

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

PREXXARTAN™ (valsartan) oral solution Initial U.S. Approval: 1996

WARNING: FETAL TOXICITY

See full prescribing information for complete boxed warning.

- When pregnancy is detected, discontinue PREXXARTAN™ as soon as possible. (5.1)
- Drugs that act directly on the renin-angiotensin system can cause injury and death to the developing fetus. (5.1)

INDICATIONS AND USAGE

PREXXARTAN™ is an angiotensin II receptor blocker (ARB) indicated for:

- **Hypertension** in adults and children six years and older, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions (1.1)
- **Heart failure** (NYHA class II-IV); PREXXARTAN™ significantly reduces hospitalization for heart failure in patients who are unable to swallow valsartan tablets (1.2)
- **Post-myocardial infarction**, stable left ventricular failure or left ventricular dysfunction following myocardial infarction; PREXXARTAN™ reduces cardiovascular mortality in patients who are unable to swallow valsartan tablets (1.3)

DOSAGE AND ADMINISTRATION

Indication	Starting Dose	Dose Range	Target Maintenance
Hypertension—adults (2.1)	40 or 80 mg twice daily orally	40 -160 mg twice daily orally	---
Hypertension—age 6 to 16 years (2.2)	0.65 mg/kg twice daily orally (up to 40 mg total)	0.65-1.35 mg/kg twice daily orally (up to 40 mg-160 mg total)	---
Heart Failure (2.3)	40 mg twice daily orally	40 mg-160 mg twice daily orally	160 mg twice daily orally
Post-Myocardial Infarction (2.4)	20 mg twice daily orally	20 mg-160 mg twice daily orally	160 mg twice daily orally

*As tolerated by patient

DOSAGE FORMS AND STRENGTHS

Oral Solution, 4 mg/mL (20 mg/5 mL)

CONTRAINDICATIONS

Known hypersensitivity
Patients with diabetes on aliskiren (4)

WARNINGS AND PRECAUTIONS

- Observe for signs and symptoms of hypotension (5.2)
- Monitor renal function and potassium in susceptible patients (5.3, 5.4)

ADVERSE REACTIONS

Hypertension: Most common adverse reactions are headache, dizziness, fatigue and abdominal pain (6.1)

Heart Failure: Most common adverse reactions are dizziness, hypotension, diarrhea, arthralgia, back pain, fatigue and hyperkalemia (6.1)

Post-Myocardial Infarction: Most common adverse reactions which caused patients to discontinue therapy are hypotension, cough and increased blood creatinine (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Carmel Biosciences, Inc. at 1-732-314-4550 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Potassium-sparing diuretics, potassium supplements or salt substitutes may lead to increases in serum potassium, and in heart failure patients, increases in serum creatinine (7.1)
- Non-Steroidal Anti-Inflammatory Drugs (NSAIDs) increase risk of renal impairment and loss of antihypertensive effect (7.2)
- Dual inhibition of the Renin-Angiotensin System (RAS): Increased risk of renal impairment, hypotension, and hyperkalemia (7.3)
- **Lithium:** Increases in serum lithium level and lithium toxicity (7.4)

USE IN SPECIFIC POPULATIONS

Lactation: Breastfeeding is not recommended. (8.2)

Pediatrics: Efficacy and safety data support use in 6-16 year old patients; use is not recommended in patients <6 years old (6.1, 8.4)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 09/2026

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FULL PRESCRIBING INFORMATION

WARNING: FETAL TOXICITY

- **When pregnancy is detected, discontinue PREXXARTAN™ as soon as possible. (5.1)**
- **Drugs that act directly on the renin-angiotensin system can cause injury and death to the developing fetus. (5.1)**

1 INDICATIONS AND USAGE

1.1 Hypertension

PREXXARTAN™ is indicated for the treatment of hypertension in adults and children six years and older, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. These benefits have been seen in controlled trials of antihypertensive drugs from a wide variety of pharmacologic classes, including the class to which valsartan principally belongs. There are no controlled trials in hypertensive patients demonstrating risk reduction with valsartan.

Control of high blood pressure should be part of comprehensive cardiovascular risk management, including, as appropriate, lipid control, diabetes management, antithrombotic therapy, smoking cessation, exercise, and limited sodium intake. Many patients will require more than one drug to achieve blood pressure goals. For specific advice on goals and management, see published guidelines, such as those of the National High Blood Pressure Education Program's Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure (JNC).

Numerous antihypertensive drugs, from a variety of pharmacologic classes and with different mechanisms of action, have been shown in randomized controlled trials to reduce cardiovascular morbidity and mortality, and it can be concluded that it is blood pressure reduction, and not some other pharmacologic property of the drugs, that is largely responsible for those benefits. The largest and most consistent cardiovascular outcome benefit has been a reduction in the risk of stroke, but reductions in myocardial infarction and cardiovascular mortality also have been seen regularly.

Elevated systolic or diastolic pressure causes increased cardiovascular risk, and the absolute risk increase per mmHg is greater at higher blood pressures, so that even modest reductions of severe hypertension can provide substantial benefit. Relative risk reduction from blood pressure reduction is similar across populations with varying absolute risk, so the absolute benefit is greater in patients who are at higher risk independent of their hypertension (e.g., patients with diabetes or hyperlipidemia), and such patients would be expected to benefit from more aggressive treatment to a lower blood pressure goal.

Some antihypertensive drugs have smaller blood pressure effects (as monotherapy) in black patients, and many antihypertensive drugs have additional approved indications and effects (e.g., on angina, heart failure, or diabetic kidney disease). These considerations may guide selection of therapy.

PREXXARTAN™ may be used alone or in combination with other antihypertensive agents.

1.2 Heart Failure

PREXXARTAN™ is indicated for the treatment of heart failure (NYHA class II-IV) to reduce the risk of hospitalization for heart failure in patients who are unable to swallow valsartan tablets. There is no evidence that valsartan provides added benefits when it is used with an adequate dose of an angiotensin converting enzyme (ACE) inhibitor [see *Warnings and Precautions* (5.2), *Clinical Pharmacology* (12.3) and *Clinical Studies* (14.2)].

1.3 Post-Myocardial Infarction

PREXXARTAN™ is indicated to reduce the risk of cardiovascular death in clinically stable patients with left ventricular failure or left ventricular dysfunction following myocardial infarction who are unable to swallow valsartan tablets [see *Warnings and Precautions* (5.2), *Clinical Pharmacology* (12.3) and *Clinical Studies* (14.3)].

