1	-0.05	0.00
Treatment Difference*	-1.2		-0.05	
P-value#	<0.01		<0.01	
Change from baseline at Week 12				
n	264	274	236	253
Median	-5.9	-5.0	-0.06	-0.02
Treatment Difference*	-0.9		-0.04	
P-value#	<0.01		0.17	

MITT population: all consented and randomized patients with valid baseline daily hot flash diary data who had taken at least 1 dose of study drug and had at least 1 day of on-treatment daily hot flash diary data.

* Treatment Difference: the difference between the median changes from baseline.

P-value was obtained from rank-ANCOVA model.

Table 5: Trial 2: Changes in the Daily Frequency and Daily Severity of VMS at Weeks 4 and 12 in Postmenopausal Females with Moderate to Severe VMS (MITT Population)

	Frequency		Severity	
	BRISDELLE	Placebo	BRISDELLE	Placebo
Baseline				
n	284	284	284	284
Median	9.9	9.6	2.5	2.5
Change from baseline at Week 4				
n	276	274	268	271
Median	-3.8	-2.5	-0.04	-0.01
Treatment Difference*	-1.3		-0.03	
P-value#	<0.01		0.04	
Change from baseline at Week 12				
n	257	244	245	236
Median	-5.6	-3.9	-0.05	0.00
Treatment Difference*	-1.7		-0.05	
P-value#	<0.01		<0.01	

MITT population: all consented and randomized patients with valid baseline daily hot flash diary data who had taken at least 1 dose of study drug and had at least 1 day of on-treatment daily hot flash diary data.

* Treatment Difference: the difference between the median changes from baseline.

P-value is obtained from rank-ANCOVA model.

Persistence of benefit at 24 weeks in Trial 2 was evaluated with a responder analysis where responders were defined as those patients who achieved $\geq 50\%$ reduction from baseline in the frequency of moderate to severe VMS at Week 24. The proportion of patients who achieved a $\geq 50\%$ reduction in the frequency of moderate to severe VMS from baseline to Week 24 was 48% in the BRISDELLE group and 36% in the placebo group at Week 24.

16 HOW SUPPLIED/STORAGE AND HANDLING

BRISDELLE (paroxetine) capsules is available as 7.5 mg pink capsules printed with black edible ink with "BRISDELLE" and "7.5 mg".

NDC 83107-027-30 blister packs of 30

Store at 20°-25°C (68°-77°F); excursions permitted to 15°-30°C (59°-86°F). Protect from light and humidity.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Medication Guide).

Instruct patients to read the Medication Guide before starting therapy with BRISDELLE and to reread it each time the prescription is renewed.

Suicidal Thoughts and Behaviors

Advise patients, their families, and their caregivers to look for the emergence of suicidal thoughts and behaviors, especially early during treatment and to alert the health care provider if such changes occur [*see Boxed Warning and Warnings and Precautions (5.1)*].

Serotonin Syndrome

Caution patients about the risk of serotonin syndrome, particularly with the concomitant use of BRISDELLE with triptans, tricyclic antidepressants, opioids, linezolid, tramadol, amphetamines, St. John's Wort, lithium, tryptophan supplements, other serotonergic agents, or antipsychotic drugs. Instruct patients to contact their health care provider, or report to the emergency room, should they experience signs or symptoms of serotonin syndrome [*see Warnings and Precautions (5.2) and Drug Interactions (7.3)*].

Potential Impact on Tamoxifen Efficacy

Caution patients that efficacy of tamoxifen may be reduced when administered concomitantly and counsel them about the likely benefit of paroxetine for treating VMS vs. the risk of possible decreased tamoxifen effectiveness [*see Warnings and Precautions (5.3)*].

Increased Risk of Bleeding

Caution patients about the concomitant use of BRISDELLE and NSAIDs, aspirin, warfarin, and other anticoagulants because combined use of drugs that interfere with serotonin reuptake has been associated with an increased risk of bleeding [*see Warnings and Precautions (5.4)*].

Angle-Closure Glaucoma

Advise patients that taking BRISDELLE can cause mild pupillary dilation, which in susceptible individuals, can lead to an episode of angle-closure glaucoma -existing [*See Warnings and Precautions (5.5)*].

Hyponatremia

Caution patients about the risk of hyponatremia, particularly elderly patients and those who are taking diuretics or are volume-depleted [*see Warnings and Precautions (5.6)*].

Bone Fracture

Inform patients that there is the possibility for an increased risk of fracture [*see Warnings and Precautions (5.7)*].

