

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use **TOLTERODINE TARTRATE EXTENDED-RELEASE CAPSULES** safely and effectively. See full prescribing information for **TOLTERODINE TARTRATE EXTENDED-RELEASE CAPSULES**.

TOLTERODINE tartrate extended-release capsules, for oral use

Initial U.S. Approval: December 2000

RECENT MAJOR CHANGES

Table with 2 columns: Contraindications, Warnings and Precautions. Rows include: Hypersensitivity to fesoterodine fumarate (4) 09/2011; Angioedema (5.1) 09/2011; Central Nervous System Effects (5.5) 08/2012.

INDICATIONS AND USAGE

Tolterodine tartrate extended-release capsule is an antimuscarinic indicated for the treatment of overactive bladder with symptoms of urge urinary incontinence, urgency, and frequency. (1)

DOSAGE AND ADMINISTRATION

- 4 mg capsules taken orally once daily with water and swallowed whole. (2.1)
- 2 mg capsules taken orally once daily with water and swallowed whole in the presence of:
- mild to moderate hepatic impairment (Child-Pugh class A or B) (2.2)
- severe renal impairment (Creatinine Clearance (Cr) 10 to 30 mL/min) (2.2)
- drugs that are potent CYP3A4 inhibitors. (2.2)
- Tolterodine tartrate extended-release capsules are not recommended for use in patients with Cr <10 mL/min. (2.2)
- Tolterodine tartrate extended-release capsules are not recommended for use in patients with severe hepatic impairment (Child-Pugh Class C). (2.2)

DOSAGE FORMS AND STRENGTHS

Capsules: 2 mg and 4 mg (3)

CONTRAINDICATIONS

Tolterodine tartrate extended-release capsules are contraindicated in patients with urinary retention, gastric retention, or uncontrolled narrow-angle glaucoma. Tolterodine tartrate extended-release capsules are also contraindicated in patients with known hypersensitivity to the drug or its ingredients, or to fesoterodine fumarate extended-release tablets which, like tolterodine tartrate extended-release capsules, are metabolized to 5-hydroxymethyl tolterodine. (4)

WARNINGS AND PRECAUTIONS

- Anaphylaxis and angioedema requiring hospitalization and emergency medical treatment have occurred with the first or subsequent doses of tolterodine tartrate extended-release capsules. (5.1)
- Urinary Retention: use caution in patients with clinically significant bladder outflow obstruction because of the risk of urinary retention. (5.2)

- Gastrointestinal Disorders: use caution in patients with gastrointestinal obstructive disorders or decreased gastrointestinal motility because of the risk of gastric retention. (5.3)
- Controlled Narrow-Angle Glaucoma: use caution in patients being treated for narrow-angle glaucoma. (5.4)
- Central Nervous System Effects: Somnolence has been reported with tolterodine tartrate extended-release capsules. Advise patients not to drive or operate heavy machinery until they know how tolterodine tartrate extended-release capsules affects them (5.5).
- Myasthenia Gravis: use caution in patients with myasthenia gravis. (5.8)
- QT Prolongation: consider observations from the thorough QT study in clinical decisions to prescribe tolterodine tartrate extended-release capsules to patients with a known history of QT prolongation or to patients who are taking Class IA (e.g., quinidine, procainamide) or Class III (e.g., amiodarone, sotalol) antiarrhythmic medications. (5.9)

ADVERSE REACTIONS

The most common adverse reactions (incidence ≥4% and >placebo) were dry mouth, headache, constipation, and abdominal pain. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Hetero Labs Limited at 1-866-495-1995 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Potent CYP3A4 Inhibitors: Coadministration may increase systemic exposure to tolterodine tartrate extended-release capsules. Reduce tolterodine tartrate extended-release capsules dose to 2 mg once daily. (7.2)
- Other Anticholinergics (antimuscarinics): Concomitant use with other anticholinergic agents may increase the frequency and/or severity of dry mouth, constipation, blurred vision, and other anticholinergic pharmacological effects. (7.6)

USE IN SPECIFIC POPULATIONS

- Pregnancy and Lactation: Tolterodine tartrate extended-release capsules should be used during pregnancy only if the potential benefit for the mother justifies the potential risk to the fetus. Tolterodine tartrate extended-release capsules should not be administered during nursing. (8.1, 8.3)
- Pediatric Use: Efficacy in the pediatric population has not been demonstrated. Safety information from a study of a total of 710 pediatric patients (486 on tolterodine tartrate extended-release capsules, 224 on placebo) is available. (8.4)
- Renal Impairment: Tolterodine tartrate extended-release capsules are not recommended for use in patients with Cr <10 mL/min. Dose adjustment in severe renal impairment (Cr: 10 to 30 mL/min). (8.6)
- Hepatic Impairment: Not recommended for use in severe hepatic impairment (Child Pugh Class C). Dose adjustment in mild to moderate hepatic impairment (Child Pugh Class A, B). (8.7)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling. Revised: 09/2016

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

Tolterodine tartrate extended-release capsules are indicated for the treatment of overactive bladder with symptoms of urge urinary incontinence, urgency, and frequency [see CLINICAL STUDIES (14)].

2 DOSAGE AND ADMINISTRATION

2.1 Dosing Information

The recommended dose of tolterodine tartrate extended-release capsules is 4 mg once daily with water and swallowed whole. The dose may be lowered to 2 mg daily based on individual response and tolerability; however, limited efficacy data are available for tolterodine tartrate extended-release capsules 2 mg [see CLINICAL STUDIES (14)].

