

PRODUCT MONOGRAPH

Pr[®]TEVA-PINDOLOL

Pindolol

5 mg, 10 mg and 15 mg Tablets

USP

Antihypertensive/Antianginal Agent

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NAME OF DRUG

TEVA-PINDOLOL

Pindolol

5 mg, 10 mg and 15 mg Tablets

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THERAPEUTIC CLASSIFICATION

Antihypertensive/Antianginal Agent

ACTIONS

TEVA-PINDOLOL (pindolol) is a β -adrenergic-receptor-blocking agent which possesses partial agonist activity (intrinsic sympathomimetic activity - I.S.A.). It is used in the treatment of hypertension and/or the prophylaxis of angina pectoris.

Hypertension

The mechanism of the antihypertensive effect of pindolol has not been established. Among the factors that may be involved are:

- a) competitive ability to antagonize catecholamine-induced tachycardia at the β -receptor sites in the heart, thus decreasing cardiac output
- b) a reduction in total peripheral resistance
- c) inhibition of the vasomotor centres
- d) inhibition of renin release by the kidneys

Angina Pectoris

The mechanism of the antianginal effect of pindolol has not been established. Pindolol may reduce the oxygen requirement of the heart at any level of effort by blocking catecholamine-induced increases in the heart rate, systolic blood pressure, and the velocity and extent of myocardial contraction. However, oxygen requirements may be increased by such actions as increases in left ventricular fibre length, end diastolic pressure and the systolic ejection period. When the net effect is beneficial in patients with angina, it manifests itself during exercise or stress by delaying the onset of pain and reducing the incidence and severity of anginal attacks.

In man, orally-administered pindolol is rapidly and almost completely absorbed ($\geq 95\%$) from the gastrointestinal tract. The mean absolute bioavailability after oral administration is about 87-92%. Plasma levels of 10 to 30 nanogram/mL are associated with its therapeutic efficacy. Following single dose administration 5 mg pindolol, the mean maximum plasma concentration (C_{\max}) of pindolol was 33.1 ± 5.2 nanogram/mL (T_{\max} 1-2 h). The elimination rate of pindolol is not dose dependent. The elimination half-life of pindolol is 3 to 4 hours and the drug has a

systemic clearance of between 400 and 500 mL/min. Approximately, 40% of pindolol is bound to plasma proteins. Pindolol is extensively and rapidly distributed throughout the body with a mean volume of distribution of 2-3 L/kg. The elimination kinetics has generally been described as a mono-exponential decay function using one compartment pharmacokinetics.

Pindolol is partially metabolized in the liver with approximately 30 to 40% of an oral dose being excreted unchanged in the urine. The remaining 60 to 70% of pindolol is metabolized in the liver forming inactive metabolites - hydroxylate, which is excreted via kidney and liver as glucuronides and ethereal sulfate. The inactive polar metabolites are excreted out with elimination half-life of 8 h. The fraction eliminated in bile is approximately 6-8%.

Approximately 80% of an oral dose is accounted for in the urine within 24 hours.

Two comparative two-way, single-dose bioavailability studies were performed on TEVA-PINDOLOL (pindolol) 15 mg tablets and VISKEN[®] (pindolol) 15 mg Tablets and on TEVA-PINDOLOL 5 mg Tablets and VISKEN[®] 5 mg Tablets. The pharmacokinetic plasma data (mean \pm standard deviation) calculated for the TEVA-PINDOLOL and VISKEN[®] Tablet formulations is tabulated below:

15 mg Formulation:

	TEVA-PINDOLOL 1 x 15 mg		VISKEN[®] 1 x 15 mg		Percentage of VISKEN[®]
AUC _T * (ng•h/mL)	399	(28)	361	(21)	111
AUC _I * (ng•h/mL)	493	(28)	446	(21)	111
C _{max} * (ng/mL)	68	(24)	63	(16)	109
T _{max} ⁺ (h)	1.16	(0.54)	1.34	(0.60)	-
T _{1/2} ⁺ (h)	4.9	(1.9)	4.9	(1.6)	-
* Geometric means (CV)					
+ Arithmetic means (SD)					

5 mg Formulation:

	TEVA-PINDOLOL 2 x 15 mg		VISKEN[®] 2 x 15 mg		Percentage of VISKEN[®]
AUC _T * (ng•h/mL)	195	(22)	195	(24)	100
AUC _I * (ng•h/mL)	214	(22)	219	(25)	98
C _{max} * (ng/mL)	37	(16)	37	(22)	100
T _{max} ⁺ (h)	1.9	(0.38)	1.31	(0.44)	-
T _{1/2} ⁺ (h)	3.03	(0.65)	3.20	(0.94)	-
* Geometric means (CV)					
+ Arithmetic means (SD)					

INDICATIONS

Hypertension

TEVA-PINDOLOL (pindolol) is indicated for the treatment of mild to moderate hypertension. Pindolol is usually used in combination with other drugs, particularly a thiazide diuretic.

However, it may be used alone as an initial agent in those patients in whom, in the judgment of the physician, treatment should be started with a β -blocker rather than a diuretic.

The combination of pindolol with a diuretic and/or peripheral vasodilator has been found to be compatible and generally more effective than pindolol alone. Limited experience with other antihypertensive agents, including methyldopa, has not shown evidence of incompatibility with pindolol.

TEVA-PINDOLOL is not recommended for the emergency treatment of hypertensive crises.

Angina Pectoris

TEVA-PINDOLOL is indicated for the prophylaxis (prevention) of angina pectoris.

CONTRAINDICATIONS

TEVA-PINDOLOL (pindolol) should not be used in the presence of:

1. sinus bradycardia (< 45 -50 beats/min)
2. second and third degree A-V block
3. right ventricular failure secondary to pulmonary hypertension
4. congestive heart failure (see **WARNINGS**)
5. cardiogenic shock
6. anesthesia with agents which produce myocardial depression, e.g. ether
7. bronchospasm, including bronchial asthma or severe chronic obstructive pulmonary disease (see **PRECAUTIONS**)
8. hypersensitivity to pindolol, to any of the excipients or cross-sensitivity to other beta blockers
9. Prinzmetal's angina (variant angina)
10. sick sinus syndrome
11. severe peripheral arterial circulatory disturbances
12. untreated pheochromocytoma

WARNINGS

a) Cardiac Failure:

Special caution should be exercised when administering TEVA-PINDOLOL (pindolol) to patients with a history of heart failure. Sympathetic stimulation is a vital component supporting circulatory function in congestive heart failure, and inhibition with β -blockade

always carries the potential hazard of further depressing myocardial contractility and precipitating cardiac failure. Pindolol may reduce but does not abolish the inotropic action of digitalis on the heart muscle. However, the positive inotropic action of digitalis may be reduced by the negative inotropic effect of pindolol when the two drugs are used concomitantly. The effects of β -blockers and digitalis are additive in depressing AV conduction. In patients without a history of cardiac failure, continued depression of the myocardium over a period of time can, in some cases, lead to cardiac failure. Therefore, at the first sign or symptom of impending cardiac failure, patients should be fully digitalised and/or given a diuretic and the response observed closely. If cardiac failure continues, despite adequate digitalization and diuretic therapy, TEVA-PINDOLOL should be immediately withdrawn.

b) Abrupt Cessation of Therapy with TEVA-PINDOLOL:

Patients with angina should be warned against abrupt discontinuation of TEVA-PINDOLOL (pindolol). There have been reports of severe exacerbation of angina, and of myocardial infarction or ventricular arrhythmias occurring in patients with angina pectoris, following abrupt discontinuation of β -blocker therapy. The last two complications may occur with or without preceding exacerbation of angina pectoris. Therefore, when discontinuation of TEVA-PINDOLOL is planned in patients with angina pectoris, the dosage should be gradually reduced over a period of about two weeks and the patient should be observed carefully. The same frequency of administration should be maintained. In situations of greater urgency, TEVA-PINDOLOL therapy should be discontinued stepwise under very close observation.

