

**CENTER FOR DRUG EVALUATION AND
RESEARCH**

APPLICATION NUMBER:

75609

DRAFT FINAL PRINTED LABELING

**Doxazosin Mesylate
Tablets**

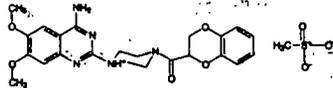
Rx Only

APPROVED

OCT 18 2000

DESCRIPTION

Doxazosin mesylate is a quinazolinone compound that is a selective inhibitor of the alpha₁ subtype of alpha adrenergic receptors. The chemical name of doxazosin mesylate is 1-(4-oxo-6,7-dimethoxy-2-quinazolinyl)-4-(1,4-benzodioxo-2-ylcarbonyl) piperazine methanesulfonate. The molecular formula for doxazosin mesylate is C₂₃H₂₆N₄O₆ • CH₃SO₃ and the molecular weight is 547.6. It has the following structure:



Doxazosin mesylate is freely soluble in dimethylsulfoxide, soluble in dimethylformamide, slightly soluble in methanol, ethanol, and water (0.8% at 25°C), and very slightly soluble in acetone and methylcyclohexane. Each doxazosin mesylate tablet (colored) for oral administration, contains 1 mg (gray), 2 mg (yellow), 4 mg (pink) and 8 mg (blue) of doxazosin as the free base.

In addition, each doxazosin mesylate tablet contains the following inactive ingredients: lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium lauryl sulfate and sodium starch glycolate. The 1 mg tablet contains FD&C Blue No. 2 Aluminum Lake, FD&C Yellow No. 6 Aluminum Lake, and FD&C Red No. 40 Aluminum Lake; the 2 mg tablet contains D&C Yellow No. 19 Aluminum Lake; the 4 mg tablet contains D&C Red No. 27 Aluminum Lake and D&C Red No. 30 Aluminum Lake; the 8 mg tablet contains FD&C Blue No. 9 Aluminum Lake.

CLINICAL PHARMACOLOGY

Pharmacokinetics

A. Single Peroral Administration (BPH)

Benign prostatic hypertrophy (BPH) is a common cause of urinary outflow obstruction in aging males. Severe BPH may lead to urinary retention and renal damage. A static and a dynamic component contribute to the symptoms and reduced urinary flow rate associated with BPH. The static component is related to an increase in prostate size caused, in part, by a proliferation of smooth muscle cells in the prostatic stroma. However, the severity of BPH symptoms and the degree of urethral obstruction do not correlate well with the size of the prostate. The dynamic component of BPH is associated with an increase in smooth muscle tone in the prostate and bladder neck. The degree of tone in this area is mediated by the alpha₁ adrenoceptor, which is present in high density in the prostatic stroma, prostatic capsule and bladder neck. Blockade of the alpha₁ receptor decreases urethral resistance and may relieve the obstruction and BPH symptoms. In the human prostate, doxazosin antagonizes phenylephrine (alpha₁ agonist)-induced contractions, *in vitro*, and blocks with high affinity to the alpha₁ adrenoceptor. The receptor subtype is thought to be the predominant functional type in the prostate. Doxazosin acts within 1-2 weeks to decrease the severity of BPH symptoms and improve urinary flow rate. Since alpha₁ adrenoceptors are of low density in the urinary bladder (apart from the bladder neck), doxazosin should maintain bladder contractility.

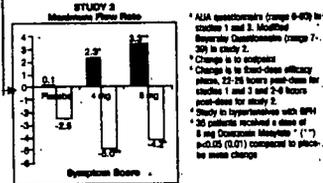
The efficacy of doxazosin was evaluated extensively in over 800 patients with BPH in double-blind, placebo-controlled trials. Doxazosin treatment was superior to placebo in improving patient symptoms and urinary flow rate. Significant relief with doxazosin was seen as early as one week into the treatment regimen, with doxazosin treated patients (N=173) showing a significant (p<0.01) increase in maximum flow rate of 0.8 mL/sec compared to a decrease of 0.3 mL/sec in the placebo group (N=41). In long-term studies improvement was maintained for up to 2 years of treatment. In 55-71% of patients, improvements above baseline were seen in both symptoms and maximum urinary flow rate.

In three placebo-controlled studies of 14-16 weeks duration obstructive symptoms (hesitation, intermittency, dribbling, weak urinary stream, incomplete emptying of the bladder) and irritative symptoms (nocturia, daytime frequency, urgency, burning) of BPH were evaluated at each visit by patient-assessed symptom questionnaires. The bothersomeness of symptoms was measured with a modified Beyerly questionnaire. Symptom severity/frequency was assessed using a modified Beyerly questionnaire or an ALA-based questionnaire. Urinary flow rate evaluations were performed at times of peak (2-4 hours post-dose) and/or trough (24 hours post-dose) plasma concentrations of doxazosin. The results from the three placebo-controlled studies (N=800) showing significant efficacy with 4 mg and 8 mg doxazosin mesylate are summarized in Table 1. In all three studies, doxazosin mesylate tablets resulted in statistically significant relief of obstructive and irritative symptoms compared to placebo. Statistically significant improvements of 2.3-3.3 mL/sec in maximum flow rate were seen with doxazosin mesylate tablets to Studies 1 and 2, compared to 0.1-0.7 mL/sec with placebo.

TABLE 1

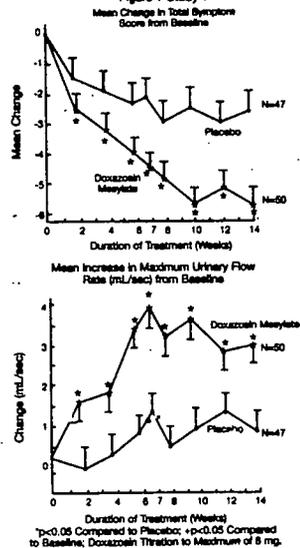
TABLE 1
SUMMARY OF EFFECTIVENESS DATA IN PLACEBO-CONTROLLED TRIALS

STUDY (Titration to maximum dose of 8 mg)*	N	SYMPTOM SCORE†		MAXIMUM FLOW RATE (mL/sec)	
		Baseline	Change	Baseline	Change
Placebo	47	15.8	-2.3	41	9.7
Doxazosin Mesylate	49	14.5	-4.9**	41	9.8
STUDY 2 (Titration to fixed dose 14 weeks)†					
Placebo	37	20.7	-2.5	30	10.8
Doxazosin Mesylate 4 mg	38	21.2	-5.0**	32	9.8
Doxazosin Mesylate 8 mg	42	19.9	-4.2*	38	10.5
STUDY 3 (Titration to fixed dose-12 weeks)					
Placebo	47	14.9	-4.7	44	9.9
Doxazosin Mesylate 4 mg	46	16.6	-5.1*	46	9.8