2 DOSAGE AND ADMINISTRATION

2.1 General Considerations

PREXXARTAN™ is not therapeutically equivalent to the tablet formulation of Diovan. The peak concentration of valsartan with PREXXARTAN™ is higher than with Diovan [see *Warnings and Precautions* (5.2) and *Clinical Pharmacology* (12.3)]. Follow dosing instructions given here.

2.2 Adult Hypertension

The recommended starting dose of PREXXARTAN™ is 40 mg or 80 mg twice daily orally when used as monotherapy in patients who are not volume-depleted. Patients requiring greater reductions in blood pressure may be started at 80 mg administered twice a day. PREXXARTAN™ may be used over a total daily dose range of 80 mg to 320 mg.

The antihypertensive effect is substantially present within 2 weeks and maximal reduction is generally attained after 4 weeks. If additional antihypertensive effect is required over the starting dose range, the total daily dose may be increased to a maximum of 320 mg or a diuretic may be added. Addition of a diuretic has a greater effect than dose increases beyond 80 mg.

PREXXARTAN™ may be administered with other antihypertensive agents.

2.3 Pediatric Hypertension 6 to 16 Years of Age

The usual recommended starting dose is 0.65 mg/kg twice daily orally (up to 40 mg total daily dose). The dosage should be adjusted according to blood pressure response and tolerability, up to a maximum dose of 1.35 mg/kg twice daily orally (maximum daily dose 160 mg). Doses higher than 1.35 mg/kg twice daily orally (or >160 mg total daily dose) have not been studied in pediatric patients 6 to 16 years old.

No data are available in pediatric patients either undergoing dialysis or with a glomerular filtration rate <30 mL/min/1.73 m² [see *Use in Specific Populations* (8.4)].

Use of PREXXARTAN™ is not recommended for patients under 6 years of age [see *Adverse Reactions* (6.1), *Pediatric Use in Specific Populations* (8.4), *Nonclinical Toxicology* (13.2), *Clinical Studies* (14.1)].

2.4 Heart Failure

The recommended starting dose of PREXXARTAN™ is 40 mg twice daily orally. Titrate to 80 mg and 160 mg twice daily orally or to the highest dose tolerated by the patient. Consider reducing the dose of concomitant diuretics. The maximum daily dose administered in clinical trials is 320 mg in divided doses.

2.5 Post-Myocardial Infarction

PREXXARTAN™ may be initiated as early as 12 hours after a myocardial infarction. The recommended starting dose of PREXXARTAN™ is 20 mg twice daily orally. Patients may be up titrated within 7 days to 40 mg twice daily orally, with subsequent titrations to a target maintenance dose of 160 mg twice daily orally, as tolerated by the patient. If symptomatic hypotension or renal dysfunction occurs, consider dosage reduction. PREXXARTAN™ may be given with other standard post-myocardial infarction treatment, including thrombolytics, aspirin, beta-blockers, and statins.

2.6 Missed Dose

If a dose of PREXXARTAN™ is missed, it should be administered as soon as possible, unless it is almost time for the next dose. The dose should not be doubled to make up for a missed dose.

3 DOSAGE FORMS AND STRENGTHS

PREXXARTAN™ (valsartan) Oral Solution contains 4 mg/mL (20 mg/5 mL) valsartan and is available as a clear, colorless, grape-flavored solution.

4 CONTRAINDICATIONS

Do not use in patients with known hypersensitivity to any component.

Do not coadminister aliskiren with PREXXARTAN™ in patients with diabetes [see *Drug Interactions* (7.3)].

5 WARNINGS AND PRECAUTIONS

5.1 Fetal Toxicity

PREXXARTAN™ can cause fetal harm when administered to a pregnant woman. Use of drugs that act on the renin-angiotensin system during the second and third trimesters of pregnancy reduces fetal renal function and increases fetal and

neonatal morbidity and death. Resulting oligohydramnios can be associated with fetal lung hypoplasia and skeletal deformations. Potential neonatal adverse effects include skull hypoplasia, anuria, hypotension, renal failure, and death. When pregnancy is detected, discontinue PREXXARTAN™ as soon as possible [see *Use in Specific Populations* (8.1)].

5.2 Hypotension

In patients with an activated renin-angiotensin system, such as volume- and/or salt-depleted patients receiving high doses of diuretics, symptomatic hypotension may occur. This condition should be corrected prior to administration of valsartan, or the treatment should start under close medical supervision.

Peak plasma concentrations of valsartan are higher following administration of PREXXARTAN™ and may result in increased risk of hypotension as compared to administration of valsartan tablets [see *Clinical Pharmacology* (12.3)]. Patients with heart failure or post-myocardial infarction patients given valsartan tablets in clinical trials commonly had some reduction in blood pressure. Only use PREXXARTAN™ in heart failure or post-myocardial infarction patients who are unable to swallow valsartan tablets. In clinical trials of valsartan tablets, discontinuation of therapy because of continuing symptomatic hypotension usually was not necessary. In controlled trials in heart failure patients, the incidence of hypotension in valsartan-treated patients was 5.5% compared to 1.8% in placebo-treated patients. In the Valsartan in Acute Myocardial Infarction Trial (VALIANT), hypotension in post-myocardial infarction patients led to permanent discontinuation of therapy in 1.4% of valsartan-treated patients and 0.8% of captopril-treated patients.

If symptomatic hypotension occurs, place the patient in the supine position and, if necessary, give an intravenous infusion of normal saline. A transient hypotensive response is not a contraindication to further treatment, which usually can be continued without difficulty once the blood pressure has stabilized.

5.3 Impaired Renal Function

Changes in renal function including acute renal failure can be caused by drugs that inhibit the renin-angiotensin system and by diuretics. Patients whose renal function may depend in part on the activity of the renin-angiotensin system (e.g., patients with renal artery stenosis, chronic kidney disease, severe congestive heart failure, or volume depletion) may be at particular risk of developing acute renal failure on valsartan. Monitor renal function periodically in these patients. Consider withholding or discontinuing therapy in patients who develop a clinically significant decrease in renal function on valsartan [see *Drug Interactions* (7)].