If angina markedly worsens or acute coronary insufficiency develops, it is recommended that treatment with TEVA-PINDOLOL be reinstituted promptly, at least temporarily.

- c) Various skin rashes and conjunctival xerosis have been reported with β -blockers, including pindolol. A severe oculo-muco-cutaneous syndrome, whose signs include conjunctivitis sicca and psoriasiform rashes, otitis, and sclerosing serositis, has occurred with the chronic use of one β -adrenergic-blocking agent (practolol). This syndrome has not been observed with pindolol. However, physicians should be alert to the possibility of such reactions and should discontinue treatment in the event that they occur. A switch to another therapeutic agent might be advisable.
- d) Sinus bradycardia may occur with the use of TEVA-PINDOLOL due to unopposed vagal activity remaining after blockade of β_1 -adrenergic-receptors. However, due to its intrinsic sympathomimetic activity (ISA) pindolol causes less bradycardia at rest than some other β -adrenergic blocking agents. If excessive bradycardia occurs the dosage of TEVA-PINDOLOL should be reduced.
- e) In patients with thyrotoxicosis, possible deleterious effects from long-term use of pindolol have not been adequately appraised. β -blockade may mask the clinical signs of continuing hyperthyroidism or complications, and give a false impression of improvement. Therefore, these patients should be carefully monitored for thyroid function. Abrupt withdrawal of

pindolol may be followed by an exacerbation of the symptoms of hyperthyroidism, including thyroid storm.

- f) **Co-medication with calcium channel blockers:** Owing to the danger of cardiac arrest, a calcium channel blocker of the verapamil type must not be administered intravenously to a patient already receiving treatment with a beta-blocker.
- g) **Psoriasis:** Since beta-blockers may aggravate psoriasis, TEVA-PINDOLOL should only be prescribed after careful consideration of benefits and risks in patients with history of psoriasis.

PRECAUTIONS

- a) Caution should be exercised in patients prone to non-allergic bronchospasm (e.g. chronic bronchitis, emphysema) since TEVA-PINDOLOL (pindolol) may block bronchodilation produced by endogenous and exogenous catecholamine stimulation of β -receptors.
- b) TEVA-PINDOLOL should be administered with caution to patients with allergic rhinitis prone to bronchospasm.

There may be increased difficulty in treating an allergic type reaction in patients on β -blockers.

In these patients, the reaction may be more severe due to pharmacologic effects of the β -blockers and problems with fluid changes. Epinephrine should be administered with caution since it may not have its usual effects in the treatment of anaphylaxis. On the one hand, larger doses of epinephrine may be needed to overcome the bronchospasm, while on the other, these doses can be associated with excessive alpha-adrenergic stimulation with consequent hypertension, reflex bradycardia and heart block and possible potentiation of bronchospasm. Alternatives to the use of large doses of epinephrine include vigorous supportive care such as fluids and the use of β -agonists, including parenteral salbutamol or isoproterenol, to overcome bronchospasm and norepinephrine to overcome hypotension.

- c) TEVA-PINDOLOL should be administered with caution to patients subject to spontaneous hypoglycemia, or to diabetic patients (especially those with labile diabetes) who are receiving insulin or oral hypoglycemic agents. β -adrenergic-blockers may mask the premonitory signs and symptoms (e.g. palpitations, tachycardia, tremor) of acute hypoglycemia whereas sweating is not inhibited. The concurrent use of beta-blockers and antidiabetic medication should always be monitored to confirm that diabetic control is well maintained.
- d) TEVA-PINDOLOL dosage should be individually adjusted when used concomitantly with other antihypertensive agents. (See **DOSAGE AND ADMINISTRATION**)

- e) Patients receiving catecholamine-depleting drugs, such as reserpine or guanethidine, should be closely monitored because the added β -adrenergic-blocking action of pindolol may produce an excessive reduction of sympathetic activity. TEVA-PINDOLOL should not be combined with other β -blockers.
- f) Appropriate laboratory tests should be performed at regular intervals during long-term treatment.
- g) The management of patients being treated with β -blockers and undergoing elective or emergency surgery is controversial. Although β -adrenergic-receptor-blockade impairs the ability of the heart to respond to β -adrenergically-mediated reflex stimuli, abrupt discontinuation of therapy with pindolol may be followed by severe complications (see **WARNINGS**).

Some patients receiving β -adrenergic-blocking agents have been subject to protracted severe hypotension during anesthesia. Difficulty in re-starting and maintaining the heart beat has also been reported.

For these reasons, in patients with angina, undergoing elective surgery, TEVA-PINDOLOL should be withdrawn gradually following the recommendation given under Abrupt Cessation of Therapy (see **WARNINGS**). According to available evidence, all clinical and physiological effects of β -blockade are not longer present 48 hours after cessation of medication.

In emergency surgery, since TEVA-PINDOLOL is a competitive inhibitor of β -adrenergic-receptor agonists, its effects may be reversed by sufficient doses of such agonists as isoproterenol or levarterenol.

h) Impaired Renal or Hepatic Function:

β -blocking agents should be used with caution in patients with impaired hepatic or renal function. Poor renal function has only minor effects on pindolol clearance, but poor hepatic function may cause blood levels of pindolol to increase substantially.