In one fixed dose study (study 2) doxazosin therapy (4-8 mg, once daily) resulted in a significant and sustained improvement in maximum urinary flow rate of 2.3-3.3 mL/sec (Table 1) compared to placebo (0.1 mL/sec). In this study, the only study in which weekly evaluations were made, significant improvement with doxazosin vs. placebo was seen after one week. The proportion of patients who responded with a maximum flow rate improvement of 2.3 mL/sec was significantly larger with doxazosin (24-42%) than placebo (13-17%). A significantly greater improvement was also seen in average flow rate with doxazosin (1.8 mL/sec) than with placebo (0.2 mL/sec). The onset and time course of symptom relief and increased urinary flow from study 1 are illustrated in Figures 1.

Figure 1- Study 1



*p<0.05 Compared to Placebo; **p<0.05 Compared to Baseline; Doxazosin Titration to Maximum of 8 mg.

In BPH patients (N=450) treated for 20 to 2 years in open-label studies, doxazosin therapy resulted in significant improvement above baseline in urinary flow rate and BPH symptoms. The significant effects of doxazosin were maintained over the entire treatment period.

TABLE 2

Mean Changes in Blood Pressure from baseline to the Mean of the Final Efficacy Phase in Normotensives (Diastolic BP<90 mmHg) in Two Double-blind, Placebo-controlled U.S. Studies with Doxazosin Mesylate Tablets 1-8 mg once daily.

Sitting BP (mmHg)	PLACEBO (N=86)		DOXAZOSIN MESYLATE (N=183)	
	Baseline	Change	Baseline	Change
Systolic	129.4	-1.6	129.4	-4.9*
Diastolic	79.2	-1.2	79.6	-2.4*
Standing BP (mmHg)	PLACEBO (N=86)		DOXAZOSIN MESYLATE (N=183)	
Systolic	128.5	-0.8	128.5	-5.3*
Diastolic	80.5	-0.7	80.4	-2.6*

*p<0.05 compared to placebo

doxazosin, other symptoms of lowered blood pressure, such as dizziness, light-headedness, or vertigo can occur, especially at initiation of therapy or at the time of dose increases.

a) Hypertension
These symptoms were common in clinical trials in hypertension, occurring in up to 23% of all patients treated and causing discontinuation of therapy in about 2%. In placebo-controlled titration trials in hypertension, orthostatic effects were minimized by beginning therapy at 1 mg per day and titrating every two weeks to 2, 4, or 8 mg per day. There was an increased frequency of orthostatic effects in patients given 8 mg or more, 10%, compared to 5% at 1-4 mg and 3% in the placebo group.

b) Benign Prostatic Hypertrophy
In placebo-controlled trials in BPH, the incidence of orthostatic hypotension with doxazosin was 0.3% and did not increase with increasing dosage (to 8 mg/day). The incidence of discontinuations due to hypotensive or orthostatic symptoms was 3.3% with doxazosin and 1% with placebo. The titration interval in these studies was one to two weeks.

Patients in occupations in which orthostatic hypotension could be dangerous should be treated with particular caution. As alpha₁ antagonists can cause orthostatic effects, it is important to evaluate standing blood pressure two minutes after standing and patients should be advised to exercise care when arising from a supine or sitting position.

If hypotension occurs, the patient should be placed in the supine position and, if this measure is inadequate, volume expansion with intravenous fluids or vasopressor therapy may be used. A transient hypotensive response is not a contraindication to further doses of doxazosin mesylate.

Information for Patients (See Patient Package Insert): Patients should be made aware of the possibility of syncope and orthostatic symptoms, especially at the initiation of therapy, and urged to avoid driving or hazardous tasks for 24 hours

8. Hypertension

The mechanism of action of doxazosin is selective blockade of the alpha₁ (postjunctional) subtype of adrenergic receptors. Studies in normal human subjects have shown that doxazosin competitively antagonized the pressor effects of phenylephrine (an alpha₁ agonist) and the systolic pressor effect of norepinephrine. Doxazosin and prazosin have similar abilities to antagonize phenylephrine. The antihypertensive effect of doxazosin results from a decrease in systemic vascular resistance. The parent compound doxazosin is primarily responsible for the antihypertensive activity. The low plasma concentrations of known active and inactive metabolites of doxazosin (2-hydroxy, 6'- and 7'-hydroxy and 6'- and 7'-O-desmethyl compounds) compared to parent drug indicate that the contribution of even the most potent compound (6'-hydroxy) to the antihypertensive effect of doxazosin in man is probably small. The 6'- and 7'-hydroxy metabolites have demonstrated antitussive properties at concentrations of 5 µM, *in vitro*.

Administration of doxazosin results in a reduction in systemic vascular resistance. In patients with hypertension there is little change in cardiac output. Maximum reductions in blood pressure usually occur 2-6 hours after dosing and are associated with a small increase in standing heart rate. Like other alpha₁ adrenergic blocking agents, doxazosin has a greater effect on blood pressure and heart rate in the standing position.

In a pooled analysis of placebo-controlled hypertension studies with about 300 hypertensive patients per treatment group, doxazosin, at doses of 1-16 mg given once daily, lowered blood pressure at 24 hours by about 108 mmHg compared to placebo in the standing position and about 89 mmHg in the supine position. Peak blood pressure effects (1-6 hours) were larger by about 60-75% (i.e., trough values were about 55-70% of peak effect), with the larger peak-trough differences seen in systolic pressures. There was no apparent difference in blood pressure responses of Caucasians and Afro-Americans or of patients above and below age 65. In these predominantly normocholesterolemic patients doxazosin produced small reductions in total serum cholesterol (2-3%), LDL cholesterol (4%), and a similarly small increase in HDL total cholesterol ratio (1%). The clinical significance of these findings is uncertain. In the same patient population, patients receiving doxazosin mesylate tablets gained a mean of 0.6 kg compared to a mean loss of 0.1 kg for placebo patients.

Pharmacokinetics

After oral administration of therapeutic doses, peak plasma levels of doxazosin occur at about 2-3 hours. Bioavailability is approximately 65%, reflecting first pass metabolism of doxazosin by the liver. The effect of food on the pharmacokinetics of doxazosin was evaluated in a crossover study with twelve hypertensive subjects. Reductions of 10% in mean maximum plasma concentration and 12% in the area under the concentration-time curve occurred when doxazosin was administered with food. Neither of these differences was statistically or clinically significant.