2.2 Dosage Adjustment in Specific Populations

For patients with mild to moderate hepatic impairment (Child-Pugh Class A or B) or severe renal impairment (Cr: 10 to 30 mL/min), the recommended dose of tolterodine tartrate extended-release capsules is 2 mg once daily. Tolterodine tartrate extended-release capsules are not recommended for use in patients with severe hepatic impairment (Child-Pugh Class C). Patients with Cr <10 mL/min have not been studied and use of tolterodine tartrate extended-release capsules in this population is not recommended [see WARNINGS AND PRECAUTIONS (5.6) and USE IN SPECIFIC POPULATIONS (8.6, 8.7)].

2.3 Dosage Adjustment in Presence of Concomitant Drugs

For patients who are taking drugs that are potent inhibitors of CYP3A4 [e.g., ketoconazole, clarithromycin, or ritonavir], the recommended dose of tolterodine tartrate extended-release capsules is 2 mg once daily [see DRUG INTERACTIONS (7.2)].

3 DOSAGE FORMS AND STRENGTHS

The 2 mg capsules are blue green colored size '4' hard gelatin capsules imprinted with 'J' on cap and '104' on body, filled with white to off-white pellets.

The 4 mg capsules are blue colored size '3' hard gelatin capsules imprinted with 'J' on cap and '105' on body, filled with white to off-white pellets.

4 CONTRAINDICATIONS

Tolterodine tartrate extended-release capsules are contraindicated in patients with urinary retention, gastric retention, or uncontrolled narrow-angle glaucoma. Tolterodine tartrate extended-release capsules are also contraindicated in patients with known hypersensitivity to the drug or its ingredients, or to fesoterodine fumarate extended-release tablets which, like tolterodine tartrate extended-release capsules, are metabolized to 5-hydroxymethyl tolterodine [see WARNINGS AND PRECAUTIONS (5.2) (5.3), (5.4)].

5 WARNINGS AND PRECAUTIONS

5.1 Angioedema

Anaphylaxis and angioedema requiring hospitalization and emergency medical treatment have occurred with the first or subsequent doses of tolterodine tartrate extended-release capsules. In the event of difficulty in breathing, upper airway obstruction, or fall in blood pressure, tolterodine tartrate extended-release capsules should be discontinued and appropriate therapy promptly provided.

5.2 Urinary Retention

Administer tolterodine tartrate extended-release capsules with caution to patients with clinically significant bladder outflow obstruction because of the risk of urinary retention [see CONTRAINDICATIONS (4)].

5.3 Gastrointestinal Disorders

Administer tolterodine tartrate extended-release capsules with caution in patients with gastrointestinal obstructive disorders because of the risk of gastric retention.

Tolterodine tartrate extended-release capsules, like other antimuscarinic drugs, may decrease gastrointestinal motility and should be used with caution in patients with conditions associated with decreased gastrointestinal motility (e.g., intestinal atony) [see CONTRAINDICATIONS (4)].

5.4 Controlled Narrow-Angle Glaucoma

Administer tolterodine tartrate extended-release capsules with caution in patients being treated for narrow-angle glaucoma [see CONTRAINDICATIONS (4)].

5.5 Central Nervous System Effects

Tolterodine tartrate extended-release capsules are associated with anticholinergic central nervous system (CNS) effects [see Adverse Reactions (6.2)] including dizziness and somnolence [see Adverse Reactions (6.1)]. Patients should be monitored for signs of anticholinergic CNS effects, particularly after beginning treatment or increasing the dose. Advise patients not to drive or operate heavy machinery until the drug's effects have been determined. If a patient experiences anticholinergic CNS effects, dose reduction or drug discontinuation should be considered.

5.6 Hepatic Impairment

The clearance of orally administered tolterodine immediate release was substantially lower in cirrhotic patients than in the healthy volunteers. For patients with mild to moderate hepatic impairment (Child-Pugh Class A or B), the recommended dose of tolterodine tartrate extended-release capsules are 2 mg once daily. Tolterodine tartrate extended-release capsules is not recommended for use in patients with severe hepatic impairment (Child-Pugh Class C) [see DOSAGE AND ADMINISTRATION (2.2) and USE IN SPECIFIC POPULATIONS (8.6)].

5.7 Renal Impairment

Renal impairment can significantly alter the disposition of tolterodine and its metabolites. The dose of tolterodine tartrate extended-release capsules should be reduced to 2 mg once daily in patients with severe renal impairment (Cr: 10 to 30 mL/min). Patients with Cr <10 mL/min have not been studied and use of tolterodine tartrate extended-release capsules in this population is not recommended [see DOSAGE AND ADMINISTRATION (2.2) and USE IN SPECIFIC POPULATIONS (8.7)].

5.8 Myasthenia Gravis

Administer tolterodine tartrate extended-release capsules with caution in patients with myasthenia gravis, a disease characterized by decreased cholinergic activity at the neuromuscular junction.

5.9 Use in Patients with Congenital or Acquired QT Prolongation

In a study of the effect of tolterodine immediate release tablets on the QT interval [see CLINICAL PHARMACOLOGY (12.2)], the effect on the QT interval appeared greater for 8 mg/day (two times the therapeutic dose) compared to 4 mg/day and was more pronounced in CYP2D6 poor metabolizers (PM) than extensive metabolizers (EMs). The effect of tolterodine 8 mg/day was not as large as that observed after four days of therapeutic dosing with the active control moxifloxacin. However, the confidence intervals overlapped.