In patients with severe renal impairment, further impairment of renal function has been only rarely observed during therapy with pindolol.

i) Anaphylactic Reaction:

Anaphylactic reactions precipitated by other agents may be particularly severe in patients taking beta-blockers, especially non-selective beta-blockers, and may be resistant to normal doses of adrenaline. Whenever possible, beta-blockers should be avoided in patients who are at increased risk for anaphylaxis.

j) Pheochromocytoma:

If patients with pheochromocytoma are treated with a beta-blocker, an alpha-blocker should always be co-administered. (See **CONTRAINDICATIONS**)

k) **Usage in Pregnancy:**

Since pindolol has not been studied in human pregnancy, the drug should not be given to pregnant women. The use of any drug in patients of child-bearing potential requires that the anticipated benefit be weighed against possible hazards. Pindolol crosses the placental barrier.

l) **Lactating Women:**

Pindolol passes in small quantities into breast milk.

m) **Fertility:**

In rats, pindolol did not cause any adverse effects on fertility or reproductive performance at a dose of 10 mg/kg, which is 17-times the human dose. While effects in animals are not always predictive of human effects, at dose levels of 30 mg/kg and greater, female rats were observed to mate less frequently than untreated animals. (See **TOXICOLOGY**)

n) **Usage in Children:**

Since there is no experience with pindolol in the treatment of pediatric age groups, TEVA-PINDOLOL is not indicated for pediatrics.

o) **Usage in Geriatric Patients:**

No evidence exists that geriatric patients require different dosages; however these patients should be treated cautiously. An excessive decrease in blood pressure or pulse rate may reduce blood supply to vital organs to inadequate levels.

p) Because dizziness or fatigue may occur during initiation of treatment with β -adrenoreceptor blocking drugs, patients driving vehicles or operating machinery should exercise caution until they have determined their individual response to treatment.

ADVERSE REACTIONS

Cardiovascular

Congestive heart failure (see **WARNINGS**), severe bradycardia (see **WARNINGS**), may occur. Syncope, lightheadedness, postural hypotension. Lengthening of PR interval, second degree AV block, palpitation, chest pains, cold extremities, Raynaud's phenomenon, claudication, hot flushes. Very rarely arrhythmia, coronary insufficiency.

Central Nervous System

Insomnia, nightmares, vivid dreams, fatigue, drowsiness, weakness, dizziness, vertigo, tinnitus, headache, mental depression, nervousness. The following adverse reactions have been reported rarely: aggressiveness, motor disorders, confusion.

Gastrointestinal

Diarrhea, constipation, flatulence, heartburn, nausea and vomiting, abdominal pain and dry mouth.

Respiratory

Shortness of breath and/or dyspnea, wheezing, bronchospasm.

Allergic, Dermatological (see WARNINGS)

Exanthema, sweating, pruritis, psoriasiform rash.

Eyes

Itching, burning, grittiness, dryness.

Miscellaneous

Muscle cramps, appetite stimulation, weight gain, urinary frequency.

Clinical Laboratory

On rare occasions, changes in the following parameters were noted: elevated transaminases, alkaline phosphatase, LDH, serum uric acid; reduced bilirubin.

Post-Market Adverse Drug Reactions

The following adverse drug reactions are, in most cases, mild and transient in nature and necessity for interruption of pindolol therapy is rarely observed (see **WARNINGS and PRECAUTIONS**).

These adverse drug reactions (Table 1) have been derived from post-marketing experience with pindolol. Because these reactions are reported voluntarily from a population of uncertain size, it is not possible to reliably estimate their frequency which is therefore categorized as not known. Adverse drug reactions are listed according to system organ classes in MedDRA. Within each system organ class, ADRs are presented in order of decreasing seriousness.

Table 1 Adverse Drug Reactions (frequency not known)

Psychiatric disorders

Sleep disorders, depression, hallucinations

Nervous system disorders

Tremor, dizziness, headache

Cardiac disorders

Bradycardia, conduction disorder, cardiac failure

Vascular disorders

Hypotension, symptoms of peripheral vascular disorders (peripheral coldness), Raynaud's-like symptoms

Respiratory, thoracic and mediastinal disorders

Bronchospasm, dyspnea

Gastrointestinal disorders

Gastrointestinal disorders (nausea, vomiting, abdominal pain and diarrhea)

Skin and subcutaneous tissue disorders

Skin reaction, hyperhidrosis, worsening of psoriasis

Musculoskeletal and connective tissue disorders

Muscle cramps

General disorders and administration site conditions

Fatigue

DRUG INTERACTIONS

Table 2 Established or Potential Drug-Drug Interactions

Product	Ref	Effect	Clinical Comment
Monoamine oxidase (MAO) inhibitors	C, T	Combining these medications may increase the risk of hypotension, orthostasis, bradycardia, and heart failure due to excessive reduction of sympathetic activity. Possibly significant hypertension may theoretically occur up to 14 days following discontinuation of the MAO inhibitor.	Monoamine oxidase (MAO) inhibitors may potentiate the pharmacologic effects of beta-blockers, which are thought to competitively antagonize catecholamines at cardiac and other peripheral adrenergic neurons. Concurrent use with beta-blockers is not recommended.
Antidiabetic Agents	T	Beta-blockers may interfere with the usual hemodynamic response to hyperglycaemia and produce a rise in blood pressure associated with severe bradycardia.	Beta-blockade reduces the release of insulin in response to hyperglycemia; therefore, it may be necessary to adjust the dose of antidiabetic drugs. Beta-blockers should be avoided in unstable diabetic patients (patients who experience wide and unpredictable fluctuations of blood glucose values and/or difficulty in stabilizing blood glucose levels) prone to episodes of hypoglycemia (see WARNINGS and PRECAUTIONS).
Calcium-channel blocking agents	CT (Carruthers 1991)	Because of their potential effect on the cardiac conduction system and contractility, the i.v. route must be avoided. Oral treatment, if judged absolutely necessary, requires careful monitoring, especially when the beta-blocker is combined with a verapamil-type calcium antagonist.	Severe reduction in blood pressure and heart failure upon the concomitant administration of dihydropyridine derivatives such as nifedipine with pindolol in patients with latent cardiac insufficiency is possible.
Anti-adrenergic agents	T	Antihypertensive effect of alpha-adrenergic blockers such as guanethidine, betanidine, reserpine, alpha-methyldopa or clonidine may be potentiated by beta-blockers, which may lead to postural hypotension.	When therapy is discontinued in patients receiving a beta-blocker and clonidine concurrently, the beta-blockers should be gradually discontinued several days before clonidine is discontinued, in order to reduce the potential risk

Product	Ref	Effect	Clinical Comment
			of a clonidine withdrawal hypertensive crisis (rebound effect). Monitoring of blood pressure is recommended during the anti-adrenergics withdrawal.
Non-steroidal anti-inflammatory drugs (NSAIDs)	T	Concomitant administration of non-steroidal anti-inflammatory drugs including COX-2 inhibitors with a beta-blocker, may decrease its antihypertensive effect, possibly as a result of the inhibition of renal prostaglandin synthesis and sodium and fluid retention caused by NSAIDs.	Anti-hypertensive effects of beta-blockers may be decreased by non-steroidal anti-inflammatory drugs, which may lead to uncontrolled hypertension. Monitoring is required.
Phenothiazines	CT	Concurrent administration of pindolol and thioridazine is reported to result in a moderate increases in the serum levels of thioridazine and two of its metabolites, as well as higher than expected serum pindolol levels.	Concurrent use with beta-blockers with phenothiazines results in an increased plasma concentration of either drug, which may lead to hypotension, ventricular tachycardia, and pigmentary retinopathy. Monitoring is required.
Sympathomimetic drugs	T	Concomitant administration of sympathomimetic drugs such as adrenaline, noradrenaline, isoprenaline, ephedrine, phenylephrine phenylpropanolamine, or xanthine derivatives with a non-selective beta-blocker may enhance the pressor response resulting in severe hypertension due to antagonistic effects.	Pindolol may antagonize the effects of sympathomimetic drugs and xanthine derivatives which may lead to severe hypertension. Monitoring is required.
Anesthetic agents	CT	Beta-blockers and certain anaesthetics may be additive in their cardio-depressant effect and may lead to protracted severe hypotension (see WARNINGS and PRECAUTIONS).	Anaesthetic agents causing myocardial depression, such as cyclopropane and trichloroethylene, are best be avoided.
Anti-arrhythmic agents	CT	Concomitant administration of beta-blockers with class I anti-arrhythmic agents such as disopyramide, tocainide, flecainide or amiodarone have a potentiating effect on atrial-conduction time and induce negative inotropic effect, which may lead to myocardial depression, cardiac failure,	Although this potentiation effect is weak for pindolol, the possibility of interactions with anti-arrhythmic agents cannot be eliminated. Monitoring is required.