Doxazosin is extensively metabolized in the liver, mainly by O-demethylation of the quinazoline nucleus or hydroxylation of the benzodioxan moiety. Although several active metabolites of doxazosin have been identified, the pharmacokinetics of these metabolites have not been characterized. In a study of two subjects administered radiolabelled doxazosin 2 mg orally and 1 mg intravenously on two separate occasions, approximately 63% of the dose was eliminated in the feces and 9% of the dose was found in the urine. On average only 4.8% of the dose was excreted as unchanged drug in the feces and only a trace of the total radioactivity in the urine was attributed to unchanged drug. At the plasma concentrations achieved by therapeutic doses approximately 98% of the circulating drug is bound to plasma proteins.

Plasma elimination of doxazosin is biphasic, with a terminal elimination half-life of about 12 hours. Single-dose studies in hypertensive patients given doxazosin doses of 2-16 mg once daily showed linear kinetics and dose proportionality. In two studies, following the administration of 2 mg orally once daily, the mean accumulation ratios (steady-state AUC vs. first dose AUC) were 1.2 and 1.7. Enterohepatic recirculation is suggested by secondary peaking of plasma doxazosin concentrations.

In a crossover study in 24 normotensive subjects, the pharmacokinetics and safety of doxazosin were shown to be similar with morning and evening dosing regimens. The area under the curve after morning dosing was, however, 11% less than that after evening dosing and the time to peak concentration after evening dosing occurred significantly later than that after morning dosing (5.6 hr vs. 3.5 hr).

The pharmacokinetics of doxazosin in young (<65 years) and elderly (≥65 years) subjects were similar for plasma half-life values and oral clearance. Pharmacokinetic studies in elderly patients and patients with renal impairment have shown no significant alterations compared to younger patients with normal renal function. Administration of a single 2 mg dose to patients with decreased (Child-Pugh Class A) showed a 40% increase in exposure to doxazosin. There are only limited data on the effects of drugs known to influence the hepatic metabolism of doxazosin (e.g., cimetidine (see PRECAUTIONS)). As with any drug wholly metabolized by the liver, use of doxazosin in patients with altered liver function should be undertaken with caution.

In two placebo-controlled studies, of normotensive and hypertensive BPW patients, in which doxazosin was administered in the morning and the afternoon interval was two weeks and one week, respectively, trough plasma concentrations of doxazosin were similar in the two populations. Linear kinetics and dose proportionality were observed.

INDICATIONS AND USAGE

A. Benign Prostatic Hypertrophy (BPH). Doxazosin mesylate tablets are indicated for the treatment of both the urinary outflow obstruction and obstructive and irritative symptoms associated with BPH obstructive symptoms (hesitancy, intermittency, dribbling, weak urinary stream, incomplete emptying of the bladder) and irritative symptoms (nocturia, daytime frequency, urgency, burning). Doxazosin may be used in all BPH patients whether hypertensive or normotensive. In patients with hypertension and BPH, both conditions were effectively treated with doxazosin monotherapy. Doxazosin provides rapid improvement in symptoms and urinary flow rate in 86-71% of patients. Sustained improvements with doxazosin were seen in patients treated for up to 14 weeks in double-blind studies and up to 2 years in open-label studies.

B. Hypertension. Doxazosin mesylate tablets are also indicated for the treatment of hypertension. Doxazosin may be used alone or in combination with diuretic, beta-adrenergic blocking agents, calcium channel blockers or angiotensin-converting enzyme inhibitors.

CONTRAINDICATIONS

Doxazosin mesylate tablets are contraindicated in patients with a known sensitivity to quinazolinones (e.g., prazosin, terazosin).

Warnings: Syncope and "first-dose" effect: doxazosin, like other alpha-1-adrenergic blocking agents, can cause marked hypotension, especially in the upright position. Marked orthostatic effects are most common with the first dose but can also occur when the effects are most common with the first dose but can also occur when there is a change in posture, or if therapy is interrupted for more than a few days. To determine the likelihood of excessive hypotension and syncope, it is essential that treatment be initiated with the 1 mg dose. The 2, 4, and 8 mg doses are not for initial therapy. Doses should then be adjusted slowly (see **DOSEAGE AND ADMINISTRATION**) with evaluation and increase in dose every two weeks to the recommended dose. Additional antihypertensive agents should be added with caution.

Patients being treated with doxazosin should be cautioned to avoid situations where injury could result should syncope occur, during both the day and night. In an early investigational study of the safety and tolerance of increasing daily doses of doxazosin in normotensives beginning at 1 mg/day, only 2 of 8 subjects could tolerate more than 2 mg/day without experiencing orthostatic hypotension. In another study of 24 healthy normotensive male subjects receiving initial doses of 2 mg/day of doxazosin, seven (29%) of the subjects experienced orthostatic hypotension between 0.5 and 6 hours after the first dose necessitating termination of the study. In this study 2 of the normotensive subjects experienced syncope. Subsequent trials in hypertensive patients also began doxazosin dosing at 1 mg/day resulting in a 4% incidence of postural side effects at 1 mg/day with no cases of syncope.

In multiple dose clinical trials in hypertension involving over 1500 hypertensive patients with dose titration every one to two weeks, syncope was reported in 0.7% of patients. None of these events occurred at the starting dose of 1 mg and 1.2% (n=664) occurred at 16 mg/day.

In placebo-controlled, clinical trials in BPH, 3 out of 655 patients (0.5%) taking doxazosin reported syncope. Two of the patients were taking 1 mg doxazosin, while one patient was taking 2 mg doxazosin when syncope occurred. In the open-label, long-term extension follow-up of approximately 450 BPH patients, there were 3 reports of syncope (0.7%). One patient was taking 2 mg, one patient was taking 8 mg and one patient was taking 12 mg when syncope occurred. In a clinical pharmacology study, one subject receiving 2 mg, experienced syncope.

If syncope occurs, the patient should be placed in a recumbent position and if necessary, oxygen should be administered.

Precautions: Rarely (probably less frequently than once in every several thousand patients), alpha-1 antagonists such as doxazosin have been associated with acute angle-closure (painful pupil) iritis, sustained for hours and unrelieved by usual iritis therapy. Because this condition can lead to permanent blindness if not promptly treated, patients must be advised about the seriousness of the condition (see **PRECAUTIONS: Information for the Patient**).