5.4 Hyperkalemia

Some patients with heart failure have developed increases in potassium. These effects are usually minor and transient, and they are more likely to occur in patients with pre-existing renal impairment. Dosage reduction and/or discontinuation of PREXXARTAN™ may be required [see *Adverse Reactions* (6.1)].

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

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

Adult Hypertension

Valsartan has been evaluated for safety in more than 4,000 patients, including over 400 treated for over 6 months, and more than 160 for over 1 year. Adverse reactions have generally been mild and transient in nature and have only infrequently required discontinuation of therapy. The overall incidence of adverse reactions with valsartan was similar to placebo.

The overall frequency of adverse reactions was neither dose-related nor related to gender, age, race, or regimen. Discontinuation of therapy due to side effects was required in 2.3% of valsartan patients and 2.0% of placebo patients. The most common reasons for discontinuation of therapy with valsartan were headache and dizziness.

The adverse reactions that occurred in placebo-controlled clinical trials in at least 1% of patients treated with valsartan and at a higher incidence in valsartan (n=2,316) than placebo (n=888) patients included fatigue (2% vs. 1%) and abdominal pain (2% vs. 1%).

In trials in which valsartan was compared to an ACE inhibitor with or without placebo, the incidence of dry cough was

significantly greater in the ACE-inhibitor group (7.9%) than in the groups who received valsartan (2.6%) or placebo (1.5%). In a 129-patient trial limited to patients who had had dry cough when they had previously received ACE inhibitors, the incidences of cough in patients who received valsartan, HCTZ, or lisinopril were 20%, 19%, and 69% respectively ($p < 0.001$).

Dose-related orthostatic effects were seen in less than 1% of patients. An increase in the incidence of dizziness was observed in patients treated with valsartan 320 mg (8%) compared to 10 to 160 mg (2% to 4%).

Pediatric Hypertension

Valsartan has been evaluated for safety in over 400 pediatric patients aged 6 to 17 years and more than 160 pediatric patients aged 6 months to 5 years. No relevant differences were identified between the adverse experience profile for pediatric patients aged 6 to 16 years and that previously reported for adult patients. Headache and hyperkalemia were the most common adverse events suspected to be study drug-related in older children (6 to 17 years old) and younger children (6 months to 5 years old), respectively. Hyperkalemia was mainly observed in children with underlying renal disease.

Neurocognitive and developmental assessment of pediatric patients aged 6 to 16 years revealed no overall clinically relevant adverse impact after treatment with valsartan for up to 1 year.

Valsartan is not recommended for pediatric patients under 6 years of age. In a study ($n=90$) of pediatric patients (1 to 5 years), two deaths and three cases of on-treatment transaminase elevations were seen in the one-year open-label extension phase. These 5 events occurred in a study population in which patients frequently had significant co-morbidities. A causal relationship to valsartan has not been established. In a second study of 6-months duration in 75 children aged 1 to 5 years, there were no deaths; one case of marked liver transaminase elevations occurred following 6 months of treatment.

Heart Failure

In the Valsartan Heart Failure Trial (Val-HeFT), comparing valsartan in total daily doses up to 320 mg ($n=2,506$) to placebo ($n=2,494$), 10% of valsartan patients discontinued for adverse reactions vs. 7% of placebo patients.

The table shows adverse reactions in double-blind short-term heart failure trials, including the first 4 months of the Valsartan Heart Failure Trial, with an incidence of at least 2% that were more frequent in valsartan-treated patients than in placebo-treated patients. All patients received standard drug therapy for heart failure, frequently as multiple medications, which could include diuretics, digitalis, beta-blockers. About 93% of patients received concomitant ACE inhibitors.

	Valsartan (n=3,282)	Placebo (n=2,740)
Dizziness	17%	9%
Hypotension	7%	2%
Diarrhea	5%	4%
Arthralgia	3%	2%
Fatigue	3%	2%
Back Pain	3%	2%
Dizziness, postural	2%	1%
Hyperkalemia	2%	1%
Hypotension, postural	2%	1%

Discontinuations occurred in 0.5% of valsartan-treated patients and 0.1% of placebo patients for each of the following: elevations in creatinine and elevations in potassium.

Other adverse reactions with an incidence greater than 1% and greater than placebo included headache, nausea, renal impairment, syncope, blurred vision, upper abdominal pain and vertigo.

From the long-term data in the Valsartan Heart Failure Trial, there did not appear to be any significant adverse reactions not previously identified.

Post-Myocardial Infarction

The table shows the percentage of patients discontinued in the valsartan and captopril-treated groups in the VALsartan In Acute myocardial infarction trial (VALIANT) with a rate of at least 0.5% in either of the treatment groups.

Discontinuations due to renal dysfunction occurred in 1.1% of valsartan-treated patients and 0.8% of captopril-treated

patients.

	Valsartan (n=4,885)	Captopril (n=4,879)
Discontinuation for adverse reaction	5.8%	7.7%
Adverse reactions		
Hypotension NOS	1.4%	0.8%
Cough	0.6%	2.5%
Blood creatinine increased	0.6%	0.4%
Rash NOS	0.2%	0.6%

Clinical Laboratory Test Findings

Creatinine: In heart failure trials, greater than 50% increases in creatinine were observed in 3.9% of valsartan-treated patients compared to 0.9% of placebo-treated patients. In post-myocardial infarction patients, doubling of serum creatinine was observed in 4.2% of valsartan-treated patients and 3.4% of captopril-treated patients.

Neutropenia: Neutropenia was observed in 1.9% of patients treated with valsartan and 0.8% of patients treated with placebo.

Blood Urea Nitrogen (BUN): In heart failure trials, greater than 50% increases in BUN were observed in 16.6% of valsartan-treated patients compared to 6.3% of placebo-treated patients [see *Warnings and Precautions (5.3)*].

6.2 Postmarketing Experience

The following additional adverse reactions have been reported in postmarketing use of valsartan. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Hypersensitivity: Angioedema has been reported. Some of these patients previously experienced angioedema with other drugs, including ACE inhibitors. Valsartan should not be re-administered to patients who have had angioedema.

Digestive: Elevated liver enzymes and very rare reports of hepatitis

Musculoskeletal: Rhabdomyolysis

Renal: Impaired renal function, renal failure

Dermatologic: Alopecia, bullous dermatitis

Blood and Lymphatic: Thrombocytopenia.

