

Tamsulosin capsules are administered concomitantly with Nifedipine, atenolol, or enalapril.

Warfarin: A definitive drug-drug interaction study between Tamsulosin Hydrochloride and warfarin was not conducted. Results from limited in vitro and in vivo studies are inconclusive. Therefore, caution should be exercised with concomitant administration of warfarin and TAMPRO™ capsules.

Digoxin and Theophylline

Dosage adjustments are not necessary when a TAMPRO capsule is administered concomitantly with digoxin or theophylline.

INDICATIONS AND USAGE

TAMPRO (Tamsulosin Hydrochloride) capsules are indicated for the treatment of the signs and symptoms of benign prostatic hyperplasia (BPH).

CONTRAINDICATIONS

TAMPRO capsules are contraindicated in patients known to be hypersensitive to Tamsulosin Hydrochloride or any component of TAMPRO capsules.

TAMPRO capsules are not indicated for the treatment of hypertension.

WARNINGS

The signs and symptoms of orthostasis (postural hypotension, dizziness and vertigo) were detected more frequently in Tamsulosin capsule-treated patients than in placebo recipients. As with other alpha-adrenergic blocking agents there is a potential risk of syncope

ADVERSE REACTIONS

Patients beginning treatment with Tamsulosin capsules should be cautioned to avoid situations where injury could result & should syncope occur. Doses of Tamsulosin higher than 0.4 mg (e.g., 0.8 mg) should not be used in combination with strong inhibitors of CYP3A4 (e.g., ketoconazole). Rarely (probably less than 1 in 50,000 patients), tamsulosin, like other alpha1 antagonists, has been associated with priapism (persistent painful penile erection unrelated to sexual activity). Because this condition can lead to permanent impotence if not properly treated, patients must be advised about the seriousness of the condition.

PREGNANCY

TAMPRO capsules are not indicated for use in women.

OVERDOSAGE

Overdosage of TAMPRO (Tamsulosin Hydrochloride) capsules can lead to hypotension.

DOSAGE AND ADMINISTRATION

TAMPRO capsules 0.4 mg once daily is recommended as the dose for the treatment of the signs and symptoms of BPH. It

should be administered approximately one-half hour following the same meal each day. For those patients who fail to respond to the 0.4 mg dose after two to four weeks of dosing, the dose of TAMPRO capsules can be increased to 0.8 mg once daily. The 0.8 mg dose should not be used in combination with strong inhibitors of CYP3A4 (e.g., ketoconazole). If TAMPRO capsules administration is discontinued or interrupted for several days at either the 0.4 mg or 0.8 mg dose, therapy should be started again with the 0.4 mg once daily dose.

INSTRUCTIONS:

Dosage as directed by the physician.

Store below 25°C.

Protect from light and moisture.

Keep all medicines out of the reach of children.

To be sold on the prescription of a registered medical practitioner only.

PRESENTATION:

TAMPRO (Tamsulosin HCl) Capsules 0.4mg are available in Alu/Pvc blister pack of 2x10's.

ہدایات:

خوراک ڈاکٹر کی ہدایت کے مطابق استعمال کریں۔

۲۵ ڈگری سینٹی گریڈ سے کم پر رکھیں۔

روشنی اور نمی سے محفوظ رکھیں۔

تمام دوائیں بچوں کی پہنچ سے دور رکھیں۔

صرف رجسٹرڈ ڈاکٹر کے نسخہ پر فروخت کریں۔

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