These observations should be considered in clinical decisions to prescribe tolterodine tartrate extended-release capsules to patients with a known history of QT prolongation or to patients who are taking Class IA (e.g., quinidine, procainamide) or Class III (e.g., amiodarone, sotalol) antiarrhythmic medications. There has been no association of Torsade de Pointes in the international post-marketing experience with tolterodine tartrate immediate release tablets or tolterodine tartrate extended-release capsules.

6 ADVERSE REACTIONS

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

17.2 FDA-Approved Patient Labeling

PATIENT INFORMATION
Tolterodine Tartrate Extended-Release Capsules (TOH-TER-oh-deen TAR-trate)
Read the Patient Information that comes with tolterodine tartrate extended-release capsules before you start using it and each time you get a refill. There may be new information. This leaflet does not take the place of talking with your doctor about your condition or your treatment. Only your doctor can determine if treatment with tolterodine tartrate extended-release capsules is right for you.
What are tolterodine tartrate extended-release capsules?
Tolterodine tartrate extended-release capsule is a prescription medicine for adults used to treat the following symptoms due to a condition called overactive bladder.
• Having a strong need to urinate with leaking or wetting accidents (urge urinary incontinence).
• Having a strong need to urinate right away (urgency).
• Having to urinate often (frequency).
• The symptoms of overactive bladder when studied in children.
What is overactive bladder?
Overactive bladder happens when you cannot control your bladder muscle. When the muscle contracts too often or too strongly, you may feel the need to urinate frequently, which are leakage of urine (urge urinary incontinence), needing to urinate right away (urgency), and needing to urinate often (frequency).
Who should not take tolterodine tartrate extended-release capsules?
Do not take tolterodine tartrate extended-release capsules if:
• You have trouble emptying your bladder (also called "urinary retention").
• Your stomach empties slowly (also called "gastric retention").
• You have an eye problem called "uncontrolled narrow-angle glaucoma".
• You are allergic to tolterodine tartrate extended-release capsules or to any of its ingredients. See the end of this leaflet for a complete list of ingredients.
• You are allergic to fesoterodine fumarate extended-release tablets, which contains fesoterodine tartrate extended-release capsules?
Before starting tolterodine tartrate extended-release capsules, tell your doctor about all of your medical conditions, including if you:
• Have any stomach or intestinal problems.
• Have trouble emptying your bladder or you have a weak urine stream.
• Have an eye problem called narrow-angle glaucoma.
• Have liver problems.
• Have kidney problems.
• Have a condition called myasthenia gravis.
• Or any family members have a rare heart condition called QT prolongation (long QT syndrome).
Are pregnant or trying to become pregnant. It is not known if tolterodine tartrate extended-release capsules could harm your unborn baby.
Are breastfeeding. It is not known if tolterodine tartrate passes into your milk and if it can harm your child.
Tell your doctor about all the medicines you take, including prescription and non-prescription medicines, vitamins, and herbal supplements. Other drugs can affect how your body handles tolterodine tartrate extended-release capsules. Your doctor may use a lower dose of tolterodine tartrate extended-release capsules if you are taking.
Certain medicines for fungus or yeast infections such as Nizoral (ketoconazole), Sporonox (itraconazole), or Monistat (miconazole).
Certain medicines for bacteria infections such as Biaxin (clarithromycin).
Certain medicines for treatment of HIV infection such as Norvir (ritonavir), Invirase (saquinavir), Reyataz (atazanavir), and Sandimmune (cyclosporine) or Velban (zalcitabine).
Know the medicines you take. Keep a list of them with you to show your doctor or pharmacist each time you get a new medicine.
How should I take tolterodine tartrate extended-release capsules?
Take tolterodine tartrate extended-release capsules exactly as prescribed. Your doctor will prescribe the dose that is right for you. Do not change your dose unless told to do so by your doctor.

Size : 250 x 450 mm
Book Folding : 35 x 35 mm
Color : Black
Pharma Code : F-4311, B-4312
Spec: Printed on 40 GSM Bible paper, front & back side printing.
Note: Pharma code position, Orientation & Product Name Size are tentative, will be change based on folding size



8.5 Geriatric Use

No overall differences in safety were observed between the older and younger patients treated with tolterodine.

In multiple-dose studies in which tolterodine immediate release 4 mg (2 mg bid) was administered, serum concentrations of tolterodine and of 5-HMT were similar in healthy elderly volunteers (aged 64 through 80 years) and healthy young volunteers (aged less than 40 years). In another clinical study, elderly volunteers (aged 71 through 81 years) were given tolterodine immediate release 2 or 4 mg (1 or 2 mg bid). Mean serum concentrations of tolterodine and 5-HMT in these elderly volunteers were approximately 20% and 50% higher, respectively, than concentrations reported in young healthy volunteers. However, no overall differences were observed in safety between older and younger patients on tolterodine in the Phase 3, 12-week, controlled clinical studies; therefore, no tolterodine dosage adjustment for elderly patients is recommended.