Vascular: Vasculitis

7 DRUG INTERACTIONS

7.1 Agents Increasing Serum Potassium

Concomitant use of valsartan with other agents that block the renin-angiotensin system, potassium-sparing diuretics (e.g., spironolactone, triamterene, amiloride), potassium supplements, salt substitutes containing potassium or other drugs that may increase potassium levels (e.g., heparin) may lead to increases in serum potassium and in heart failure patients to increases in serum creatinine. If co-medication is considered necessary, monitoring of serum potassium is advisable.

7.2 Non-Steroidal Anti-Inflammatory Agents Including Selective Cyclooxygenase-2 Inhibitors (COX-2 Inhibitors)

In patients who are elderly, volume-depleted (including those on diuretic therapy), or with compromised renal function, coadministration of NSAIDs, including selective COX-2 inhibitors, with angiotensin II receptor antagonists, including valsartan, may result in deterioration of renal function, including possible acute renal failure. These effects are usually reversible. Monitor renal function periodically in patients receiving valsartan and NSAID therapy.

The antihypertensive effect of angiotensin II receptor antagonists, including valsartan, may be attenuated by NSAIDs, including selective COX-2 inhibitors.

7.3 Dual Blockade of the Renin-Angiotensin System (RAS)

Dual blockade of the RAS with angiotensin receptor blockers, ACE inhibitors, or aliskiren is associated with increased

risks of hypotension, hyperkalemia, and changes in renal function (including acute renal failure) compared to monotherapy. Most patients receiving the combination of two RAS inhibitors do not obtain any additional benefit compared to monotherapy [see *Clinical Studies (14.3)*]. In general, avoid combined use of RAS inhibitors. Closely monitor blood pressure, renal function and electrolytes in patients on PREXXARTAN™ and other agents that affect the RAS.

Do not coadminister aliskiren with PREXXARTAN™ in patients with diabetes. Avoid use of aliskiren with PREXXARTAN™ in patients with renal impairment (GFR < 60 mL/min).

7.4 Lithium

Increases in serum lithium concentrations and lithium toxicity have been reported during concomitant administration of lithium with angiotensin II receptor antagonists, including valsartan. Monitor serum lithium levels during concomitant use.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

PREXXARTAN™ can cause fetal harm when administered to a pregnant woman. Use of drugs that act on the renin-angiotensin system during the second and third trimesters of pregnancy reduces fetal renal function and increases fetal and neonatal morbidity and death. Most epidemiologic studies examining fetal abnormalities after exposure to antihypertensive use in the first trimester have not distinguished drugs affecting the renin-angiotensin system from other antihypertensive agents. Published reports include cases of anhydramnios and oligohydramnios in pregnant women treated with valsartan (see *Clinical Considerations*). Studies in rats and rabbits with valsartan showed fetotoxicity only at maternally toxic doses (see *Data*).

When pregnancy is detected, consider alternative drug treatment and discontinue PREXXARTAN™ as soon as possible.

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2 to 4%, and 15 to 20%, respectively.

Clinical Considerations

Disease-associated maternal and/or embryo/fetal risk

Hypertension in pregnancy increases the maternal risk for pre-eclampsia, gestational diabetes, premature delivery, and delivery complications (e.g., need for cesarean section, and post-partum hemorrhage). Hypertension increases the fetal risk for intrauterine growth restriction and intrauterine death. Pregnant women with hypertension should be carefully monitored and managed accordingly.

Fetal/Neonatal Adverse Reactions

Oligohydramnios in pregnant women who use drugs affecting the renin-angiotensin system in the second and third trimesters of pregnancy can result in the following: reduced fetal renal function leading to anuria and renal failure, fetal lung hypoplasia, skeletal deformations, including skull hypoplasia, hypotension and death. In the unusual case that there is no appropriate alternative to therapy with drugs affecting the renin-angiotensin system for a particular patient, apprise the mother of the potential risk to the fetus.

In patients taking PREXXARTAN™ during pregnancy, perform serial ultrasound examinations to assess the intra-amniotic environment. Fetal testing may be appropriate, based on the week of gestation. Patients and physicians should be aware, however, that oligohydramnios may not appear until after the fetus has sustained irreversible injury. If oligohydramnios is observed, consider alternative drug treatment. Closely observe neonates with histories of *in utero* exposure to PREXXARTAN™ for hypotension, oliguria, and hyperkalemia. In neonates with a history of *in utero* exposure to PREXXARTAN™, if oliguria or hypotension occurs, support blood pressure and renal perfusion. Exchange transfusions or dialysis may be required as a means of reversing hypotension and replacing renal function.

Data

Animal Data

No teratogenic effects were observed when valsartan was administered to pregnant mice and rats at oral doses of up to 600 mg/kg/day (9 and 18 times the maximum recommended human dose (MRHD) on a mg/m² basis) and to pregnant rabbits at oral doses of up to 10 mg/kg/day.

In rats, oral valsartan administered at maternally toxic doses (600 mg/kg/day) during organogenesis or late gestation and lactation, resulted in decreased fetal and pup weight, pup survival and delayed developmental milestones. In rabbits administered maternally toxic doses of 5 and 10 mg/kg/day, fetotoxicity was observed.

8.2 Lactation

Risk Summary

There are no data on the presence of PREXXARTAN™ in human milk, the effects on the breastfed infant, or the effects on milk production. Valsartan is present in rat milk (*see Data*). Because of the potential for valsartan to affect postnatal renal development in nursing infants, advise a nursing woman not to breastfeed during treatment with PREXXARTAN™.

Data

Valsartan was detected in the milk of lactating rats 15 minutes after oral administration of a 3 mg/kg dose.

8.4 Pediatric Use

PREXXARTAN™ is not recommended for pediatric patients under 6 years of age due to safety findings for which a relationship to treatment could not be excluded [*see Adverse Reactions (6.1)*]. Furthermore, it is unknown whether post-natal use of valsartan before maturation of renal function is complete has long-term deleterious effects on the kidney. In humans, nephrogenesis is thought to be complete around birth; however, maturation of other aspects of kidney function (such as glomerular filtration and tubular function) may continue until approximately 2 years of age.