Product	Ref	Effect	Clinical Comment
		hypotension, bradycardia, AV block and asystole.	
Digitalis glycosides	T	Beta-blockers and digitalis glycosides may be additive in their depressant effect on myocardial conduction, particularly through the atrioventricular node.	Concomitant administration of digitalis glycosides may induce serious bradycardia or heart block and thus should be avoided.
Ergot alkaloid	T	Administration with beta-blockers may enhance the vasoconstrictive effect of ergot alkaloids.	Concomitant administration with beta-blockers with ergot alkaloid may enhance the vasoconstriction, which leads to hypertension.
Cimetidine	CT	Cimetidine is a moderate inhibitor of multiple cytochrome enzymes such as CYP2D6, CYP3A4, CYP2C19, CYP2E1, CYP2C9, and CYP1A2. Concomitant administration of cimetidine may inhibit the hepatic metabolism of pindolol resulting in increased plasma concentrations of pindolol, which may lead to hypotension.	Monitoring is required.
Fingolimod	CT	Bradycardia	Concomitant use of fingolimod with beta blockers may potentiate bradycardic effects and is not recommended. Where such coadministration is considered necessary, appropriate monitoring at treatment initiation, i.e. at least overnight monitoring, is recommended.

Legend: C = Case Study; CT = Clinical Trial; T = Theoretical

SYMPTOMS AND TREATMENT OF OVERDOSAGE

An overdosage of beta-blocker may lead to pronounced bradycardia, hypotension, cardiac failure, cardiogenic shock, conduction abnormalities, cardiac arrest, dyspnea, bronchospasm, vomiting, hypoglycemia, depressed levels of consciousness, generalized convulsions, coma and death. In rare circumstances, overdose of beta-blockers with intrinsic sympathomimetic activity (ISA), like TEVA-PINDOLOL, may present with tachycardia and hypertension. Concomitant ingestion of alcohol, antihypertensives, antidepressants, or antiarrhythmic may aggravate the signs and symptoms of overdose.

If overdosage occurs, in all cases therapy with TEVA-PINDOLOL (pindolol) should be discontinued and the patient observed closely. If required, the following therapeutic measures are suggested:

1. Bradycardia: atropine or another anticholinergic drug.
2. Heart block (second or third degree): isoproterenol or transvenous cardiac pacemaker.
3. Congestive heart failure: conventional therapy.
4. Hypotension: (depending on associated factors) epinephrine rather than isoproterenol or norepinephrine may be useful in addition to atropine and digitalis (see **PRECAUTIONS** concerning the use of epinephrine).
5. Bronchospasm: aminophylline or isoproterenol.
6. Hypoglycemia: intravenous glucose.

It should be remembered that pindolol is a competitive antagonist of isoproterenol and hence large doses of isoproterenol can be expected to reverse many of the effects of excessive doses of TEVA-PINDOLOL. However, the complications of excess isoproterenol should not be overlooked.

DOSAGE AND ADMINISTRATION

a) Hypertension

TEVA-PINDOLOL (pindolol) is usually used in conjunction with other antihypertensive agents, particularly a thiazide diuretic but may be used alone (see **INDICATIONS**).

TEVA-PINDOLOL should be taken with meals.

The dosage of TEVA-PINDOLOL must always be adjusted to the individual requirements of the patients in accordance with the following guidelines:

TEVA-PINDOLOL therapy should be initiated with doses of 5 mg in the morning with breakfast and 5 mg with the evening meal. If an adequate response is not achieved after one to two weeks, the dose should be increased to 10 mg twice a day.

If after one to two additional weeks an adequate response is not observed, dosage may be increased to 15 mg twice a day (30 mg/day).

Doses greater than 30 mg daily must be given on a t.i.d. schedule.

Patients who show a satisfactory response to TEVA-PINDOLOL at daily doses of 10 to 20 mg may be maintained by giving the required total dose once daily in the morning with breakfast.

The usual maintenance dose is within the range of 15 to 45 mg daily which should not be exceeded. However, during long-term therapy, some patients may be maintained on smaller doses of TEVA-PINDOLOL.

b) **Angina Pectoris**

The dosage of TEVA-PINDOLOL must always be adjusted to the individual requirements of the patient.

In angina, TEVA-PINDOLOL should be administered on a three or four times per day dosing regimen. TEVA-PINDOLOL therapy should be initiated with doses of 5 mg three times a day taken with meals. If after one to two weeks an adequate response is not observed, dosage may be increased. The usual maintenance dose is 15 mg up to the maximum of 40 mg per day.

c) **Special Populations**

Patients with impaired renal function/hepatic function

Patients with impaired renal or hepatic function may usually be treated with normal doses. Only in severe cases may a reduction of the daily dose be necessary (see **PHARMACOLOGY – Special Populations**).

Pediatric patients

Since the efficacy and safety of TEVA-PINDOLOL has not been established in children, TEVA-PINDOLOL is not indicated for pediatrics.

Geriatric patients

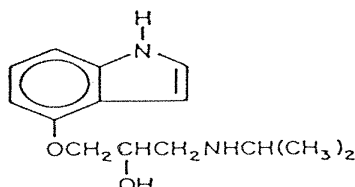
No evidence exists that geriatric patients require different dosages; however these patients should be treated cautiously (see **PRECAUTIONS** and **PHARMACOLOGY – Special Populations**).

PHARMACEUTICAL INFORMATION

Trade Name: TEVA-PINDOLOL

Proper Name: Pindolol

Structural Formula:



Molecular Formula: C₁₄H₂₀N₂O₂

Molecular Weight: 248.3

Chemical Name: 1-(Indol-4-yloxy)-3-(isopropylamino)-2-propanol

Description: Pindolol is a white to off-white crystalline powder having a faint odour. It is practically insoluble in water, slightly soluble in methanol and very slightly soluble in chloroform.