8.6 Renal Impairment

Renal impairment can significantly alter the disposition of tolterodine immediate release and its metabolites. In a study conducted in patients with creatinine clearance between 10 and 30 mL/min, tolterodine and 5-HMT levels were approximately 2 to 3 fold higher in patients with renal impairment than in healthy volunteers. Exposure levels of other metabolites of tolterodine (e.g., tolterodine acid, *N*-dealkylated tolterodine acid, *N*-dealkylated tolterodine, and *N*-dealkylated hydroxy tolterodine) were significantly higher (10 to 30 fold) in renally impaired patients as compared to the healthy volunteers. The recommended dose for patients with severe renal impairment (CrCl: 10 to 30 mL/min) is tolterodine tartrate extended-release capsules 2 mg daily. Patients with CrCl < 10 mL/min have not been studied and use of tolterodine tartrate extended-release capsules in this population is not recommended. [see **DOSE AND ADMINISTRATION** (2.2) and **WARNINGS AND PRECAUTIONS** (5.6)]. Tolterodine tartrate extended-release capsules have not been studied in patients with mild to moderate renal impairment (CrCl 30 to 80 mL/min).

8.7 Hepatic Impairment

Liver impairment can significantly alter the disposition of tolterodine immediate release. In a study of tolterodine immediate release conducted in cirrhotic patients (Child-Pugh Class A and B), the elimination half-life of tolterodine immediate release was longer in cirrhotic patients (mean, 7.8 hours) than in healthy, young, and elderly volunteers (mean, 2 to 4 hours). The clearance of orally administered tolterodine immediate release was substantially lower in cirrhotic patients (1 ± 1.7 L/h/kg) than in the healthy volunteers (5.7 ± 3.8 L/h/kg). The recommended dose for patients with mild to moderate hepatic impairment (Child-Pugh Class A or B) is tolterodine tartrate extended-release capsules 2 mg once daily. Tolterodine tartrate extended-release capsules are not recommended for use in patients with severe hepatic impairment (Child-Pugh Class C) [see **DOSE AND ADMINISTRATION** (2.2) and **WARNINGS AND PRECAUTIONS** (5.4)].

8.8 Gender

The pharmacokinetics of tolterodine immediate release and 5-HMT are not influenced by gender. Mean C_{max} of tolterodine immediate release (1.6 mcg/L in males versus 2.2 mcg/L in females) and the active 5-HMT (2.2 mcg/L in males versus 2.5 mcg/L in females) are similar in males and females who were administered tolterodine immediate release 2 mg. Mean AUC values of tolterodine (6.7 mcg•h/L in males versus 7.8 mcg•h/L in females) and 5-HMT (10 mcg•h/L in males versus 11 mcg•h/L in females) are also similar. The elimination half-life of tolterodine immediate release for both males and females is 2.4 hours, and the half-life of 5-HMT is 3 hours in females and 3.3 hours in males.

8.9 Race

Pharmacokinetic differences due to race have not been established.

10 OVERDOSAGE

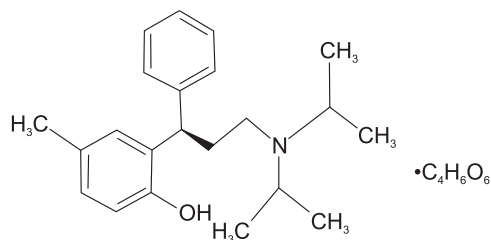
Overdosage with tolterodine tartrate extended-release capsules can potentially result in severe central anticholinergic effects and should be treated accordingly.

ECG monitoring is recommended in the event of overdosage. In dogs, changes in the QT interval (slight prolongation of 10% to 20%) were observed at a suprapharmacologic dose of 4.5 mg/kg, which is about 68 times higher than the recommended human dose. In clinical trials of normal volunteers and patients, QT interval prolongation was observed with tolterodine immediate release at doses up to 8 mg (4 mg bid) and higher doses were not evaluated [see **WARNINGS AND PRECAUTIONS** (5.9) and **CLINICAL PHARMACOLOGY** (12.2)].

A 27-month-old child who ingested 5 to 7 tolterodine immediate release 2 mg tablets was treated with a suspension of activated charcoal and was hospitalized overnight with symptoms of dry mouth. The child fully recovered.

11 DESCRIPTION

Tolterodine tartrate extended-release capsules contain tolterodine tartrate. The active moiety, tolterodine, is a muscarinic receptor antagonist. The chemical name of tolterodine tartrate is (R)-2-[3-[Bis (1-methylethyl) amino]-1-phenylpropyl]-4-Methyl phenol [R-(R*, R*)]-2, 3-dihydroxybutanedioate (1:1) (salt). The empirical formula of tolterodine tartrate is C₂₇H₃₇NO₆•C₄H₈O₆. Its structure is:



Tolterodine tartrate is a white to off white crystalline powder with a molecular weight of 475.57. The pK_a value is 9.24 and the solubility in water is 12 mg/mL. It is soluble in dimethylformamide and in methanol. The partition coefficient (Log P) value is 2.6.

Tolterodine tartrate extended-release capsules, 2 mg and 4 mg for oral administration contain 2 mg and 4 mg of tolterodine tartrate, respectively. Inactive ingredients are aqueous ethylcellulose dispersion, dehydrated alcohol, dibasic calcium phosphate anhydrous, ethyl acrylate and methyl methacrylate copolymer dispersion, ethyl cellulose, FD & C Blue 2, gelatin, hydroxypropyl methylcellulose, microcrystalline cellulose, talc and titanium dioxide. Additionally 2 mg capsules contain iron oxide yellow.