Screening Patients for Bipolar Disorder and Monitoring for Mania/Hypomania

Advise patients, their families, and their caregivers to observe for signs of activation of mania/hypomania [*see Warnings and Precautions (5.8)*].

Pregnancy

Advise patients to notify their physician if they become pregnant during therapy [*see Contraindications (4.5) and Use in Specific Populations (8.1)*].

Drug Interactions

Advise patients to inform their healthcare provider if they are taking, or plan to take, any prescription or over-the-counter drugs, including herbal supplements, because there is a potential for interaction with paroxetine [see *Drug Interactions (7)*].

Sexual Dysfunction

Advise patients that use of BRISDELLE may cause symptoms of sexual dysfunction in female patients. Inform patients that they should discuss any changes in sexual function and potential management strategies with their healthcare provider [see *Warnings and Precautions (5.10)*].

Allergic Reactions

Advise patients to seek medical attention if they develop an allergic reaction such as rash, hives, swelling, or difficulty breathing [see *Adverse Reactions (6.2)*].

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Manufactured for:
Legacy Pharma Inc.
George Town, Grand Cayman
KY1-9012



MEDICATION GUIDE

BRISDELLE® (bris-del)
(paroxetine)
capsules

Read this Medication Guide that comes with BRISDELLE before you start taking it and each time you get a refill. There may be new information. This information does not take the place of talking to your healthcare provider about your medical condition or treatment. Talk with your healthcare provider if there is something you do not understand or want to learn more about.

What is the most important information I should know about BRISDELLE?

BRISDELLE may cause serious side effects, including:

- **Increased risk of suicidal thoughts or actions.** BRISDELLE and related antidepressant medicines may increase suicidal thoughts or actions within the first few months of treatment.
 - Depression or other serious mental illnesses are the most important causes of suicidal thoughts or actions.
 - Watch for these changes and call your healthcare provider right away if you notice:
 - New or sudden changes in mood, behavior, actions, thoughts, or feelings, especially if severe.

- Pay close attention to any changes when BRISDELLE is started.
- Keep all follow-up visits with your healthcare provider and call between visits if you are worried about symptoms.

Call your healthcare provider right away or go to the nearest emergency room if you have any of the following symptoms, especially if they are new, worse, or worry you:

- attempts to commit suicide
- acting on dangerous impulses
- acting aggressive or violent
- thoughts about suicide or dying
- new or worse depression
- new or worse anxiety or panic attacks
- feeling agitated, restless, angry or irritable
- trouble sleeping
- an increase in activity or talking more than what is normal for you
- other unusual changes in behavior or mood

What is BRISDELLE?

- BRISDELLE is a prescription medicine used to treat moderate to severe hot flashes associated with menopause.
- BRISDELLE is not any for psychiatric problems such as depression, obsessive compulsive disorder, panic disorder, generalized anxiety disorder, social anxiety disorder, and post-traumatic stress disorder.
- BRISDELLE is not for use in children.

Who should not take BRISDELLE?

Do not take BRISDELLE if you:

- **take a Monoamine Oxidase Inhibitor (MAOI).** Ask your healthcare provider or pharmacist if you are not sure if you take an MAOI, including the antibiotic linezolid or the intravenous methylene blue.
 - Do not take an MAOI within 14 days of stopping BRISDELLE unless directed to do so by your healthcare provider.
 - Do not start BRISDELLE if you stopped taking an MAOI in the last 14 days unless directed to do so by your healthcare provider.
 - **People who take BRISDELLE close in time to an MAOI may have a serious or life-threatening side effect called serotonin syndrome. Get medical help right away if you have any of these symptoms:**
 - high fever
 - uncontrolled muscle spasms
 - stiff muscles
 - rapid changes in heart rate or blood pressure
 - confusion
 - loss of consciousness (pass out)
 - nausea, vomiting, or diarrhea
- **take thioridazine.** Do not take thioridazine together with BRISDELLE because this can cause serious heart rhythm problems or sudden death.

- **take the antipsychotic medicine pimozide.** Do not take pimozide together with BRISDELLE because this can cause serious heart problems.
- **are allergic to paroxetine or any of the ingredients in BRISDELLE. See the end of this Medication Guide for a complete list of ingredients in BRISDELLE.**
- **are pregnant or become pregnant.** BRISDELLE can harm your unborn baby. Call your healthcare provider if you become pregnant while taking BRISDELLE.

What should I tell my healthcare provider before taking BRISDELLE?

Before taking BRISDELLE, tell your healthcare provider about all of your medical conditions, including if you:

- have liver problems
- have kidney problems
- have or had seizures or convulsions
- have or have a family history of bipolar disorder, mania or hypomania
- have low sodium levels in your blood
- have or had bleeding problems
- have glaucoma (high pressure in the eye)
- have bone problems

Tell your healthcare provider about all the medicines that you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. BRISDELLE and some medicines may interact with each other, may not work as well, or may cause serious side effects when taken together.