AVAILABILITY

- 5 mg: White, round, flat with bevelled edge, compressed tablets; on one side single scored, novo over 5 engraved on the reverse. Available in bottles of 100 and 500.
- 10 mg: White, round, flat with bevelled edge compressed tablets, on one side single scored, novo over 10 engraved on the reverse. Available in bottles of 100 and 500.
- 15 mg: White, round, flat with bevelled edge compressed tablets, on one side single scored, novo over 15 engraved on the reverse. Available in bottles of 100 and 500.

Non-medicinal Ingredients:

5 mg, 10 mg and 15 mg tablets: colloidal silicon dioxide, magnesium stearate, microcrystalline cellulose, pregelatinized starch, sodium lauryl sulfate and sodium starch glycolate.

STORAGE RECOMMENDATIONS

TEVA-PINDOLOL (pindolol) tablets should be protected from light and stored at 15-30°C in well closed containers.

PHARMACOLOGY

Effects on the Cardiovascular System

Pindolol, in the non-anesthetized dog, produced a 70% inhibition of tachycardia and changes in blood pressure induced by isoproterenol at doses of 0.05 mg/kg i.v. and 2 mg/kg i.v. respectively. Complete antagonism was observed following pindolol doses of 0.1 to 5 mg/kg i.v. In the anesthetized dog, 0.2 to 2.0 mg/kg i.v. produced dose dependent decreases in blood pressure; heart rate changes were unrelated to dose and were reduced by 12% after a dose of 0.2 mg/kg i.v. and 4% after i.v. injection of 2 mg/kg.

In the anesthetized dog, 0.2 to 1 mg/kg i.a. antagonized the vasodilation induced by isoproterenol, whereas transient 25 to 40% reductions in vascular resistance were observed after intra-arterial doses of 50 and 200 mg/kg. Intravenous doses of 2 mg/kg of pindolol elicited peripheral vasodilation and an associated reduction in total peripheral resistance.

In vivo studies on the guinea pig atrium showed that pindolol produced dose dependent antagonism of epinephrine-induced positive inotropy and chronotropy.

In five healthy volunteers given a single oral dose of 10 mg of pindolol, antagonism of isoproterenol-induced tachycardia and changes in blood pressure and heart rate were observed 30 minutes after ingestion and persisted for 24 hours.

In ten hypertensive patients receiving pindolol for 16 months in divided doses of 20 to 40 mg, blood pressure reduction was associated with statistically significant reduction in forearm and total systemic vascular resistance at rest and during stress testing. Venous tone was significantly reduced during and after exercise. No significant change was reported in cardiac output following prolonged use (see **ACTIONS**).

Pindolol has little membrane stabilizing activity being approximately 1/12 that of quinidine in prolonging the relative refractory period of cardiac cells in the isolated guinea pig atrium. A concentration of up to 5% pindolol was devoid of local anesthetic effects when applied to the cornea of the eye.

Pindolol possesses partial agonist (intrinsic sympathomimetic) activity. Long-lasting increases in myocardial activity manifested by positive chronotropic actions were observed following i.v. infusions of pindolol at doses of 0.16 mcg/kg to 2.5 mg/kg in reserpinized, adrenalectomized and vagotomized cat.

Pindolol decreases the basal rate of myocardial oxygen consumption and blocks increases mediated by increased sympathetic nervous system activity.

Pindolol has antiarrhythmic activity. At doses of 8 mg/kg in the anesthetized dog, pindolol increased the dose of ouabain required to produce ventricular arrhythmia. In guinea pigs and dogs, it delayed the onset of ouabain-induced ventricular arrhythmia and in the dog produced reversion to sinus rhythm.

Pindolol has been reported to reduce plasma renin activity in some patients. However, plasma renin may remain unchanged or increase following treatment. There does not appear to be any significant relationship between the antihypertensive activity of pindolol and changes in plasma renin activity.

Effects on Pulmonary Function

In a study of 58 hypertensive patients with normal respiratory function who received oral doses of 15, 30, or 60 mg of pindolol, no significant changes were observed in forced expiratory volume (FEV), maximum voluntary ventilation rate, maximum expiratory flow rate and maximum mid-expiratory flow rate.

Decreased FEV₁ has, however, been reported in other studies.

Other Effects

Electroencephalographic changes, following oral doses of 5 and 10 mg in healthy volunteers, consisted of theta and fast beta and decreases in alpha activity. In rats given 5.2 mg/kg s.c., pindolol blocked tetrabenzine-induced ptosis but not catalepsy. In mice at doses of 1 to 30 mg/kg i.v., pindolol antagonized reserpine-induced hypothermia.

Special populations

Geriatrics

The elderly population may show higher plasma concentrations of pindolol as a combined result of a decreased metabolism of the drug in elderly population, a decreased hepatic blood flow and a decreased renal elimination.

Pregnancy

The elimination half-life of pindolol does not differ significantly between pregnant and non-pregnant patients. (See **PRECAUTIONS**)

Transplacental distribution of pindolol is not stereoselective. Pregnancy may alter the pharmacokinetic disposition of pindolol, suggesting an increase in the distribution volume and total clearance.

Patients with hepatic / renal impairment

Patients with impaired renal or hepatic function may usually be treated with normal doses. Only in severe cases may a reduction of the daily dose be necessary. The plasma half-life of pindolol is increased up to 11.5 hours, depending on severity, in patient with renal impairment and is increased up to 30 hours, depending on severity, in patients with liver cirrhosis.

TOXICOLOGY

a) Acute Toxicity

Species	Route	LD ₅₀ (mg/kg)
Mouse	i.v.	29 ± 1.2
Mouse	p.o.	200 ± 22
Rat	i.v.	35 ± 1.7
Rat	p.o.	260 ± 36
Rabbit	i.v.	10 ± 0.9
Rabbit	p.o.	650 ± 102
Dog	p.o.	≥ 30

b) Subacute:

Species/ Strain	Sex M/F	No. of groups	N/group	Dose mg/kg/day	Route	Duration	Toxic effects
Rat	40/40	4	10M/10F	0, 16, 66, 246	Oral	13 weeks	At 246 mg/kg there was a mortality rate of 20%. Arrest of spermatogenesis in males and hypoplastic uteri in females were observed at doses of 66 and 246 mg/kg. Doses of 16, 66 and 246 mg/kg slightly to moderately increased SGPT levels, and reduced food intake, the efficiency of food utilization and organ and body weights. Treated animals had a slightly higher incidence of infection than controls. Granular inclusions in liver and adrenal cells and increased numbers of fat droplets in renal tubule cells were seen at doses of 246 mg/kg. Similar but less prominent changes were found at 66 mg/kg. There were isolated incidences of thymus involution, contraction of the seminal vesicles and prostatic atrophy. Green discolouration of the urine was observed.
Rats	40/40	4	10M/10F	0, 5, 25, 130	Oral	26 weeks	At 130 mg/kg/day, decreased body weight and cyanosis were observed.
Dogs/ Beagle	8/8	4	2M/2F	0, 5, 20, 80 (6 days/week)	Oral	13 weeks	At 80 mg/kg/day, convulsions, gastrointestinal disturbances, mydriasis, erythema secondary to cutaneous vasodilation were observed. Food intake and body weight were reduced.