Both the 2 mg and 4 mg strengths capsule shells are imprinted with a pharmaceutical grade printing ink that contains butyl alcohol, dehydrated alcohol, isopropyl alcohol, potassium hydroxide, propylene glycol, shellac, strong ammonia solution and titanium dioxide.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Tolterodine acts as a competitive antagonist of acetylcholine at postganglionic muscarinic receptors. Both urinary bladder contraction and salivation are mediated via cholinergic muscarinic receptors.

After oral administration, tolterodine is metabolized in the liver, resulting in the formation of 5-hydroxymethyl tolterodine (5-HMT), the major pharmacologically active metabolite. 5-HMT, which exhibits an antimuscarinic activity similar to that of tolterodine, contributes significantly to the therapeutic effect. Both tolterodine and 5-HMT exhibit a high specificity for muscarinic receptors, since both show negligible activity or affinity for other neurotransmitter receptors and other potential cellular targets, such as calcium channels.

12.2 Pharmacodynamics

Tolterodine has a pronounced effect on bladder function. Effects on urodynamic parameters before and 1 and 5 hours after a single 6.4 mg dose of tolterodine immediate release were determined in healthy volunteers. The main effects of tolterodine at 1 and 5 hours were an increase in residual urine, reflecting an incomplete emptying of the bladder, and a decrease in detrusor pressure. These findings are consistent with an antimuscarinic action on the lower urinary tract.

Cardiac Electrophysiology

The effect of 2 mg BID and 4 mg BID of tolterodine tartrate immediate release tablets on the QT interval was evaluated in a 4-way crossover, double-blind, placebo- and active-controlled (moxifloxacin 400 mg QD) study in healthy male (N=25) and female (N=23) volunteers aged 18 to 55 years. Study subjects [approximately equal representation of CYP2D6 extensive metabolizers (EMs) and poor metabolizers (PMs)] completed sequential 4-day periods of dosing with moxifloxacin 400 mg QD, tolterodine 2 mg BID, tolterodine 4 mg BID, and placebo. The 4 mg BID dose of tolterodine IR (two times the highest recommended dose) was chosen because this dose results in tolterodine exposure similar to that observed upon coadministration of tolterodine 2 mg BID with potent CYP3A4 inhibitors in patients who are CYP2D6 poor metabolizers [see **DRUG INTERACTIONS** (7.2)]. QT interval was measured over a 12-hour period following dosing, including the time of peak plasma concentration (T_{max}) of tolterodine and at steady state (Day 4 of dosing).

Table 2 summarizes the mean change from baseline to steady state in corrected QT interval (QT_c) relative to placebo at the time of peak tolterodine (1 hour) and moxifloxacin (2 hour) concentrations. Both Fridericia's (QT_{cF}) and a population-specific (QT_{cP}) method were used to correct QT interval for heart rate. No single QT correction method is known to be more valid than others. QT interval was measured manually and by machine, and data from both are presented. The mean increase of heart rate associated with a 4 mg/day dose of tolterodine in this study was 2 beats/minute and 6.3 beats/minute with 8 mg/day tolterodine. The change in heart rate with moxifloxacin was 0.5 beats/minute.

Table 2. Mean (CI) change in QT_c from baseline to steady state (Day 4 of dosing) at T_{max} (relative to placebo)

Drug/Dose	N	QT _{cF} (msec) (manual)	QT _{cF} (msec) (machine)	QT _{cP} (msec) (manual)	QT _{cP} (msec) (machine)
Tolterodine 2 mg BID*	48	5.01 (0.28, 9.74)	1.16 (-2.29, 5.30)	4.45 (-0.37, 9.26)	2 (-1.81, 5.81)
Tolterodine 4 mg BID*	48	11.84 (7.11, 16.58)	5.63 (1.48, 9.77)	10.31 (5.49, 15.12)	8.34 (4.53, 12.15)
Moxifloxacin 400 mg QD†	45	19.26‡ (15.49, 23.03)	8.90 (4.77, 13.03)	19.10‡ (15.32, 22.89)	9.29 (5.34, 13.24)

* At T_{max} of 1 hr; 95% Confidence Interval.

† At T_{max} of 2 hr; 90% Confidence Interval.

‡ The effect on QT interval with 4 days of moxifloxacin dosing in this QT trial may be greater than typically observed in QT trials of other drugs.

The reason for the difference between machine and manual read of QT interval is unclear.

The QT effect of tolterodine immediate release tablets appeared greater for 8 mg/day (two times the therapeutic dose) compared to 4 mg/day. The effect of tolterodine 8 mg/day was not as large as that observed after four days of therapeutic dosing with the active control moxifloxacin. However, the confidence intervals overlapped.

Tolterodine's effect on QT interval was found to correlate with plasma concentration of tolterodine. There appeared to be a greater QT_c interval increase in CYP2D6 poor metabolizers than in CYP2D6 extensive metabolizers after tolterodine treatment in this study.

This study was not designed to make direct statistical comparisons between drugs or dose levels. There has been no association of Torsade de Pointes in the international post-marketing experience with tolterodine tartrate immediate release tablets or tolterodine tartrate extended-release capsules [see **WARNINGS AND PRECAUTIONS** (5.7)].