PRECAUTIONS

General: Prostate Cancer: Carcinoma of the prostate causes many of the symptoms associated with BPH and the two disorders frequently co-exist. Carcinoma of the prostate should therefore be ruled out prior to commencing therapy with doxazosin. Dehydration: While syncope is the most severe orthostatic effect of doxazosin, other symptoms of lowered blood pressure, such as dizziness, light-headedness, or vertigo can occur, especially at initiation of therapy or at the time of dose increase.

a) **Hypertension**
These symptoms were common in clinical trials in hypertension, occurring in up to 23% of all patients treated and causing discontinuation of therapy in about 2%. In placebo-controlled trials in hypertension, orthostatic effects were minimized by beginning therapy at 1 mg per day and titrating every two weeks to 2, 4, or 8 mg per day. There was an increased frequency of orthostatic effects in patients given 8 mg or more, 10%, compared to 5% at 1-4 mg and 3% in the placebo group.

b) **Benign Prostatic Hyperplasia**
In placebo-controlled trials in BPH, the incidence of orthostatic hypotension with doxazosin was 0.3% and did not increase with increasing dosage (to 8 mg/day). The incidence of discontinuation due to hypotensive or orthostatic symptoms was 1.3% with doxazosin and 1% with placebo. The titration interval in these studies was one to two weeks. Patients in occupations in which orthostatic hypotension could be dangerous should be treated with particular caution. As alpha-1 antagonists can cause orthostatic effects, it is important to evaluate standing blood pressure two minutes after standing and patients should be advised to exercise care when arising from a supine or sitting position. If hypotension occurs, the patient should be placed in the supine position and, if the measure is inadequate, volume expansion with intravenous fluids or vasoconstrictor therapy may be used. A transient hypotensive response is not a contraindication to further doses of doxazosin therapy. Patients should be made aware of the possibility of syncope and/or orthostatic symptoms, especially at the initiation of therapy, and urged to avoid driving or hazardous work for 24 hours.

after the first dose, after a dosage increase, and after interruption of therapy when treatment is resumed. They should be cautioned to avoid situations where injury could result should syncope occur during initiation of doxazosin therapy. They should also be advised of the need to sit or lie down when symptoms of lowered blood pressure occur, although these symptoms are not always orthostatic, and to be careful when rising from a sitting or lying position. If dizziness, lightheadedness, or palpitations are bothersome they should be reported to the physician, so that dose adjustment can be considered. Patients should also be told that drowsiness or somnolence can occur with doxazosin or any selective alpha₁ adrenoceptor antagonist, requiring caution in people who must drive or operate heavy machinery.

Patients should be advised about the possibility of priapism as a result of treatment with alpha₁ antagonists. Patients should know that this adverse event is very rare. If they experience priapism, it should be brought to immediate medical attention for, if not treated promptly, it can lead to permanent erectile dysfunction (impotence).

Drug/Laboratory Test Interactions: Doxazosin does not affect the plasma concentration of prostate specific antigen in patients treated for up to 3 years. Both doxazosin, an alpha₁ inhibitor, and finasteride, a 5-alpha reductase inhibitor, are highly protein bound and metabolized hepatically. There is no definitive controlled clinical experience on the concurrent use of alpha₁ inhibitors and 5-alpha reductase inhibitors at this time.

Impaired Liver Function: Doxazosin enalapril should be administered with caution to patients with evidence of impaired hepatic function or to patients receiving drugs known to influence hepatic metabolism (see CLINICAL PHARMACOLOGY).
Laboratory/Investigations: Analysis of hematologic data from hypertensive patients receiving doxazosin in controlled hypertension clinical trials showed that the mean WBC (N=74) and mean neutrophil counts (N=119) were decreased by 2.4% and 1.0%, respectively, compared to placebo. A phenomenon not seen with other alpha₁ blocking drugs. In BPH patients, the incidence of clinically significant WBC abnormalities was 0.4% (2/452) with doxazosin and 0% (0/147) with placebo, with no statistically significant difference between the two treatment groups. A search through a data base of 2400 hypertensive patients and 665 BPH patients revealed 4 hypertensives in which drug-related neutropenia could not be ruled out and one BPH patient in which drug related leukopenia could not be ruled out. Two hypertensives had a single low value on the last day of treatment. Two hypertensives had stable, non-progressive neutrophil counts in the 1000/mm³ (cubed) range over periods of 20 and 40 weeks. One BPH patient had a decrease from a WBC count of 4800/mm³ (cubed) to 2700/mm³ (cubed) at the end of the study; there was no evidence of clinical impairment. In cases where follow-up was available the WBC and neutrophil counts returned to normal after discontinuation of doxazosin. No patients became symptomatic as a result of the low WBC or neutrophil counts.

Drug Interactions: Most (80%) of plasma doxazosin is protein bound. *In vitro* data in human plasma indicate that doxazosin has no effect on protein binding of digoxin, warfarin, phenytoin or indomethacin. There is no information on the effect of other highly plasma protein bound drugs on doxazosin binding. Doxazosin has been administered without any evidence of an adverse drug interaction to patients receiving thyroid diuretics, beta-blocking agents, and non-steroidal anti-inflammatory drugs. In a placebo-controlled trial in hypertensives, the administration of a single 1 mg dose of doxazosin on day 1 of a 4-day regimen of oral cimetidine (400 mg twice daily) resulted in a 10% increase in mean AUC of doxazosin (p<0.005), and a slight but not statistically significant increase in mean C_{max} and mean half-life of doxazosin. The clinical significance of this increase in doxazosin AUC is unknown.

In clinical trials, doxazosin enalapril tablets have been administered to patients on a variety of concomitant medications; while no formal interaction studies have been conducted, no interactions were observed. Doxazosin enalapril tablets have been used with the following drugs or drug classes: 1) analgesic/anti-inflammatory (e.g., acetaminophen, aspirin, codeine and codeine combinations, ibuprofen, indomethacin); 2) antibiotics (e.g., erythromycin, trimethoprim and sulfamethoxazole, ampicillin); 3) anticholinergics (e.g., chlorpheniramine); 4) cardiovascular agents (e.g., atenolol, hydrochlorothiazide, and furosemide); 5) corticosteroids; 6) gastrointestinal agents (e.g., antacids); 7) hypoglycemics and insulin; 8) sedatives and tranquilizers (e.g., diazepam); 9) cold and flu remedies.