The antihypertensive effects of valsartan have been evaluated in two randomized, double-blind clinical studies in pediatric patients from 1-5 and 6-16 years of age [*see Clinical Studies (14.1)*]. The pharmacokinetics of valsartan have been evaluated in pediatric patients 1 to 16 years of age [*see Clinical Pharmacology (12.3)*]. Valsartan was generally well tolerated in children 6-16 years and the adverse experience profile was similar to that described for adults.

In children and adolescents with hypertension where underlying renal abnormalities may be more common, renal function and serum potassium should be closely monitored as clinically indicated.

No data are available in pediatric patients either undergoing dialysis or with a glomerular filtration rate less than 30 mL/min/1.73m².

There is limited clinical experience with valsartan in pediatric patients with mild to moderate hepatic impairment [*see Warnings and Precautions (5.3)*].

8.5 Geriatric Use

In the controlled clinical trials of valsartan, 1,214 (36.2%) hypertensive patients treated with valsartan were ≥65 years and 265 (7.9%) were ≥75 years. No overall difference in the efficacy or safety of valsartan was observed in this patient population, but greater sensitivity of some older individuals cannot be ruled out.

Exposure [measured by area under the curve (AUC)] to valsartan is higher by 70% in the elderly than in the young, however no dosage adjustment is necessary [*see Clinical Pharmacology (12.3)*].

Of the 2,511 patients with heart failure randomized to valsartan in the Valsartan Heart Failure Trial, 45% (1,141) were 65 years of age or older. In the VALsartan In Acute Myocardial iNfarcTion Trial (VALIANT), 53% (2,596) of the 4,909 patients treated with valsartan and 51% (2,515) of the 4,885 patients treated with valsartan + captopril were 65 years of age or older. There were no notable differences in efficacy or safety between older and younger patients in either trial.

8.6 Renal Impairment

Safety and effectiveness of valsartan in patients with severe renal impairment (glomerular filtration rate less than 30 mL/min/1.73m²) have not been established. No dose adjustment is required in patients with mild (glomerular filtration rate 60 to 90 mL/min/1.73m²) or moderate (glomerular filtration rate 30 to 60 mL/min/1.73m²) renal impairment.

8.7 Hepatic Impairment

No dose adjustment is necessary for patients with mild-to-moderate liver disease. No dosing recommendations can be provided for patients with severe liver disease.

10 OVERDOSAGE

Limited data are available related to overdosage in humans. The most likely manifestations of overdosage would be hypotension and tachycardia; bradycardia could occur from parasympathetic (vagal) stimulation. Depressed level of consciousness, circulatory collapse and shock have been reported. If symptomatic hypotension should occur, institute supportive treatment.

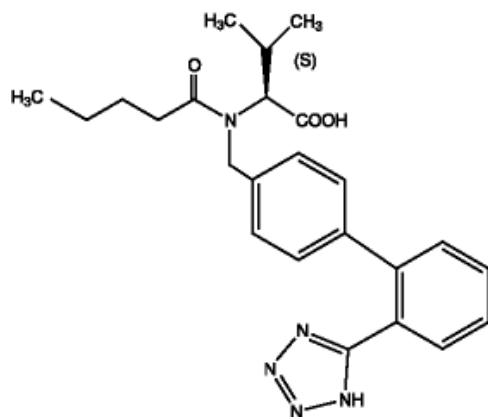
Valsartan is not removed from the plasma by hemodialysis.

Valsartan was without grossly observable adverse effects at single oral doses up to 2000 mg/kg in rats and up to 1000 mg/kg in marmosets, except for salivation and diarrhea in the rat and vomiting in the marmoset at the highest dose (60 and 31 times, respectively, the MRHD dose on a mg/m^2 basis). (Calculations assume an oral dose of 320 mg/day and a 60-kg patient).

11 DESCRIPTION

PREXXARTAN™ (valsartan) is a nonpeptide, orally active, and specific angiotensin II receptor blocker acting on the AT₁ receptor subtype.

Valsartan is chemically described as *N*-(1-oxopentyl)-*N*-[[2'-(1*H*-tetrazol-5-yl) [1,1'-biphenyl]-4-yl]methyl]-L-valine. Its empirical formula is C₂₄H₂₉N₅O₃, its molecular weight is 435.5, and its structural formula is:



Valsartan is a white to practically white fine powder. It is soluble in ethanol and methanol and slightly soluble in water.

PREXXARTAN™ is formulated at a concentration of 4 mg/mL valsartan in a grape flavored aqueous solution for oral administration. The inactive ingredients are: grape flavor, methylparaben NF, poloxamer 188, potassium sorbate, propylene glycol NF, purified water USP, sodium citrate dihydrate USP, and sucralose NF.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Angiotensin II is formed from angiotensin I in a reaction catalyzed by angiotensin-converting enzyme (ACE, kininase II). Angiotensin II is the principal pressor agent of the renin-angiotensin system, with effects that include vasoconstriction, stimulation of synthesis and release of aldosterone, cardiac stimulation, and renal reabsorption of sodium. Valsartan blocks the vasoconstrictor and aldosterone-secreting effects of angiotensin II by selectively blocking the binding of angiotensin II to the AT₁ receptor in many tissues, such as vascular smooth muscle and the adrenal gland. Its action is therefore independent of the pathways for angiotensin II synthesis.

There is also an AT₂ receptor found in many tissues, but AT₂ is not known to be associated with cardiovascular homeostasis. Valsartan has much greater affinity (about 20,000-fold) for the AT₁ receptor than for the AT₂ receptor. The increased plasma levels of angiotensin II following AT₁ receptor blockade with valsartan may stimulate the unblocked AT₂ receptor. The primary metabolite of valsartan is essentially inactive with an affinity for the AT₁ receptor about one- 200th (1/200th) that of valsartan itself.

Blockade of the renin-angiotensin system with ACE inhibitors, which inhibit the biosynthesis of angiotensin II from angiotensin I, is widely used in the treatment of hypertension. ACE inhibitors also inhibit the degradation of bradykinin, a reaction also catalyzed by ACE. Because valsartan does not inhibit ACE (kininase II), it does not affect the response to bradykinin. Whether this difference has clinical relevance is not yet known. Valsartan does not bind to or block other hormone receptors or ion channels known to be important in cardiovascular regulation.

Blockade of the angiotensin II receptor inhibits the negative regulatory feedback of angiotensin II on renin secretion, but the resulting increased plasma renin activity and angiotensin II circulating levels do not overcome the effect of valsartan on blood pressure.