12.3 Pharmacokinetics

Absorption: In a study with ¹⁴C-tolterodine solution in healthy volunteers who received a 5 mg oral dose, at least 77% of the radiolabeled dose was absorbed. C_{max} and area under the concentration-time curve (AUC) determined after dosing of tolterodine immediate release are dose-proportional over the range of 1 to 4 mg. Based on the sum of unbound serum concentrations of tolterodine and 5-HMT ("active moiety"), the AUC of tolterodine extended-release 4 mg daily is equivalent to tolterodine immediate release 4 mg (2 mg bid). C_{max} and C_{min} levels of tolterodine extended release are about 75% and 150% of tolterodine immediate release, respectively. Maximum serum concentrations of tolterodine extended release are observed 2 to 6 hours after dose administration.

Effect of Food: There is no effect of food on the pharmacokinetics of tolterodine extended release.

Distribution: Tolterodine is highly bound to plasma proteins, primarily α₁-acid glycoprotein. Unbound concentrations of tolterodine average 3.7% ± 0.13% over the concentration range achieved in clinical studies. 5-HMT is not extensively protein bound, with unbound fraction concentrations averaging 36% ± 4%. The blood to serum ratio of tolterodine and 5-HMT averages 0.6 and 0.8, respectively, indicating that these compounds do not distribute extensively into erythrocytes. The volume of distribution of tolterodine following administration of a 1.28 mg intravenous dose is 113 ± 26.7 L.

Metabolism: Tolterodine is extensively metabolized by the liver following oral dosing. The primary metabolic route involves the oxidation of the 5-methyl group and is mediated by the cytochrome P450 2D6 (CYP2D6) and leads to the formation of a pharmacologically active metabolite, 5-HMT. Further metabolism leads to formation of the 5-carboxylic acid and *N*-dealkylated 5-carboxylic acid metabolites, which account for 51% ± 14% and 29% ± 6.3% of the metabolites recovered in the urine, respectively.

Variability in Metabolism: A subset of individuals (approximately 7% of Caucasians and approximately 2% of African Americans) are poor metabolizers for CYP2D6, the enzyme responsible for the formation of 5-HMT from tolterodine. The identified pathway of metabolism for these individuals ("poor metabolizers") is dealkylation via cytochrome P450 3A4 (CYP3A4) to *N*-dealkylated tolterodine. The remainder of the population is referred to as "extensive metabolizers." Pharmacokinetic studies revealed that tolterodine is metabolized at a slower rate in poor metabolizers than in extensive metabolizers; this results in significantly higher serum concentrations of tolterodine and in negligible concentrations of 5-HMT.

Excretion: Following administration of a 5 mg oral dose of ¹⁴C-tolterodine solution to healthy volunteers, 77% of radioactivity was recovered in urine and 17% was recovered in feces in 7 days. Less than 1% (< 2.5% in poor metabolizers) of the dose was recovered as intact tolterodine, and 5% to 14% (< 1% in poor metabolizers) was recovered as 5-HMT.

A summary of mean (± standard deviation) pharmacokinetic parameters of tolterodine extended release and 5-HMT in extensive (EM) and poor (PM) metabolizers is provided in Table 3. These data were obtained following single and multiple doses of tolterodine extended release administered daily to 17 healthy male volunteers (13 EM, 4 PM).

Table 3. Summary of Mean (±SD) Pharmacokinetic Parameters of Tolterodine Extended Release and its Active Metabolite (5-Hydroxymethyl Tolterodine) in Healthy Volunteers

	Tolterodine				5-Hydroxymethyl Tolterodine			
	t _{max} * (h)	C _{max} (mcg/L)	C _{avg} (mcg/L)	t _{1/2} (h)	t _{max} * (h)	C _{max} (mcg/L)	C _{avg} (mcg/L)	t _{1/2} (h)
Single dose								
4 mg†								
EM	4 (2-6)	1.3 (0.8)	0.8 (0.57)	8.4 (3.2)	4 (3-6)	1.6 (0.5)	1 (0.32)	8.8 (5.9)
Multiple dose								
4 mg								
EM	4 (2-6)	3.4 (4.9)	1.7 (2.8)	6.9 (3.5)	4 (2-6)	2.7 (0.90)	1.4 (0.6)	9.9 (4)
PM	4 (3-6)	19 (16)	13 (11)	18 (16)	‡	‡	‡	‡

C_{max} = Maximum serum concentration; t_{max} = Time of occurrence of C_{max};

C_{avg} = Average serum concentration; t_{1/2} = Terminal elimination half-life.

* Data presented as median (range).

† Parameter dose-normalized from 8 to 4 mg for the single-dose data.

‡ = not applicable.

Drug Interactions:

Potent CYP2D6 inhibitors: Fluoxetine is a selective serotonin reuptake inhibitor and a potent inhibitor of CYP2D6 activity. In a study to assess the effect of fluoxetine on the pharmacokinetics of tolterodine immediate release and its metabolites, it was observed that fluoxetine significantly inhibited the metabolism of tolterodine immediate release in extensive metabolizers, resulting in a 4.8-fold increase in tolterodine AUC. There was a 52% decrease in C_{max} and a 20% decrease in AUC of 5-hydroxymethyl tolterodine (5-HMT), the pharmacologically active metabolite of tolterodine. Fluoxetine thus alters the pharmacokinetics in patients who would otherwise be CYP2D6 extensive metabolizers of tolterodine immediate release to resemble the pharmacokinetic profile in poor metabolizers. The sums of unbound serum concentrations of tolterodine immediate release and 5-HMT are only 25% higher during the interaction. No dose adjustment is required when tolterodine and fluoxetine are co-administered.