If you take BRISDELLE, you should not take any other medicines that contain paroxetine, including Paxil, Paxil CR and Pexeva.

Especially tell your healthcare provider if you take:

- triptans used to treat migraine headache
- medicines used to treat mood, anxiety, psychotic or thought disorders, including MAOIs, SSRIs, tricyclics, lithium, buspirone, or antipsychotics
- tramadol, fentanyl, meperidine, methadone, or other opioids
- over-the-counter supplements such as tryptophan or St. John's Wort
- amphetamines
- thioridazine
- pimozide
- tamoxifen
- atomoxetine
- cimetidine
- digoxin
- theophylline
- medicines to treat irregular heart rate (like propafenone, flecainide, and encainide)
- medicines used to treat schizophrenia
- certain medicines used to treat HIV infection
- the blood thinner warfarin

- nonsteroidal anti-inflammatory drugs (NSAIDs) (like ibuprofen, naproxen, or aspirin)
- certain medicines used to treat seizures (like phenobarbital and phenytoin)
- other drugs containing paroxetine, the medicine in BRISDELLE

Ask your healthcare provider if you are not sure if you are taking any of these medications.

Your healthcare provider or pharmacist can tell you if it is safe to take BRISDELLE with your other medicines. Do not start or stop any medicine while taking BRISDELLE without talking to your healthcare provider first.

How should I take BRISDELLE?

- Take BRISDELLE exactly as your healthcare provider tells you to take it.
- Take BRISDELLE 1 time each day at bedtime.
- BRISDELLE may be taken with or without food.
- If you take too much BRISDELLE, call your healthcare provider or Poison Help line right away, or go to the nearest emergency room right away.

What are the possible side effects of BRISDELLE?

BRISDELLE may cause serious side effects, including:

- See “What is the most important information I should know about BRISDELLE?”
- **Serotonin syndrome. This condition can be life-threatening and can happen when you take BRISDELLE with certain other medicines. Call your healthcare provider or go to the nearest hospital emergency room right away if you have any of the following symptoms of serotonin syndrome:**
 - agitation (nervousness), hallucinations, coma or other changes in mental status
 - coordination problems or muscle twitching (small movements of the muscles that you cannot control)
 - racing heartbeat, high or low blood pressure
 - sweating or fever
 - nausea, vomiting, or diarrhea
 - muscle rigidity
 - dizziness
 - flushing
 - tremors
 - seizures
- **Reduced effectiveness of tamoxifen.** Tamoxifen (a medicine used to treat breast cancer) may not work as well if it is taken while you take BRISDELLE. If you are taking tamoxifen, tell your healthcare provider before starting BRISDELLE.
- **Abnormal bleeding.** BRISDELLE may increase your risk of bleeding or bruising, especially if you take the blood thinner warfarin or non-steroidal anti-inflammatory drugs (NSAIDs) like ibuprofen, naproxen, or aspirin.
- **Visual problems (angle-closure glaucoma).** BRISDELLE may cause a visual problem called angle-closure glaucoma. Symptoms may include:

- eye pain
- changes in vision
- swelling or redness in or around the eye

Only some people are at risk for these problems. You may want to undergo an eye examination to see if you are at risk and receive preventative treatment if you are.

- **Low salt (sodium) levels in the blood.** Elderly people and those taking diuretics or are dehydrated are at greater risk of experiencing low salt levels in their blood. Symptoms may include:
 - headache
 - weakness or feeling unsteady
 - confusion, problems concentrating or thinking or memory problems
 - seeing or hearing things that are not there (hallucinations)
 - fainting (syncope)
 - seizures
 - stopping breathing (respiratory arrest)
- **Bone fractures.** Women who take BRISDELLE may have a higher risk of bone fractures. Contact your healthcare provider if you have pain in a bone.
- **Manic episodes. Manic episodes may happen in people with bipolar disorder who take BRISDELLE. Symptoms may include:**
 - greatly increased energy
 - severe trouble sleeping
 - racing thoughts
 - reckless behavior
 - unusually grand ideas
 - excessive happiness or irritability
 - talking more or faster than usual
- **Seizures**
- **Sexual problems (dysfunction).** Taking selective serotonin reuptake inhibitors (SSRIs), including BRISDELLE, may cause symptoms of sexual problems. Symptoms in females may include:
 - decreased sexual drive
 - delayed or inability to have an orgasm

The most common side effects of BRISDELLE include:

- headache
- tiredness
- nausea and vomiting

Tell your healthcare provider if you have any side effect that bothers you or does not go away. These are not all the possible side effects of BRISDELLE. For more information, ask your healthcare provider or pharmacist.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088

How should I store BRISDELLE?