Species/ Strain	Sex M/F	No. of groups	N/group	Dose mg/kg/day	Route	Duration	Toxic effects
Dogs/ Beagle	12/12	4	3M/3F	0, 5, 15, 45	Oral	26 weeks	At 45 mg/kg, the mortality rate was 50%. Hepatocyte swelling, and the presence of intracellular hyaline droplets and lipochrome pigment in hepatocytes and Kupffer cells were seen at 15 and 45 mg/day and a few single sporadic degenerating liver cells were observed. Green discolouration of the urine was seen at 15 and 45 mg/kg/day. One dog in each group given 5, 15, and 45 mg/kg/day showed transient increases in alkaline phosphatase. In the 45 mg/kg/day group, convulsions, gastrointestinal disturbances, arrest of spermatogenesis, weight loss and reduced adreno-cortical lipids were observed.
Rats	30/30	3	10M/10F	0, 1, 3	i.v.	4 weeks	None
Dogs/ Beagle	2/2 4/4	1 2	2M/2F 2M/2F	0 1.5	i.v.	4 weeks	1.5 mg/kg: Erythema secondary to cutaneous vasodilation
Rats	5/5 10/10	1 1	5M/5F 10M/10F	0 5	i.m.	4 weeks	5 mg/kg: Slight irritant effect at injection site

c) Chronic Toxicity:

Species/ Strain	Sex M/F	No. of groups	N/group	Dose mg/kg/day	Route	Duration (years)	Toxic effects
Rats	120/120	4	30M/30F	0, 2, 14, 98	Oral	2	Green discolouration of the urine at 98 m/kg. At 2, 14 and 98 mg/kg, deposition of a greenish brown pigment in Kupffer cells of the liver.
Dogs/ Beagle	16/16	4	4M/4F	0, 2, 6, 18	Oral	2	Tachycardia of 1 week duration. Erythema secondary to cutaneous vasodilatation which was not dose dependent. Emesis and soft stools.
Monkey/ Rhesus	9/9	3	3M/3F	0, 2.5, 25	Oral	1	At 2.5 mg/kg, heart rate was slowed 15 to 20%. Bradycardia was seen at 25 mg/kg. Green discolouration of the urine at 25 mg/kg.

d) Disposition of Pigment

Oral administration of pindolol to rats at a dose of 200 mg/kg/day for 26 weeks resulted in the deposition of a melanin-like pigment in the liver, spleen, adrenal gland and subcutaneous tissue. Partial disappearance of this pigment from Kupffer cells in the liver occurred within four weeks following discontinuation of pindolol.

In dogs given oral doses of 5, 15, and 45 mg/kg/day for 26 weeks, dose related increases in hepatocyte lipid content were observed.

However, despite the pigment deposition and increased lipid content, all tests done for hepatic, splenic and adrenal function were normal. The significance of pigment and lipid changes is unknown.

e) Teratology and Reproduction Studies

i. *Teratology*

The parameters studied in the rat and rabbit teratology studies were the following: total number of pregnancies, implantations, viable fetuses, dead fetuses, total prenatal deaths, abnormal fetuses in % of living fetuses.

Rat:

Doses of 30 and 100 mg/kg were administered orally to groups of 20 pregnant rats (Sandoz Closed Strain) on days 7-16 of gestation. Treatment with pindolol did not adversely affect any of the parameters studied.

Rabbit:

Doses of 8, 23 and 80 mg/kg were administered orally to groups of respectively, 13, 16 and 15 pregnant rabbits (Swiss Hare Strain) on days 6-18 of gestation. None of the parameters studied was significantly affected.

ii. *Reproduction*

Rat: Doses of 10, 30 and 100 mg/kg were administered orally to groups of 15 male (Sandoz Closed Strain) and 30 female (Carworth Wistar CFE Strain) rats. Males were treated for 70 days prior to and during the mating period. The females were treated for up to 15 days prior to mating, during mating, and throughout the gestation and lactation period to 21 days postpartum, with an interim sacrifice at Day 13 of gestation.

Spermatogenesis and fertility were reduced at doses of 30 but not 100 mg/kg/day. Tubular atrophy in the testes was found in male rats treated with doses of 30 and 100 mg/kg/day.

There was significantly greater mortality in the offspring of females treated with 100 mg/kg/day in the first four-day postpartum period and in pups of females receiving 30 mg/kg/day during the 4 to 21-day postpartum interval. This increased mortality may be

consequence of deficits in maternal rearing behaviour, inhibition of lactation or the presence of the drug in maternal milk.

f) Carcinogenicity Studies

Mouse: Pindolol was administered to 50 male and 50 female mice (Sandoz OFI Strain) at dietary levels of approximately 124 mg/kg/day for 82 weeks, with an equal number of mice serving as controls. The incidence of nodules and masses observed at necropsy were comparable in the treated and control groups. This strain of mice was previously shown to be susceptible to chemical carcinogenesis.

Rat: Pindolol was administered to 50 male and 50 female rats (Sandoz OFA Strain) at a mean dose of 50 mg/kg/day for 83 weeks. A similar group of 100 rats served as a control. Mortality and incidence of tumor were comparable in the treated and untreated groups. This strain of rat was previously shown to be susceptible to chemically (2AAF) induced carcinogenesis.

REFERENCES

1. Aellig WH. β -adrenoceptor blocking activity and duration of action of pindolol and propranolol in healthy volunteers. *Br J Clin Pharmacol* 3: 251-257, 1976.
2. Alexandre JM, Menard J, Chevillard C, and Schmitt H. Increased plasma renin activity induced in rats by physostigmine and effects of α - and β -receptors blocking drugs thereon. *Eur J Pharmacol* 12 :127-131, 1970.
3. Arstila M, Kallio V, and Wendelin H. Propranolol and LB 46 (prinodolol) in angina pectoris. A comparative long-term ergometric study. *Ann Clin Res* 5: 91-100, 1973.
4. Atterhog JH, Duner H, and Pernow B. Experience with pindolol, a β -receptor blocker, in the treatment of hypertension. *Am J Med* 60: 872-876, 1976.
5. Binak K, Sirmaci N, Tavsanoğlu S, and Harmanci N. β -adrenergic blocker LB-46 and isosorbide dinitrate synergism in angina pectoris. *New Istanbul Contr Clin Sci* 11: 8-23, 1974.
6. Bjerle P and Jacobsson KA. Pindolol (Visken*) and alprenolol (Aptin) in angina pectoris: a comparative clinical study with special reference to ergometer working capacity tests. *Curr Ther Res* 16: 101-116, 1974.
7. Buhler FR, Marbet G, Patel U, and Burkart F. Renin-suppressive potency of various β -adrenergic blocking agents at supine rest and during upright exercise. *Clin Sci & Molecular Med* 48 Suppl. 2: 61s-64s, 1975.
8. Clark BJ. Pharmacology of β -adrenoceptor blocking agents. In β -adrenoreceptor Blocking Agents: The pharmacological basis of clinical use. Saxena PR, Forsyth RP (ed), Amsterdam, Holland Publishing Co., 1976, 45-76.
9. Clark B and Saameli K. Pharmacological properties of a new β -receptor blocking agent. *Triangle* 2: 300-308, 1970.
10. Cocco G, Strozzi C, Chu D, Amrein R, and Castagnoli E. Therapeutic effects of pindolol and nifedipine in patients with stable angina pectoris and asymptomatic resting ischemia. *Eur J Card* 10: 59-69, 1979.
11. Frishman W, Kostis J, Strom J, Hossler M, Elkayam U, Goldner S, Silverman R, Davis R, Weinstein J, and Sonnenblick E. Clinical pharmacology of the new β -adrenergic blocking drugs. Part 6. A comparison of pindolol and propranolol in treatment of patients with angina pectoris. The role of intrinsic sympathomimetic activity. *American Heart Journal* 98: 526-535, 1979.
12. Frithz G and Nordgren L. Pindolol and alprenolol in angina pectoris: a comparative clinical study. *Curr Ther Res* 17: 133-138, 1975.