Potent CYP3A4 inhibitors: The effect of a 200 mg daily dose of ketoconazole on the pharmacokinetics of tolterodine immediate release was studied in 8 healthy volunteers, all of whom were CYP2D6 poor metabolizers. In the presence of ketoconazole, the mean C_{max} and AUC of tolterodine increased by 2- and 2.5-fold, respectively. Based on these findings, other potent CYP3A4 inhibitors may also lead to increases of tolterodine plasma concentrations.

For patients receiving ketoconazole or other potent CYP3A4 inhibitors such as itraconazole, miconazole, clarithromycin, ritonavir, the recommended dose of tolterodine tartrate extended-release capsules is 2 mg daily [see **DOSE AND ADMINISTRATION** (2.3)].

Warfarin: In healthy volunteers, coadministration of tolterodine immediate release 4 mg (2 mg bid) for 7 days and a single dose of warfarin 25 mg on day 4 had no effect on prothrombin time, Factor VII suppression, or on the pharmacokinetics of warfarin.

Oral Contraceptives: Tolterodine immediate release 4 mg (2 mg bid) had no effect on the pharmacokinetics of an oral contraceptive (ethinyl estradiol 30 mcg/levonorgestrel 150 mcg) as evidenced by the monitoring of ethinyl estradiol and levonorgestrel over a 2-month period in healthy female volunteers.

Diuretics: Coadministration of tolterodine immediate release up to 8 mg (4 mg bid) for up to 12 weeks with diuretic agents, such as indapamide, hydrochlorothiazide, triamterene, bendroflumethiazide, chlorothiazide, methichlorothiazide, or furosemide, did not cause any adverse electrocardiographic (ECG) effects.

Effect of tolterodine on other drugs metabolized by Cytochrome P450 enzymes: Tolterodine immediate release does not cause clinically significant interactions with other drugs metabolized by the major drug-metabolizing CYP enzymes. *In vivo* drug-interaction data show that tolterodine immediate release does not result in clinically relevant inhibition of CYP1A2, 2D6, 2C9, 2C19, or 3A4 as evidenced by lack of influence on the marker drugs caffeine, debrisoquine, S-warfarin, and omeprazole. *In vitro* data show that tolterodine immediate release is a competitive inhibitor of CYP2D6 at high concentrations (K_i 1.05 μM), while tolterodine immediate release as well as the 5-HMT are devoid of any significant inhibitory potential regarding the other isoenzymes.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenicity studies with tolterodine were conducted in mice and rats. At the maximum tolerated dose in mice (30 mg/kg/day), female rats (20 mg/kg/day), and male rats (30 mg/kg/day), exposure margins were approximately 6 to 9 times, 7 times, and 11 times the clinical exposure to the pharmacologically active components of tolterodine tartrate extended-release capsules (based on AUC of tolterodine and its 5-HMT metabolite). At these exposure margins, no increase in tumors was found in either mice or rats.

No mutagenic or genotoxic effects of tolterodine were detected in a battery of *in vitro* tests, including bacterial mutation assays (Ames test) in 4 strains of *Salmonella typhimurium* and in 2 strains of *Escherichia coli*, a gene mutation assay in L5178Y mouse lymphoma cells, and chromosomal aberration tests in human lymphocytes. Tolterodine was also negative *in vivo* in the bone marrow micronucleus test in the mouse.

In female mice treated for 2 weeks before mating and during gestation with 20 mg/kg/day (about 9 to 12 times the clinical exposure via AUC), neither effects on reproductive performance or fertility were seen. In male mice, a dose of 30 mg/kg/day did not induce any adverse effects on fertility.

14 CLINICAL STUDIES

Tolterodine tartrate extended-release capsules 2 mg were evaluated in 29 patients in a Phase 2 dose-effect study. Tolterodine tartrate extended-release capsules 4 mg were evaluated for the treatment of overactive bladder with symptoms of urge urinary incontinence and frequency in a randomized, placebo-controlled, multicenter, double-blind, Phase 3, 12-week study. A total of 507 patients received tolterodine tartrate extended-release capsules 4 mg once daily in the morning and 508 received placebo. The majority of patients were Caucasian (95%) and female (81%), with a mean age of 61 years (range, 20 to 93 years). In the study, 642 patients (42%) were 65 to 93 years of age. The study included patients known to be responsive to tolterodine immediate release and other anticholinergic medications, however, 47% of patients never received prior pharmacotherapy for overactive bladder. At study entry, 97% of patients had at least 5 urge incontinence episodes per week and 91% of patients had 8 or more micturitions per day.

The primary efficacy assessment was change in mean number of incontinence episodes per week at week 12 from baseline. Secondary efficacy measures included change in mean number of micturitions per day and mean volume voided per micturition at week 12 from baseline.

Patients treated with tolterodine tartrate extended-release capsules experienced a statistically significant decrease in number of urinary incontinence per week from baseline to last assessment (week 12) compared with placebo as well as a decrease in the average daily urinary frequency and an increase in the average urine volume per void.

Mean change from baseline in weekly incontinence episodes, urinary frequency, and volume voided between placebo and tolterodine tartrate extended-release capsules are summarized in Table 4.