Cardiac Toxicity in Animals: An increased incidence of myocardial necrosis or fibrosis was displayed by Sprague-Dawley rats after 6 months of dietary administration at concentrations calculated to provide 80 mg doxazosin/kg/day and after 12 months of dietary administration at concentrations calculated to provide 40 mg doxazosin/kg/day (AUC exposure in rats 8 times the human AUC exposure with a 12 mg/day therapeutic dose). Myocardial fibrosis was observed in both rats and mice treated in the same manner with 40 mg doxazosin/kg/day for 18 months (exposure 8 times human AUC exposure in rats and somewhat equivalent to human C_{max} exposure in mice). No cardiotoxicity was observed at lower doses (up to 10 or 20 mg/kg/day, depending on the study) in either species. These lesions were not observed after 12 months of oral dosing in dogs at maximum doses of 20 mg/kg/day (maximum plasma concentrations (C_{max}) in dogs 14 times the C_{max} exposure in humans receiving a 12 mg/day therapeutic dose) and in Wistar rats at doses of 100 mg/kg/day (C_{max} exposure 15 times human C_{max} exposure with a 12 mg/day therapeutic dose). There is no evidence that similar lesions occur in humans.

Cardiogenesis, Mutagenesis, Impairment of Fertility: Chronic dietary administration (up to 24 months) of doxazosin at maximally tolerated doses of 40 mg/kg/day in rats and 120 mg/kg/day in mice revealed no evidence of carcinogenic potential. The highest doses evaluated in the rat and mouse studies are associated with AUCs (a measure of systemic exposure) that are 8 times and 4 times, respectively, the human AUC at a dose of 18 mg/day.

Mutagenicity studies revealed no drug- or metabolite-related effects at either chromosomal or subchromosomal levels.

Studies in rats showed reduced fertility in males treated with doxazosin at oral doses of 20 (but not 5 or 10) mg/kg/day, about 4 times the AUC exposures obtained with a 12 mg/day human dose. This effect was reversible within two weeks of drug withdrawal. There have been no reports of any effects of doxazosin on male fertility in humans.

Pregnancy Teratogenic Effects, Pregnancy Category C: Studies in pregnant rabbits and rats at daily oral doses of up to 41 and 20 mg/kg, respectively (plasma drug concentrations 10 and 4 times human C_{max} and AUC exposures with a 12 mg/day therapeutic dose), have revealed no evidence of harm to the fetus. A dosage regimen of 52 mg/kg/day in the rabbit was associated with reduced fetal survival. There are no adequate and well-controlled studies in pregnant women. Because animal reproduction studies are not always predictive of human response, doxazosin should be used during pregnancy only if clearly needed. Radioactivity was found to cross the placenta following oral administration of labeled doxazosin to pregnant rats.

Neonatal/Postnatal Effects: In perinatal studies in rats, postnatal development at maternal doses of 40 or 50 mg/kg/day of doxazosin (8 times human AUC exposures with a 12 mg/day therapeutic dose) was delayed as evidenced by slower body weight gain and slightly later specification of anatomical features and reflexes.

Nursing Mothers: Studies in lactating rats (after a single oral dose of 1 mg/kg of [¹⁴C] doxazosin) indicate that doxazosin accumulates in rat breast milk with a maximum concentration about 20 times greater than the maternal plasma concentration. It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when doxazosin is administered to a nursing mother.

Pediatric Use: The safety and effectiveness of doxazosin as an antihypertensive agent have not been established in pediatric patients.

Use in Elderly: The safety and effectiveness profile of doxazosin in BPH was similar in the elderly (age ≥65 years) and younger (age <65 years) patients.

ADVERSE REACTIONS

A. Single Prescription Hypertensives

The incidence of adverse events has been ascertained from worldwide clinical trials in 965 BPH patients. The incidence rates presented below (Table 3) are based on combined data from seven placebo-controlled trials involving once daily administration of doxazosin in doses of 1-18 mg in hypertensives and 0.5-8 mg in normotensives. The adverse events when the incidence in the doxazosin group was at least 1% are summarized in Table 3. No significant differences in the incidence of adverse events compared to placebo was seen except for dizziness, fatigue, hypotension, edema and eye pain. Dizziness and dyspnea appeared to be dose-related.

TABLE 4
ADVERSE REACTIONS DURING
PLACEBO-CONTROLLED STUDIES
BENIGN PROSTATIC HYPERPLASIA

Body System	Docosozin (N=665)	PLACEBO (N=300)
BODY AS A WHOLE		
Back pain	1.8%	2.0%
Chest pain	1.2%	0.7%
Fatigue	8.0%	1.7%
Headache	8.9%	8.0%
Influenza-like symptoms	1.1%	1.0%
Pain	2.0%	1.0%
CARDIOVASCULAR SYSTEM		
Hypotension	1.7%*	0.0%
Palpitation	1.2%	0.3%
DIGESTIVE SYSTEM		
Abdominal Pain	2.4%	2.0%
Diarrhea	2.3%	2.0%
Dyspepsia	1.7%	1.7%
Nausea	1.5%	0.7%
METABOLIC AND NUTRITIONAL DISORDERS		
Edema	2.7%*	0.7%
NERVOUS SYSTEM		
Dizziness**	15.0%*	9.0%
Mouth Dry	1.4%	0.3%
Somnolence	3.0%	1.0%
RESPIRATORY SYSTEM		
Dyspnea	8.8%*	0.3%
Respiratory Disorders	1.1%	0.7%
SPECIAL SENSES		
Vision Abnormal	1.4%	0.7%
URINARY SYSTEM		
Incontinence	1.1%	1.0%
Urinary Tract Infection	1.4%	2.3%
SKIN & APPENDAGES		
Sweating Increased	1.1%	1.0%
PSYCHIATRIC DISORDERS		
Anxiety	1.1%	0.3%
Insomnia	1.2%††	0.2%

*p < 0.05 for treatment difference. **Includes vertigo.
 In these placebo-controlled studies of 665 docosozin patients, treated for a mean of 85 days, additional adverse reactions have been reported. These are less than 1% and not distinguishable from those that occurred in the placebo group. Adverse reactions with an incidence of less than 1% but of clinical interest are: (docosozin vs. placebo): Cardiovascular system: angina pectoris (0.8% vs. 0.7%), postural hypotension (0.2% vs. 0.3%), syncope (0.5% vs. 0.0%), tachycardia (0.3% vs. 0.0%); Urinary system: dysuria (0.5% vs. 1.3%), and Psychiatric disorders: libido decreased (0.8% vs. 0.3%). The safety profile in patients treated for up to three years was similar to that in the placebo-controlled studies. The majority of adverse experiences with docosozin were mild.