12.2 Pharmacodynamics

Valsartan inhibits the pressor effect of angiotensin II infusions. An oral dose of 80 mg inhibits the pressor effect by about 80% at peak with approximately 30% inhibition persisting for 24 hours. No information on the effect of larger doses is available.

Removal of the negative feedback of angiotensin II causes a 2- to 3-fold rise in plasma renin and consequent rise in angiotensin II plasma concentration in hypertensive patients. Minimal decreases in plasma aldosterone were observed after administration of valsartan; very little effect on serum potassium was observed.

In multiple-dose studies in hypertensive patients with stable renal insufficiency and patients with renovascular hypertension, valsartan had no clinically significant effects on glomerular filtration rate, filtration fraction, creatinine clearance, or renal plasma flow.

In multiple-dose studies in hypertensive patients, valsartan had no notable effects on total cholesterol, fasting triglycerides, fasting serum glucose, or uric acid.

12.3 Pharmacokinetics

For an equivalent dose, PREXXARTAN™ has 86% higher peak concentration (C_{max}) and 25% higher area under the plasma concentration over time curve (AUC) for valsartan compared to Diovan. AUC and C_{max} of valsartan increase approximately linearly with increasing dose over the clinical dosing range. Valsartan does not accumulate appreciably in plasma following repeated administration.

Absorption

PREXXARTAN™ C_{max} is achieved 0.7 to 3.7 hours after dosing.

Effect of Food

High-fat, high-calorie meal decreased the AUC of PREXXARTAN™ by about 8% and C_{max} by about 44%.

Distribution:

The steady state volume of distribution of valsartan after intravenous administration is small (17 L), indicating that valsartan does not distribute into tissues extensively. Valsartan is highly bound to serum proteins (95%), mainly serum albumin.

Elimination

Following intravenous administration, plasma clearance of valsartan is about 2 L/h. Renal clearance of valsartan is 0.62 L/h (about 30% of total body clearance). Valsartan shows bi-exponential decay kinetics following intravenous administration, with an average elimination half-life of about 6 hours.

Metabolism:

The primary metabolite, accounting for about 9% of dose, is valeryl 4-hydroxy valsartan. *In vitro* metabolism studies involving recombinant CYP 450 enzymes indicated that the CYP 2C9 isoenzyme is responsible for the formation of valeryl-4-hydroxy valsartan. Valsartan does not inhibit CYP 450 isozymes at clinically relevant concentrations. CYP 450 mediated drug interaction between valsartan and co-administered drugs are unlikely because of the low extent of metabolism.

Excretion

Valsartan, when administered as an oral solution, is primarily recovered in feces (about 83% of dose) and urine (about 13% of dose). The recovery is mainly as unchanged drug, with only about 20% of dose recovered as metabolites.

Following intravenous administration, plasma clearance of valsartan is about 2 L/h and its renal clearance is 0.62 L/h

(about 30% of total clearance).

The apparent clearance of valsartan following oral administration is approximately 4.5 L/h in heart failure patients. Age does not affect the apparent clearance in heart failure patients.

Specific Populations:

Geriatric Patients: Exposure (measured by AUC) to valsartan is higher by 70% and the half-life is longer by 35% in the elderly than in the young [see *Use in Specific Populations* (8.5)].

Pediatric Patients: In a study of pediatric hypertensive patients (n=26, 1 to 16 years of age) given single doses of a suspension of valsartan (mean: 0.9 to 2 mg/kg), the clearance (L/h/kg) of valsartan for children was similar to that of adults receiving the same formulation.

Male and Female Patients: Pharmacokinetics of valsartan does not differ significantly between males and females.

Patients with Renal Impairment: There is no apparent correlation between renal function (measured by creatinine clearance) and exposure (measured by AUC) to valsartan in patients with different degrees of renal impairment (down to creatinine clearance of 10 mL/min). Valsartan is not removed from the plasma by hemodialysis. [see *Use in Specific Populations* (8.6)].

Patients with Hepatic Impairment: On average, patients with mild-to-moderate chronic liver disease have twice the exposure (measured by AUC values) to valsartan of healthy volunteers (matched by age, sex, and weight) [see *Use in Specific Populations* (8.7)].

Drug Interaction Studies

No clinically significant pharmacokinetic interactions were observed when valsartan was coadministered with nebivolol, amlodipine, atenolol, cimetidine, digoxin, furosemide, glyburide, hydrochlorothiazide, or indomethacin. The valsartan-atenolol combination was more antihypertensive than either component, but it did not lower the heart rate more than atenolol alone.

Co-administration of valsartan and warfarin did not change the pharmacokinetics of valsartan or the time-course of the anticoagulant properties of warfarin.

Transporters: The results from an *in vitro* study with human liver tissue indicate that valsartan is a substrate of the hepatic uptake transporter OATP1B1 and the hepatic efflux transporter MRP2. Coadministration of inhibitors of the uptake transporter (rifampin, cyclosporine) or efflux transporter (ritonavir) may increase the systemic exposure to valsartan.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

There was no evidence of carcinogenicity when valsartan was administered in the diet to mice and rats for up to 2 years at doses up to 160 and 200 mg/kg/day, respectively. These doses in mice and rats are about 2.6 and 6 times, respectively, the MRHD on a mg/m² basis. (Calculations assume an oral dose of 320 mg/day and a 60-kg patient).

Mutagenicity assays did not reveal any valsartan-related effects at either the gene or chromosome level. These assays included bacterial mutagenicity tests with *Salmonella* (Ames) and *E coli*; a gene mutation test with Chinese hamster V79 cells; a cytogenetic test with Chinese hamster ovary cells; and a rat micronucleus test.

Valsartan had no adverse effects on the reproductive performance of male or female rats at oral doses up to 200 mg/kg/day. This dose is 6 times the MRHD on a mg/m² basis (Calculations assume an oral dose of 320 mg/day and a 60-kg patient).