13. Giudicelli JF, Schmitt H, and Boissier JR. Studies on d1-4-(2-hydroxy-3-isopropyl aminopropoxy)-indole (LB-46), a new potent β -adrenergic blocking drug. *J Pharmacol Exp Ther* 168: 116-126, 1969.
14. Gugler R, Bodem G, and Dengler HJ. Pharmacokinetics of β -blocking agents in relation to their anti-hypertensive effect. In: 4th Meeting of the International Society of Hypertension. Sydney, Australia, Feb. 24-26, 1976. The International Society of Hypertension and The National Heart Foundation of Australia, 1976, p55.
15. Gugler R, Herold W, and Dengler HJ. Pharmacokinetics of pindolol in man. *Eur J Clin Pharmacol* 7: 17-24, 1974.
16. Javor T. Treatment of angina pectoris with Visken*. *Cardiol Hungarica Suppl 2A*: 67-71, 1973.
17. Lancaster R, Goodwin TJ, and Peart WS. The effect of pindolol on plasma renin activity and blood pressure in hypertensive patients. *Br J Clin Pharmacol* 3: 453-460, 1976.
18. Leary WP and Asmal AC. Treatment of coexistent angina pectoris and hypertension with pindolol. *S Afr Med J* 49: 11-12, 1975.
19. Malmborg RO, Isacsson SO, and Kallivroussis G. The effect of β -blockade and/or physical training in patients with angina pectoris. *Curr Ther Res* 16: 171-183, 1974.
20. Meier J. Pindolol: A pharmacokinetic comparison with other β -adrenoceptor blocking agents. *Curr Med Res Opin* 4 Suppl. 5: 31-38, 1977.
21. Meier J and Nuesch E. Pindolol, a β -adrenoceptor blocking agent with a negligible first-pass effect. *Br J Clin Pharmacol* 4: 371-372, 1977.
22. Meurice E, Remy-Schmetz J, Denys WJ, and Spruyt L. Essai préliminaire d'un β -bloqueur (pindolol) chez trois schizophrènes et trois malades psychiatriques non psychotiques. Différence de sensibilité de la fréquence cardiaque. *Acta Psychiat Belg* 74: 294-304, 1974.
23. Nager F and Favre H. Double-blind study with propranolol and LB-46 in angina pectoris. In: *Coronary Heart Disease. International Symposium*, Frankfurt, Jan. 22-24, 1970. Kaltenbach M and Lichtlen P (ed), Stuttgart, Georg Thieme-Verlag, 1971, pp. 224-226.
24. Nair DV. A double-blind study of a new β -adrenergic blocking agent LB-46 [4(2 hydroxy 3 isopropyl-aminopropoxy)-indole] in the treatment of angina pectoris. *Kerala Med J* 12: No. 4, 155-164, 1971.
25. Nigam PD and Malhotra AS. Prolonged action of pindolol. *Br Med J* I: 742-743, 1973.

26. Ohnhaus EE, Nuesch E, Meier J, and Kalberer F. Pharmacokinetics of unlabelled and ^{14}C -labelled pindolol in uraemia. *Eur J Clin Pharmacol* 7: 25-29, 1974.
27. Roubicek J. Effect of β -adrenoceptor blocking drugs on EEG. *Br J Clin Pharmacol* 3: 661-665, 1976.
28. Singh BN and Vaughan Williams EM. Effects on cardiac muscle of the β -adrenoceptor blocking drugs INPEA and LB-46 in relation to their local anesthetic action on nerve. *Br J Pharmacol* 43: 10-22, 1971.
29. Sainani GS and Mukherjee AK. A double-blind trial of LB-46 (Visken*) in angina pectoris. *Indian Heart J* 24: Suppl. 1: pp192-196, 1972.
30. Thorpe P. Pindolol (Visken) and angina pectoris: a double-blind multicentre trial. *NZ Med J* 76: 171-173, 1972.
31. Vetter W, Zaruba K, Armbruster H, Beckernoff R, Vetter H, Nussberger J, Schmied U, Reck G, Fontana A, and Siegenthaler W. Effect of propranolol and pindolol in renin secretion in normal supine man. *Klin Wschr* 53: 709-711, 1975.
32. Visken (pindolol) Product Monograph, Tribute Pharmaceuticals Canada Inc. Control No. 190690, Date of Revision: May 24, 2016.

PART III: CONSUMER INFORMATION

PrTEVA-PINDOLOL (pindolol)

This leaflet is Part III of a three-part ‘Product Monograph’ published when TEVA-PINDOLOL was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about TEVA-PINDOLOL. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION

What the medication is used for:

TEVA-PINDOLOL belongs to the group of medicines known as beta-adrenergic blocking agents (beta-blockers). It is used to treat high blood pressure, also called hypertension. It is also used to prevent a type of chest pain called Angina Pectoris.

What it does:

TEVA-PINDOLOL works by reducing your heart's workload in order to prevent chest pain and heart attacks.

With its ability to dilate blood vessels it is used to treat high blood pressure.

When it should not be used:

Do not take TEVA-PINDOLOL if you:

- are allergic to pindolol or any other beta-blockers or any of the inactive ingredients of TEVA-PINDOLOL listed under “What the non-medicinal ingredients are”,
- suffer from, or have suffered in the past from, bronchial asthma, or other lung diseases or trouble in breathing;
- suffer from a severe heart disease,
- have had an alteration in the structure and function of the right ventricle of the heart caused by a primary disorder of respiratory system (called Cor pulmonale),
- suffer from chest pain, mainly occurring at rest;
- suffer from irregular or unusually slow heartbeat (less than 45 to 50 beats per minute),
- have had sudden loss of consciousness in the past;
- suffer from severe blood flow disturbances of your blood vessels causing paleness or poor circulation in the arms and legs (cold hands or feet),
- suffer from pheochromocytoma (a type of tumor of the adrenal glands).

What the medicinal ingredient is:

Pindolol

What the non-medicinal ingredients are:

The non-medicinal ingredients are colloidal silicon dioxide, magnesium stearate, microcrystalline cellulose, pregelatinized starch, sodium lauryl sulfate and sodium starch glycolate.