B. Hypertension
 Docosozin has been administered to approximately 4000 hypertensive patients, of whom 1679 were included in the hypertension clinical development program. In that program, minor adverse effects were frequent, but led to discontinuation of treatment in only 7% of patients. In placebo-controlled studies adverse effects occurred in 49% and 40% of patients in the docosozin and placebo groups, respectively, and led to discontinuation in 2% of patients in each group. The major reasons for discontinuation were postural effects (2%), edema, malaise/fatigue, and some heart rate disturbances, each about 0.7%. In controlled hypertension clinical trials directly comparing docosozin to placebo there was no significant difference in the incidence of side effects, except for dizziness (including postural), weight gain, somnolence and fatigue/taste. Postural effects and edema appeared to be dose related. The prevalence rates presented below are based on combined data from placebo-controlled studies involving once daily administration of docosozin at doses ranging from 1-18 mg. Table 4 summarizes those adverse experiences (possibly/probably related) reported for patients in these hypertension studies where the prevalence rate in the docosozin group was at least 0.5% or where the reaction is of particular interest.

TABLE 4

TABLE 4
ADVERSE REACTIONS DURING
PLACEBO-CONTROLLED STUDIES*

	HYERTENSION	
	DOXAZOSIN (N=330)	PLACEBO (N=330)
CARDIOVASCULAR SYSTEM		
Dizziness	19%	0%
Vertigo	2%	1%
Postural Hypotension	0.3%	0%
Edema	4%	3%
Palpitation	2%	2%
Arrhythmia	1%	0%
Hypotension	1%	0%
Tachycardia	0.3%	1%
Peripheral Ischemia	0.3%	0%
SKIN & APPENDAGES		
Rash	1%	1%
Pruritus	1%	1%
MUSCULOSKELETAL SYSTEM		
Arthralgia/Arthritis	1%	0%
Muscle Weakness	1%	0%
Myalgia	1%	0%
CENTRAL & PERIPHERAL N.S.		
Headache	14%	16%
Paresthesia	1%	1%
Kinetic Disorders	1%	0%
Ataxia	1%	0%
Hypertonia	1%	0%
Muscle Cramps	1%	0%
AUTONOMIC		
Mouth Dry	2%	2%
Flushing	1%	0%
SPECIAL SENSES		
Vision Abnormal	2%	1%
Conjunctivitis/Eye Pain	1%	1%
Tinnitus	1%	0.3%
PSYCHIC		
Somnolence	5%	1%
Nervousness	2%	2%
Depression	1%	1%
Insomnia	1%	1%
Sexual Dysfunction	2%	1%
GASTROINTESTINAL		
Nausea	3%	4%
Diarrhea	2%	3%
Constipation	1%	1%
Dyspepsia	1%	1%
Flatulence	1%	1%
Abdominal Pain	0%	2%
Vomiting	0%	1%
RESPIRATORY		
Rhinitis	3%	1%
Dyspnea	1%	1%
Epistaxis	1%	0%
URINARY		
Polyuria	2%	0%
Urinary Incontinence	1%	0%
Micturition Frequency	0%	2%
GENERAL		
Fatigue/Weakness	12%	0%
Chest Pain	2%	2%
Aches	1%	1%
Face Edema	1%	0%
Pain	2%	2%

Additional adverse reactions have been reported, but these are, in general, not distinguishable from symptoms that might have occurred in the absence of exposure to doxazosin. The following adverse reactions occurred with a frequency of between 0.5% and 1%: syncope, hypoaesthesia, increased sweating, agitation, increased weight. The following additional adverse reactions were reported by <0.5% of 3360 patients who received doxazosin in controlled or open, short- or long-term clinical studies, including international studies: Cardiovascular System: angina pectoris, myocardial infarction, cerebrovascular accident; Autonomic Nervous System: parosmia; Metabolic: thirst, gout, hypokalemia, Heretotopic lymphadenopathy, purpura; Reproductive System: breast pain; Skin Disorders: alopecia, dry skin, eczema; Central Nervous System: paresis, tremor, twitching, confusion, migraine, impaired concentration; Psychiatric: paranoia, anorexia, emotional lability, abnormal thinking, depersonalization; Special Senses: anosmia, vertigo, taste perversion, photophobia, abnormal lacrimation; Gastrointestinal System: increased appetite, anorexia, fecal incontinence, gastroenteritis; Respiratory System: bronchospasm, sinusitis, coughing, pharyngitis; Urinary System: renal calculus; General Body System: hot flashes, back pain, infection, insect/bug, decreased weight, influenza-like symptoms.

Doxazosin has not been associated with any clinically significant changes in routine biochemical tests. No clinically relevant adverse effects were noted on sodium, potassium, serum glucose, uric acid, blood urea nitrogen, creatinine or liver function tests. Doxazosin has been associated with decreases in white blood cell counts (see PRECAUTIONS).

Experiences with doxazosin overdosage is limited. Two adolescents who each intentionally ingested 40 mg doxazosin mesylate tablets with diazepam or paracetamol, were treated with gastric lavage with activated charcoal and made full recoveries. A two-year-old child who accidentally ingested 4 mg doxazosin mesylate tablet was treated with gastric lavage and remained normotensive during the five-hour emergency room observation period. A 67-month-old child accidentally received a crushed 1 mg tablet of doxazosin mesylate and was reported to have been asleep. A 22-year-old female with chronic renal failure, epilepsy and depression intentionally ingested 80 mg doxazosin mesylate (blood level 0.9 ppm); normal values in hypertensives 0.02 ppm/L; death was attributed to a gross mal seizure resulting from hypotension. A 38-year-old female who ingested 70 mg doxazosin mesylate, alcohol and Deltaine® (Paracetamol) developed hypotension which responded to fluid therapy.

The oral LD_{50} of doxazosin is greater than 1000 mg/kg in mice and rats. The most likely manifestation of overdosage would be hypotension, for which the usual treatment would be intravenous infusion of fluid. As doxazosin is highly protein bound, dialysis would not be indicated.

DOXAZOSIN ADMINISTRATION

DOXAZOSIN SHOULD BE INDIVIDUALIZED. The initial dosage of doxazosin mesylate tablets in patients with hypertension and/or BPH is 1 mg given once daily in the a.m. or p.m. This starting dose is intended to minimize the frequency of postural hypotension and first dose syncope associated with doxazosin mesylate tablets. Postural effects are most likely to occur between 2 and 6 hours after a dose. Therefore blood pressure measurements should be taken during this time period after the first dose and with each increase in dose. If doxazosin mesylate tablet administration is discontinued for several days, therapy should be restarted using the initial dosing regimen.