13.2 Animal Toxicology and/or Pharmacology

Daily oral dosing of neonatal/juvenile rats with valsartan at doses as low as 1 mg/kg/day (about 10% of the maximum recommended pediatric dose on a mg/m² basis) from postnatal day 7 to postnatal day 70 produced persistent, irreversible kidney damage. These kidney effects in neonatal rats represent expected exaggerated pharmacological effects that are observed if rats are treated during the first 13 days of life. This period coincides with 36 weeks of gestation in humans, which could occasionally extend up to 44 weeks after conception in humans. In humans, nephrogenesis is thought to be complete around birth; however, maturation of other aspects of kidney function (such as glomerular filtration and tubular function) may continue until approximately 2 years of age. It is unknown whether post-natal use of valsartan before

maturation of renal function is complete has long-term deleterious effects on the kidney [*see Use in Specific Populations (8.4)*].

14 CLINICAL STUDIES

14.1 Hypertension

Studies evaluating the antihypertensive effects of valsartan were conducted with a formulation that is not therapeutically equivalent to PREXXARTAN™ [*see Clinical Pharmacology (12.3)*].

Adult Hypertension

The antihypertensive effects of valsartan were demonstrated principally in 7 placebo-controlled, 4- to 12-week trials (1 in patients over 65 years) of dosages from 10 to 320 mg/day in patients with baseline diastolic blood pressures of 95-115 mmHg. The studies allowed comparison of once-daily and twice-daily regimens of 160 mg/day; comparison of peak and trough effects; comparison (in pooled data) of response by gender, age, and race; and evaluation of incremental effects of hydrochlorothiazide.

Administration of valsartan to patients with essential hypertension results in a significant reduction of sitting, supine, and standing systolic and diastolic blood pressure, usually with little or no orthostatic change.

In most patients, after administration of a single oral dose, onset of antihypertensive activity occurs at approximately 2 hours, and maximum reduction of blood pressure is achieved within 6 hours. The antihypertensive effect persists for 24 hours after dosing, but there is a decrease from peak effect at lower doses (40 mg) presumably reflecting loss of inhibition of angiotensin II. At higher doses, however (160 mg), there is little difference in peak and trough effect. During repeated dosing, the reduction in blood pressure with any dose is substantially present within 2 weeks, and maximal reduction is generally attained after 4 weeks. In long-term follow-up studies (without placebo control), the effect of valsartan appeared to be maintained for up to 2 years. The antihypertensive effect is independent of age, gender or race. The latter finding regarding race is based on pooled data and should be viewed with caution because antihypertensive drugs that affect the renin-angiotensin system (that is, ACE inhibitors and angiotensin-II blockers) have generally been found to be less effective in low-renin hypertensives (frequently blacks) than in high-renin hypertensives (frequently whites). In pooled, randomized, controlled trials of valsartan that included a total of 140 blacks and 830 whites, valsartan and an ACE-inhibitor control were generally at least as effective in blacks as whites. The explanation for this difference from previous findings is unclear.

Abrupt withdrawal of valsartan has not been associated with a rapid increase in blood pressure.

The blood pressure-lowering effect of valsartan and thiazide-type diuretics are approximately additive.

The 7 studies of valsartan monotherapy included over 2,000 patients randomized to various doses of valsartan and about 800 patients randomized to placebo. Doses below 80 mg were not consistently distinguished from those of placebo at trough, but doses of 80, 160 and 320 mg produced dose-related decreases in systolic and diastolic blood pressure, with the difference from placebo of approximately 6-9/3-5 mmHg at 80 to 160 mg and 9/6 mmHg at 320 mg. In a controlled trial the addition of HCTZ to valsartan 80 mg resulted in additional lowering of systolic and diastolic blood pressure by approximately 6/3 and 12/5 mmHg for 12.5 and 25 mg of HCTZ, respectively, compared to valsartan 80 mg alone.

Patients with an inadequate response to 80 mg once daily were titrated to either 160 mg once daily or 80 mg twice daily, which resulted in a similar response in both groups.

In controlled trials, the antihypertensive effect of once-daily valsartan 80 mg was similar to that of once-daily enalapril 20 mg or once-daily lisinopril 10 mg.

There are no trials of valsartan demonstrating reductions in cardiovascular risk in patients with hypertension, but at least one pharmacologically similar drug has demonstrated such benefits.

There was essentially no change in heart rate in valsartan-treated patients in controlled trials.

Pediatric Hypertension

Children Between 6 to 16 Years of Age

In a clinical study involving 261 hypertensive pediatric patients 6 to 16 years of age, patients who weighed less than 35 kg received 10, 40 or 80 mg of valsartan daily (low, medium and high doses), and patients who weighed greater than or equal to 35 kg received 20, 80, and 160 mg of valsartan daily (low, medium and high doses). Renal and urinary disorders, and

essential hypertension with or without obesity were the most common underlying causes of hypertension in children enrolled in this study. At the end of 2 weeks, valsartan reduced both systolic and diastolic blood pressure in a dose-dependent manner. Overall, the three dose levels of valsartan (low, medium and high) significantly reduced systolic blood pressure by 8, 10, and 12 mmHg from the baseline, respectively. Patients were re-randomized to either continue receiving the same dose of valsartan or were switched to placebo. In patients who continued to receive the medium and high doses of valsartan, systolic blood pressure at trough was 4 and 7 mmHg lower than patients who received the placebo treatment. In patients receiving the low dose of valsartan, systolic blood pressure at trough was similar to that of patients who received the placebo treatment. Overall, the dose-dependent antihypertensive effect of valsartan was consistent across all the demographic subgroups.

14.2 Heart Failure

The Valsartan Heart Failure Trial (Val-HeFT) was a multinational, double-blind study in which 5,010 patients with NYHA class II (62%) to IV (2%) heart failure and LVEF less than 40%, on baseline therapy chosen by their physicians, were randomized to placebo or valsartan (titrated from 40 mg twice daily to the highest tolerated dose or 160 mg twice daily) and followed for a mean of about 2 years. The VAL-HeFT study was conducted with a formulation of valsartan that is not therapeutically equivalent to PREXXARTAN™ [see *Clinical Pharmacology* (12.3)]. Although Val-HeFT's primary goal was to examine the effect of valsartan when added to an ACE inhibitor, about 7% were not receiving an ACE inhibitor. Other background therapy included diuretics (86%), digoxin (67%), and beta-blockers (36%). The population studied was 80% male, 46% 65 years or older and 89% Caucasian. At the end of the trial, patients in the valsartan group had a blood pressure that was 4 mmHg systolic and 2 mmHg diastolic lower than the placebo group. There were two primary end points, both assessed as time to first event: all-cause mortality and heart failure morbidity, the latter defined as all-cause mortality, sudden death with resuscitation, hospitalization for heart failure, and the need for intravenous inotropic or vasodilatory drugs for at least 4 hours. These results are summarized in the following table.