What Dosage Forms it comes in:

Tablets; 5, 10 and 15 mg

WARNINGS AND PRECAUTIONS

BEFORE you use TEVA-PINDOLOL talk to your doctor or pharmacist if:

- If you suffer from any of the following:
 - diabetes mellitus (sugar diabetes), and you are taking insulin or diabetes medicine by mouth,
 - severe kidney disease,
 - liver disease,
 - milder forms of circulatory disturbances of blood vessels (symptoms such as paleness, cold hands or feet),
 - overactive thyroid,
 - chronic bronchitis and emphysema,
 - psoriasis (a type of skin disease characterized by thickened patches of red/silver skin).
- you are taking or have recently taken any other medicines, including medicines obtained without a prescription.
- you are pregnant, or plan to become pregnant.
- you are breastfeeding (the active ingredient may pass into your milk and harm your baby).
- you are having any kind of surgery or dental treatment.
- you are experiencing eye problems (symptoms like dry, gritty or burning eyes).

If any of these apply to you, **tell your doctor before taking TEVA-PINDOLOL**. The doctor will take these things into account before and during your treatment with TEVA-PINDOLOL and your doctor may need to monitor you more closely while you are using this medicine.

Driving and using machines

In some people, TEVA-PINDOLOL may cause dizziness or tiredness. If this happens to you, do not drive, use machinery, or perform other tasks that need full attention.

INTERACTIONS WITH THIS MEDICATION

Other medicines and TEVA-PINDOLOL may interfere with each other. These medicines include:

- medicines used to treat high blood pressure, chest pain (angina pectoris), disturbances of heart rhythm,
- digoxin and digitalis glycosides, a heart medicine,
- medicines containing adrenaline or similar substances, such as nose and eye drops, cough medicines, or remedies for the common cold (substances that raise blood pressure),
- insulin or antidiabetic medicines taken by mouth,
- medicines used to relieve pain or inflammation (especially non-steroidal anti-inflammatory drugs),
- ergot alkaloids, a class of medicines used in the prevention and treatment of migraine headaches,

- monoamine oxidase inhibitors (antidepressant drugs);
- medicine used to relieve heartburns and gastrointestinal ulcer (called cimetidine),
- medicines used to treat anxiety and/or panic disorder, schizophrenia or psychosis.
- Fingolimod, a medicine used to treat multiple sclerosis.

PROPER USE OF THIS MEDICATION

Follow your doctor's instructions carefully. Do not exceed the recommended dose.

How much TEVA-PINDOLOL to take

The dose of this medicine will be different for different patients. Also, the number of doses you take each day and the time allowed between doses depend on the medical problem for which you are taking TEVA-PINDOLOL.

Treatment is usually started with the lowest dose. Depending on how you respond to the treatment, your doctor may suggest a higher or lower dose.

In general the average daily dosage for the tablets is in the following ranges:

Usual adult dose:

- High blood pressure
5 to 15 mg once daily in the morning.
Doses of 20 mg should be taken in 2 divided doses in the morning and the evening.
- Angina pectoris (chest pain) and Increased heart rate (tachycardia)
10 to 30 mg daily in 3 or 4 divided doses.
- Hyperkinetic heart syndrome
10 to 20 mg daily

Take TEVA-PINDOLOL at the same time each day. Swallow the tablet(s) with a little liquid with food. Do not stop taking this drug without your doctor's advice.

This drug is not recommended for children under 18 years of age.

Patients at the age of 65 or over may experience more side effects than young patients. Close monitoring by your doctor may be required.

If you have questions about how long to take TEVA-PINDOLOL, talk to your doctor or your pharmacist.

Overdose:

In case of drug overdose, contact a health care practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

If you accidentally take a lot more TEVA-PINDOLOL tablets than your doctor has prescribed, contact or go to the nearest hospital emergency department or your doctor immediately, or make sure that someone else can contact them for you.

Signs and symptoms caused by taking more of TEVA-PINDOLOL can be: Abnormally low heartbeat, low blood pressure, low blood sugar; insufficient action of the heart to pump blood which may lead to loss of consciousness; irregular heart beat; heart failure; difficulty breathing with wheezing or coughing; shortness of breath; vomiting; convulsions (seizures); heart attack.

Missed Dose:

If you forget to take a dose of TEVA-PINDOLOL, take the missed dose as you notice it. However, if it is almost time for the next dose, skip the missed dose and go back to your regular dosage schedule. Do not double doses.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

As with all medicines, patients taking TEVA-PINDOLOL may experience side effects. Keep track of your side effects and consult the table below for appropriate action.

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

Symptom/effect	Talk with your doctor or pharmacist		Stop taking drug and seek immediate medical attention
	Only if severe	In all cases	
Unknown frequency:			
Allergic reaction: rash, hives, swelling of the face, lips, tongue or throat, difficulty swallowing or breathing			√
Slow heartbeat, heart rhythm disturbance, heart failure or sudden and oppressive chest pain			√
Coldness, burning, tingling or numbness in arms or legs		√	
Difficulty breathing with wheezing or coughing		√	
Hallucination (seeing or hearing things that are not there)		√	
Low blood pressure, shortness of breath, tiredness, dizziness,	√		

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

Symptom/effect	Talk with your doctor or pharmacist		Stop taking drug and seek
headache, trembling, nausea, vomiting, abdominal pain, diarrhea, muscle cramp, sleep disturbance, depression, skin reaction, excessive sweating, aggravation of psoriasis (thick patches of red/silver skin)			

This is not a complete list of side effects. For any unexpected effects while taking TEVA-PINDOLOL, contact your doctor or pharmacist.

HOW TO STORE IT

- Do not use after the expiry date shown on the box.
- Store at 15-30°C in the original package, protect from light.
- Keep out of the reach and sight of children.

REPORTING SIDE EFFECTS

You can help improve the safe use of health products for Canadians by reporting serious and unexpected side effects to Health Canada. Your report may help to identify new side effects and change the product safety information.

3 ways to report:

- Online at MedEffect (<http://hc-sc.gc.ca/dhp-mps/medeff/index-eng.php>);
- By calling 1-866-234-2345 (toll-free);
- By completing a Consumer Side Effect Reporting Form and sending it by:
 - Fax to 1-866-678-6789 (toll-free), or
 - Mail to: Canada Vigilance Program
Health Canada
Postal Locator 0701E
Ottawa, ON
K1A 0K9

Postage paid labels and the Consumer Side Effect Reporting Form are available at MedEffect (<http://hc-sc.gc.ca/dhp-mps/medeff/index-eng.php>).

NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

This document plus the full product monograph, prepared for health professionals can be found by contacting Teva Canada Limited by:

Phone: 1-800-268-4127 ext. 3;
Email: druginfo@tevacanada.com; or
Fax: 1-416-335-4472

Reporting Side Effects to Teva Canada Limited:

Phone: 1-800-268-4127 ext. 3;
Email: PhV@tevacanada.com; or
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