- Store BRISDELLE at room temperature between 68°F to 77°F (20°C to

25°C).

- Keep BRISDELLE out of the light.
- Keep BRISDELLE dry.

Keep BRISDELLE and all medicines out of the reach of children

General information about the safe and effective use of BRISDELLE.

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use BRISDELLE for a condition for which it was not prescribed. Do not give BRISDELLE to other people, even if they have the same condition. It may harm them. You may ask your healthcare provider or pharmacist for information about BRISDELLE that is written for healthcare professionals.

What are the ingredients in BRISDELLE?

Active ingredient: paroxetine

Inactive ingredients: dibasic calcium phosphate, sodium starch glycolate, magnesium stearate, gelatin, titanium dioxide, FD&C Yellow #6, FD&C Red #3, FD&C Red #40, shellac and black iron oxide.

Manufactured for:

Legacy Pharma Inc., George Town, Grand Cayman

BRISDELLE is a registered trademark of Legacy Pharma Inc.

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For more information about BRISDELLE go to www.BRISDELLE.com

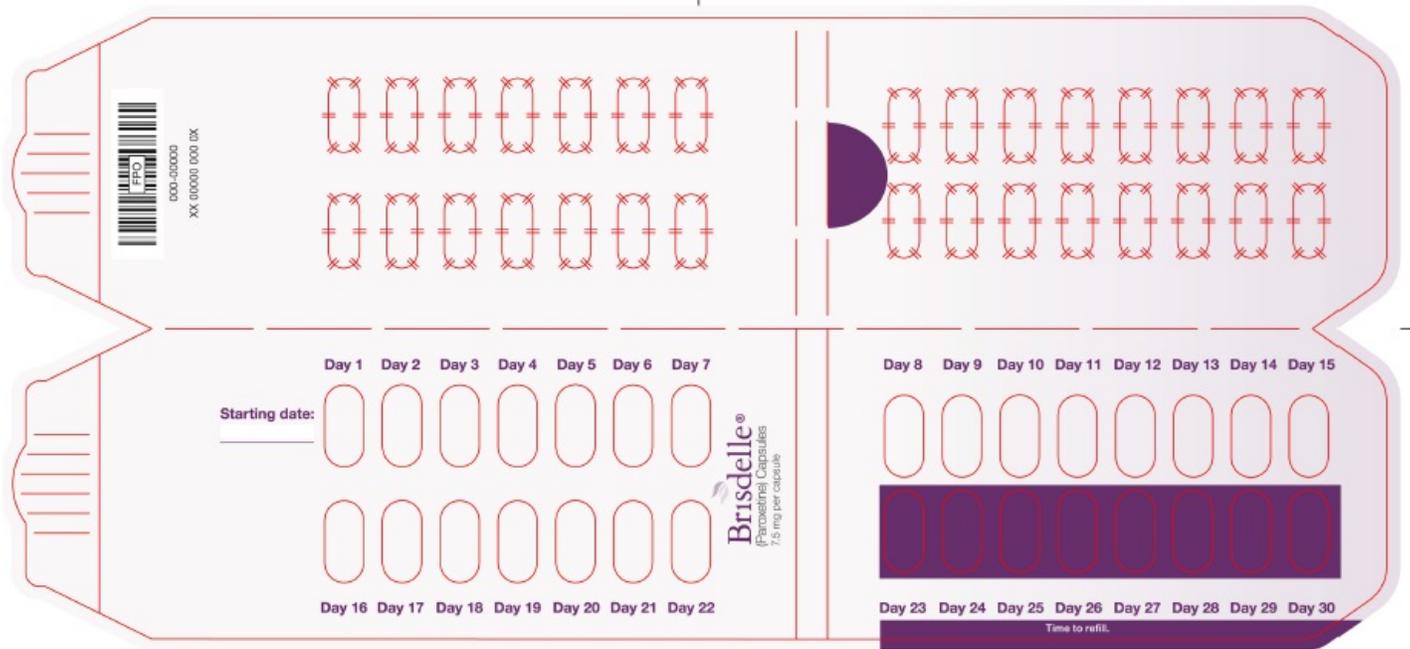
MDG90757001

This Medication Guide has been approved by the U.S. Food and Drug Administration
02/2025

Revised:

PRINCIPAL DISPLAY PANEL

BRISDELLE (paroxetine) Capsules is available as 7.5 mg - NDC 83107-027-30 - 30 Caps Blister Label



BRISDELLE (paroxetine) Capsules is available as 7.5 mg - NDC 83107-027-30 - 30 Caps Blister Carton

Brisdelle®

(Paroxetine) Capsules
7.5 mg per capsule



NDC 83107-027-30

Rx Only
30 capsules
For Oral Use

Find out more at
www.Brisdelle.com

Brisdelle®

(Paroxetine) Capsules
7.5 mg per capsule

Helpful Information:

- Days 1-30 are numbered to help you keep track of taking your medication daily as prescribed
- Write down your starting date on the provided line next to Day 1
- Take one Brisdelle® capsule daily at bedtime
 - Each capsule is the same, so don't worry if you take them out of sequence
- Refill your prescription during your last week or as advised by your doctor

Please read
full Prescribing
Information
with Medication
Guide inside. →

Each capsule contains the same amount of active ingredient.

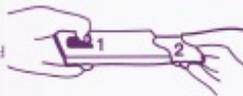
3. PRESS AND
HOLD HERE

2. PULL OUT HERE

CHILD-RESISTANT CONTAINER

OPENING INSTRUCTIONS:

1. Use thumb/nail to push the button gently
2. While holding button down, pull out the medication card
3. To close, fold the medication card and slide it back into the package to reengage child-resistant feature



Brisdelle® (Paroxetine) Capsules 7.5 mg per capsule

Manufactured for
LEGACY
Legacy Pharmaceuticals
George Town, Grand Cayman
KY1-8818
1-800-227-7151

Brisdelle®
(Paroxetine) Capsules
7.5 mg per capsule



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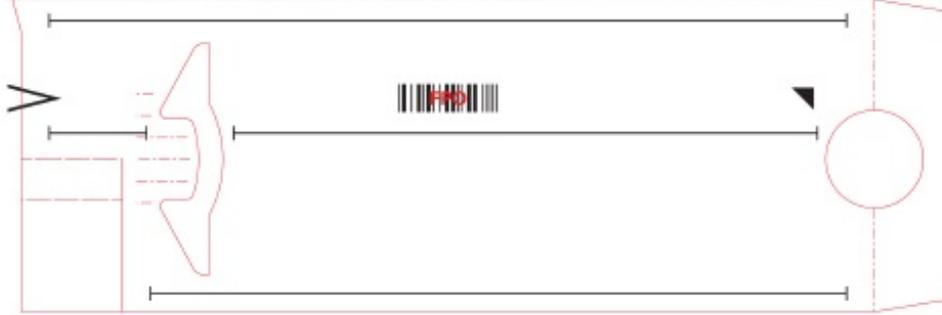
Store this medication in the child-resistant container in which it was dispensed. Keep this and all medications out of the reach of children. Store at 20°-25°C (68°-77°F), excursions permitted to 15°-30°C (59°-86°F) [See USP controlled room temperature]. Protect from light and humidity. Each capsule contains paroxetine mesylate equivalent to 7.5 mg of paroxetine.



Attention PHARMACIST
Each patient must receive the ordered Medication Guide.
To the PATIENT
Read the enclosed Medication Guide before using. Dosing: Take one capsule (7.5 mg) once a day, at bedtime, as prescribed.
by your doctor.

AFFIX PHARMACY LABEL HERE

Brisdelle® (Paroxetine) Capsules 7.5 mg per capsule



paroxetine capsule

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:83107-027
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PAROXETINE MESYLATE (UNII: M711N184JE) (PAROXETINE - UNII:41VRH5220H)	PAROXETINE	7.5 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
GELATIN (UNII: 2G86QN327L)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 3 (UNII: PN2ZH5LOQY)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SHELLAC (UNII: 46N107B71O)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	16mm
Flavor		Imprint Code	BRISDELLE;7;5;mg
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83107-027-30	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	02/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA204516	02/18/2025	

Labeler - Legacy Pharma USA, Inc. (118831776)

Establishment			
Name	Address	ID/FEI	Business Operations
Packaging Coordinators, LLC		078525133	pack(83107-027) , label(83107-027)

Establishment			
Name	Address	ID/FEI	Business Operations
Norwich Pharmaceuticals, Inc.		132218731	manufacture(83107-027) , analysis(83107-027) , pack(83107-027) , label(83107-027)

Establishment			
Name	Address	ID/FEI	Business Operations
Excella GmbH		329809800	api manufacture(83107-027) , analysis(83107-027) , manufacture(83107-027)

Revised: 4/2025

Legacy Pharma USA, Inc.