A. BENIGN PROSTATIC HYPERPLASIA 1-mg once daily. The initial dosage of doxazosin mesylate tablets is 1 mg, given once daily in the a.m. or p.m. Depending on the individual patient's urodynamic and BPH symptomatology, dosage may then be increased to 2 mg and thereafter to 4 mg and 8 mg once daily, the maximum recommended dose for BPH. The recommended titration interval is 1-2 weeks. Blood pressure should be evaluated routinely in these patients.

B. HYPERTENSION 1-16 mg once daily. The initial dosage of doxazosin mesylate tablets is 1 mg given once daily. Depending on the individual patient's existing blood pressure response (based on measurements taken at 2-6 hours post-dose and 24 hours post-dose), dosage may then be increased to 2 mg and thereafter if necessary to 4 mg, 8 mg and 16 mg to achieve the desired reduction in blood pressure. Increases in dose beyond 4 mg increase the likelihood of excessive postural effects including syncope, postural dizziness/vertigo and postural hypotension. At a titrated dose of 16 mg once daily the frequency of postural effects is about 12% compared to 2% for placebo.

HOW SUPPLIED

Doxazosin mesylate tablets are available as:
1 mg, gray, capsule-shaped tablets, debossed "ETH287" on one side and "1" bisect "mg" on the other side in bottles of 100 tablets NDC 58177-268-04; bottles of 1000 tablets NDC 58177-268-09.

2 mg, yellow, capsule-shaped tablets, debossed "ETH287" on one side and "2" bisect "mg" on the other side in bottles of 100 tablets NDC 58177-267-04; bottles of 1000 tablets NDC 58177-267-09.

4 mg, pink, capsule-shaped tablets, debossed "ETH287" on one side and "4" bisect "mg" on the other side in bottles of 100 tablets NDC 58177-269-04; bottles of 1000 tablets NDC 58177-269-09.

8 mg, black, capsule-shaped tablets, debossed "ETH287" on one side and "8" bisect "mg" on the other side in bottles of 100 tablets NDC 58177-269-04; bottles of 1000 tablets NDC 58177-269-09.

Recommended Storage: Store at controlled room temperature 15° to 30°C (59° to 86°F)

Manufactured by
KV Pharmaceutical Co. for
ETHEX Corporation
St. Louis, MO 63043-2413


 NDC 58177-268-04
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
4 mg
 B Only
 100 Tablets
 P3148 5/00

*Each tablet contains doxazosin mesylate equivalent to 4 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.
 Manufactured by KV Pharmaceutical Co. for ETHEX Corporation, St. Louis, MO 63043-2413.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).


 NDC 58177-269-04
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
8 mg
 B Only
 100 Tablets
 P3152 5/00

*Each tablet contains doxazosin mesylate equivalent to 8 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.
 Manufactured by KV Pharmaceutical Co. for ETHEX Corporation, St. Louis, MO 63043-2413.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).


 NDC 58177-268-04
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
1 mg
 B Only
 100 Tablets
 P3133 5/00

*Each tablet contains doxazosin mesylate equivalent to 1 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.
 Manufactured by KV Pharmaceutical Co. for ETHEX Corporation, St. Louis, MO 63043-2413.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).


 NDC 58177-267-04
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
2 mg
 B Only
 100 Tablets
 P3141 5/00

*Each tablet contains doxazosin mesylate equivalent to 2 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.
 Manufactured by KV Pharmaceutical Co. for ETHEX Corporation, St. Louis, MO 63043-2413.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).


 NDC 58177-266-09
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
1 mg
 B Only
 1000 Tablets
 P3135 5/00

*Each tablet contains doxazosin mesylate equivalent to 1 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.
 Manufactured by KV Pharmaceutical Co. for ETHEX Corporation, St. Louis, MO 63043-2413.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).


 NDC 58177-267-09
APPROVED
OCT 18 2000
DOXAZOSIN MESYLATE
 Tablets
2 mg

*Each tablet contains doxazosin mesylate equivalent to 2 mg doxazosin.
 Dispense in a tight container as defined in the USP/NF.

ETHEX ETHEX ETHEX ETHEX

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information. Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).



58177-266-09 6

*Each tablet contains doxazosin mesylate equivalent to 1 mg doxazosin
Dispense in a tight container as defined in the USP/NF.

Manufactured by KV Pharmaceutical Co. for ETHEX Corporation St. Louis, MO 63043-2413

NDC 58177-266-09

Doxazosin Mesylate

Tablets
1 mg*
Rx Only
1000 Tablets

OCT 18 2000
APPROVED

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information.
Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).

P3135 5/00



*Each tablet contains doxazosin mesylate equivalent to 4 mg doxazosin

Dispense in a tight container as defined in the USP/NF.



N 3 58177-268-09 0

Manufactured by KV Pharmaceutical Co. for ETHEX Corporation St. Louis, MO 63043-2413

NDC 58177-268-09

Doxazosin Mesylate

Tablets
4 mg*
Rx Only
1000 Tablets

OCT 18 2000
APPROVED

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information.

Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).

P3150 5/00



*Each tablet contains doxazosin mesylate equivalent to 8 mg doxazosin

Dispense in a tight container as defined in the USP/NF.



N 3 58177-269-09 7

Manufactured by KV Pharmaceutical Co. for ETHEX Corporation St. Louis, MO 63043-2413

NDC 58177-269-09

Doxazosin Mesylate

Tablets
8 mg*
Rx Only
1000 Tablets

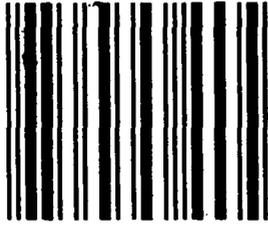
OCT 18 2000
APPROVED

DOSAGE AND ADMINISTRATION: See accompanying package insert for full prescribing information.

Store at controlled room temperature 15°C - 30°C (59°F - 86°F) (see USP).

P3157 5/00





Doxazosin Mesylate

Tablets

Patient Package Insert

OCT 18 2000:
APPROVED

PATIENT INFORMATION ABOUT DOXAZOSIN MESYLATE TABLETS

FOR BENIGN PROSTATIC HYPERPLASIA (BPH)

Read this leaflet:

- before you start taking doxazosin mesylate tablets
- each time you get a new prescription.

You and your doctor should discuss this treatment and your BPH symptoms before you start taking doxazosin mesylate tablets and at your regular checkups. This leaflet does NOT take the place of discussions with your doctor.