Table 4. 95% Confidence Intervals (CI) for the Difference between Tolterodine Tartrate Extended-Release Capsules (4 mg daily) and Placebo for Mean Change at Week 12 from Baseline*

	Tolterodine Tartrate Extended-Release Capsules (n=507)	Placebo (n=508)†	Treatment Difference, vs. Placebo (95% CI)
Number of incontinence episodes/week			
Mean Baseline	22.1	23.3	-4.8‡
Mean Change from Baseline	-11.8 (SD 17.8)	-6.9 (SD 15.4)	(-6.9, -2.8)
Number of micturitions/day			
Mean Baseline	10.9	11.3	-0.6‡
Mean Change from Baseline	-1.8 (SD 3.4)	-1.2 (SD 2.9)	(-1, -0.2)
Volume voided per micturition (mL)			
Mean Baseline	141	136	20‡
Mean Change from Baseline	34 (SD 51)	14 (SD 41)	(14, 26)

SD = Standard Deviation.

* Intent-to-treat analysis.

† 1 to 2 patients missing in placebo group for each efficacy parameter.

‡ The difference between tolterodine tartrate extended-release capsules and placebo was statistically significant.

16 HOW SUPPLIED/STORAGE AND HANDLING

Tolterodine tartrate extended-release capsules, 2 mg are blue green colored size '4' hard gelatin capsules imprinted with 'J' on cap and '104' on body, filled with white to off-white pellets. They are supplied as follows:

- Bottles of 30 capsules NDC 31722-607-30
- Bottles of 90 capsules NDC 31722-607-90
- Bottles of 500 capsules NDC 31722-607-05

Tolterodine tartrate extended-release capsules 4 mg are blue colored size '3' hard gelatin capsule imprinted with 'J' on cap and '105' on body, filled with white to off-white pellets. They are supplied as follows:

- Bottles of 30 capsules NDC 31722-608-30
- Bottles of 90 capsules NDC 31722-608-90
- Bottles of 500 capsules NDC 31722-608-05

Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from light.

17 PATIENT COUNSELING INFORMATION

See FDA-Approved Patient Labeling (17.2).

17.1 Information for Patients

Patients should be informed that antimuscarinic agents such as tolterodine tartrate extended-release capsules may produce the following effects: blurred vision, dizziness, or drowsiness. Patients should be advised to exercise caution in decisions to engage in potentially dangerous activities until the drug's effects have been determined.



Manufactured for: Camber Pharmaceuticals, Inc. Piscataway, NJ 08854

By: HETERO™ Hetero Labs Limited Jeedimetla, Hyderabad - 500 055, India

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Take tolterodine tartrate extended-release capsules once a day with liquid. Swallow the whole capsule. Tell your doctor if you cannot swallow a capsule.

- Tolterodine tartrate extended-release capsules can be taken with or without food.
- Take tolterodine tartrate extended-release capsules the same time each day.
- If you miss a dose of tolterodine tartrate extended-release capsules, begin taking tolterodine tartrate extended-release capsules again the next day. Do not take 2 doses of tolterodine tartrate extended-release capsules in the same day.
- If you took more than your prescribed dose of tolterodine tartrate extended-release capsules, call your doctor or poison control center, or go to the hospital emergency room.

What are possible side effects of tolterodine tartrate extended-release capsules?

Tolterodine tartrate extended-release capsules may cause allergic reactions that may be serious. Symptoms of a serious allergic reaction may include swelling of the face, lips, throat, or tongue. If you experience these symptoms, you should stop taking tolterodine tartrate extended-release capsules and get emergency medical help right away.

The most common side effects with tolterodine tartrate extended-release capsules are:

- Dry mouth
- Constipation
- Headache
- Stomach pain

Medicines like tolterodine tartrate extended-release capsules can cause blurred vision, dizziness, and drowsiness.

Do not drive, operate machinery, or do other dangerous activities until you know how tolterodine tartrate extended-release capsules affects you.

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

These are not all the side effects with tolterodine tartrate extended-release capsules. For a complete list, ask your doctor or pharmacist.

How do I store tolterodine tartrate extended-release capsules?

- Store tolterodine tartrate extended-release capsules at room temperature, 20° to 25°C (68° to 77°F). [see USP Controlled Room Temperature]. Protect from light. Keep in a dry place.
- Keep tolterodine tartrate extended-release capsules and all medicines out of the reach of children.

General information about tolterodine tartrate extended-release capsules

Medicines are sometimes prescribed for conditions that are not listed in this information. If you have questions about this medicine, ask your doctor. You can ask your doctor or pharmacist for information about tolterodine tartrate extended-release capsules that is written for health professionals.

What are the ingredients in tolterodine tartrate extended-release capsules?

Active ingredients: tolterodine tartrate

Inactive ingredients are aqueous ethylcellulose dispersion, dehydrated alcohol, dibasic calcium phosphate anhydrous, ethyl acrylate and methyl methacrylate copolymer dispersion, ethyl cellulose, FD & C Blue 2, gelatin, hydroxypropyl methylcellulose, microcrystalline cellulose, talc, and titanium dioxide. Additionally 2 mg capsules contain iron oxide yellow.

Both the 2 mg and 4 mg strengths capsule shells are imprinted with a pharmaceutical grade printing ink that contains butyl alcohol, dehydrated alcohol, isopropyl alcohol, potassium hydroxide, propylene glycol, shellac, strong ammonia solution and titanium dioxide.

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