Doxazosin mesylate is used to treat both benign prostatic hyperplasia (BPH) and high blood pressure (hypertension). This leaflet describes doxazosin mesylate as treatment for BPH (although you may be taking doxazosin mesylate for both your BPH and high blood pressure).

What is BPH?

BPH is an enlargement of the prostate gland. This gland surrounds the tube that drains the urine from the bladder. The symptoms of BPH can be caused by a tensing of the enlarged muscle in the prostate gland which blocks the passage of urine. This can lead to such symptoms as:

- a weak or start-and-stop stream when urinating
- a feeling that the bladder is not completely emptied after urination
- a delay or difficulty in the beginning of urination
- a need to urinate often during the day and especially at night
- a feeling that you must urinate immediately.

TREATMENT OPTIONS FOR BPH

The four main treatment options for BPH are:

- If you are not bothered by your symptoms, you and your doctor may decide on a program of "watchful waiting." It is not an active treatment like taking medication or surgery but involves having regular checkups to see if your condition is getting worse or causing problems.
- Treatment with doxazosin mesylate or other similar drugs. Doxazosin mesylate is the medication your doctor has prescribed for you. See "What doxazosin mesylate Does," below.
- Treatment with the medication class of 5-alpha reductase inhibitors (e.g., Proscar[®]) It can cause the prostate to shrink. It may take 6 months or more for the full benefit of finasteride to be seen.

- Various surgical procedures: Your doctor can describe these procedures to you: The best procedure for you depends on your BPH symptoms and medical condition.

WHAT DOXAZOSIN MESYLATE DOES

Doxazosin mesylate tablets work on a specific type of muscle found in the prostate, causing it to relax. This in turn decreases the pressure within the prostate, thus improving the flow of urine and your symptoms.

- Doxazosin mesylate helps relieve the symptoms of BPH (weak stream, start-and-stop stream, a feeling that your bladder is not completely empty, delay in beginning of urination, need to urinate often during the day and especially at night, and feeling that you must urinate immediately). It does not change the size of the prostate. The prostate may continue to grow; however, a larger prostate is not necessarily related to more symptoms or to worse symptoms. Doxazosin mesylate can decrease your symptoms and improve urinary flow, without decreasing the size of the prostate.
- If doxazosin mesylate is helping you, you should notice an effect within 1 to 2 weeks after you start your medication. Doxazosin mesylate has been studied in over 900 patients for up to 2 years and the drug has been shown to continue to work during long-term treatment. Even though you take doxazosin mesylate and it may help you, doxazosin mesylate may not prevent the need for surgery in the future.
- Doxazosin mesylate does not affect PSA levels. PSA is the abbreviation for Prostate Specific Antigen. Your doctor may have done a blood test called PSA. You may want to ask your doctor more about this if you have had a PSA test done.

OTHER IMPORTANT FACTS

- You should see an improvement of your symptoms within 1 to 2 weeks. In addition to your other regular checkups you will need to continue seeing your doctor regularly to check your progress regarding your BPH and to monitor your blood pressure.
- Doxazosin mesylate is not a treatment for prostate cancer. Your doctor has prescribed doxazosin mesylate for your BPH and not for prostate cancer; however, a man can have BPH and prostate cancer at the same time. Doctors usually recommend that men be checked for prostate cancer once a year when they turn 50 (or 40 if a family member has had prostate cancer). A higher incidence of prostate cancer has been noted in men of African-American descent. These checks should continue even if you are taking doxazosin mesylate.

How To Take Doxazosin Mesylate and What You Should Know While Taking Doxazosin Mesylate for BPH

Doxazosin Mesylate Can Cause a Sudden Drop in Blood Pressure After the VERY FIRST DOSE. You may feel dizzy, faint or "light-headed," especially after you stand up from a lying or sitting position. This is more likely to occur after you've taken the first few doses or if you increase your dose, but can occur at any time while you are taking the drug. It can also occur if you stop taking the drug and then restart treatment. If you feel very dizzy, faint or "light-headed" you should contact your doctor. Your doctor will discuss with you how often you need to visit and how often your blood pressure should be checked. Your blood pressure should be checked when you start taking doxazosin mesylate even if you do not have high blood pressure (hypertension). Your doctor will discuss with you the details of how blood pressure is measured.

Blood Pressure Measurement: Whatever equipment is used, it is usual for your blood pressure to be measured in the following way: measure your blood pressure after lying quietly on your back for five minutes. Then, after standing for two minutes measure your blood pressure again. Your doctor will discuss with you what other times during the day your blood pressure should be taken, such as two to six hours after a dose, before bedtime or after waking up in the morning. Note that moderate to high-intensity exercise can, over a period of time, lower your average blood pressure.

You can take doxazosin mesylate either in the morning or at bedtime and it will be equally effective. If you take doxazosin mesylate at bedtime but need to get up from bed to go to the bathroom, get up slowly and cautiously until you are sure how the medication affects you. It is important to get up slowly from a chair or bed at any time until you learn how you react to doxazosin mesylate. You should not drive or do any hazardous tasks until you are used to the effects of the medication. If you begin to feel dizzy, sit or lie down until you feel better.

- You will start with a 1 mg dose of doxazosin mesylate once daily. Then the once daily dose will be increased as your body gets used to the effects of the medication. Follow your doctor's instructions about how to take doxazosin mesylate. You must take it every day at the dose prescribed. Talk with your doctor if you don't take it for a few days for some reason; you may then need to restart the medication at a 1 mg dose, increase your dose gradually and again be cautious about possible dizziness. Do not share doxazosin mesylate with anyone else; it was prescribed only for you.
- Other side effects you could have while taking doxazosin mesylate, in addition to lowering of the blood pressure, include dizziness, fatigue (tiredness), swelling of the feet and shortness of breath. Most side effects are mild. However, you should discuss any unexpected effects you notice with your doctor.
- **WARNING:** Extremely rarely, doxazosin mesylate and similar medications have caused painful erection of the penis, sustained for hours and unrelieved by sexual intercourse or masturbation. This condition is serious, and if untreated it can be followed by permanent inability to have an erection. If you have a prolonged abnormal erection, call your doctor or go to an emergency room as soon as possible.
- Keep doxazosin mesylate and all medicines out of the reach of children.

FOR MORE INFORMATION ABOUT DOXAZOSIN MESYLATE AND BPH TALK WITH YOUR DOCTOR, NURSE, PHARMACIST OR OTHER HEALTH CARE PROVIDER.

Manufactured by
KV Pharmaceutical Co. for
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St. Louis, MO 63043

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