	Placebo (N=2,499)	Valsartan (N=2,511)	Hazard Ratio (95% CI*)	Nominal p-value
All-cause mortality	484 (19.4%)	495 (19.7%)	1.02 (0.90–1.15)	0.8
HF morbidity	801 (32.1%)	723 (28.8%)	0.87 (0.79–0.97)	0.009

* CI = Confidence Interval

Although the overall morbidity result favored valsartan, this result was largely driven by the 7% of patients not receiving an ACE inhibitor, as shown in the following table.

	Without ACE Inhibitor		With ACE Inhibitor	
	Placebo (N=181)	Valsartan (N=185)	Placebo (N=2,318)	Valsartan (N=2,326)
Events (%)	77 (42.5%)	46 (24.9%)	724 (31.2%)	677 (29.1%)
Hazard ratio (95% CI)	0.51 (0.35, 0.73)		0.92 (0.82, 1.02)	
p-value	0.0002		0.0965	

The modest favorable trend in the group receiving an ACE inhibitor was largely driven by the patients receiving less than the recommended dose of ACE inhibitor. Thus, there is little evidence of further clinical benefit when valsartan is added to an adequate dose of ACE inhibitor.

Secondary end points in the subgroup not receiving ACE inhibitors were as follows.

	Placebo (N=181)	Valsartan (N=185)	Hazard Ratio (95% CI)
Components of HF morbidity			
All-cause mortality	49 (27.1%)	32 (17.3%)	0.59 (0.37, 0.91)
Sudden death with resuscitation	2 (1.1%)	1 (0.5%)	0.47 (0.04, 5.20)

CHF therapy	1 (0.6%)	0 (0.0%)	–
CHF hospitalization	48 (26.5%)	24 (13.0%)	0.43 (0.27, 0.71)
Cardiovascular mortality	40 (22.1%)	29 (15.7%)	0.65 (0.40, 1.05)
Non-fatal morbidity	49 (27.1%)	24 (13.0%)	0.42 (0.26, 0.69)

In patients not receiving an ACE inhibitor, valsartan-treated patients had an increase in ejection fraction and reduction in left ventricular internal diastolic diameter (LVIDD).

Effects were generally consistent across subgroups defined by age and gender for the population of patients not receiving an ACE inhibitor. The number of black patients was small and does not permit a meaningful assessment in this subset of patients.

14.3 Post-Myocardial Infarction

The VALsartan In Acute myocardial iNfarcTion trial (VALIANT) was a randomized, controlled, multinational, double-blind study in 14,703 patients with acute myocardial infarction and either heart failure (signs, symptoms or radiological evidence) or left ventricular systolic dysfunction (ejection fraction $\leq 40\%$ by radionuclide ventriculography or $\leq 35\%$ by echocardiography or ventricular contrast angiography). The VALIANT study was conducted with a formulation of valsartan that is not therapeutically equivalent to PREXXARTAN™ [see *Clinical Pharmacology* (12.3)]. Patients were randomized within 12 hours to 10 days after the onset of myocardial infarction symptoms to one of three treatment groups: valsartan (titrated from 20 or 40 mg twice daily to the highest tolerated dose up to a maximum of 160 mg twice daily), the ACE inhibitor, captopril (titrated from 6.25 mg three times daily to the highest tolerated dose up to a maximum of 50 mg three times daily), or the combination of valsartan plus captopril. In the combination group, the dose of valsartan was titrated from 20 mg twice daily to the highest tolerated dose up to a maximum of 80 mg twice daily; the dose of captopril was the same as for monotherapy. The population studied was 69% male, 94% Caucasian, and 53% were 65 years of age or older. Baseline therapy included aspirin (91%), beta-blockers (70%), ACE inhibitors (40%), thrombolytics (35%) and statins (34%). The mean treatment duration was 2 years. The mean daily dose of Valsartan in the monotherapy group was 217 mg.

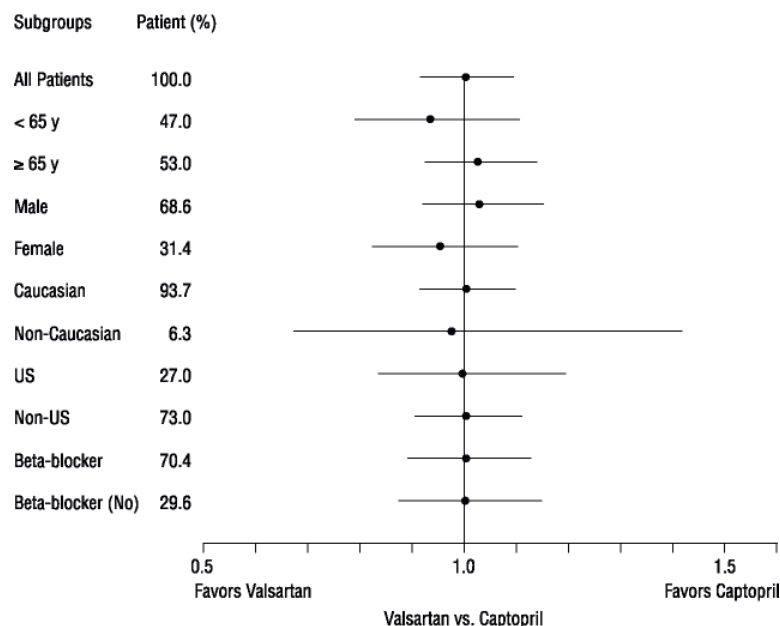
The primary endpoint was time to all-cause mortality. Secondary endpoints included (1) time to cardiovascular (CV) mortality, and (2) time to the first event of cardiovascular mortality, reinfarction, or hospitalization for heart failure. The results are summarized in the following table.

	Valsartan vs. Captopril (N=4,909) (N=4,909)			Valsartan + Captopril vs. Captopril (N=4,885) (N=4,909)		
	No. of Deaths Valsartan/Captopril	Hazard Ratio CI	p-value	No. of Deaths Comb/Captopril	Hazard Ratio CI	p-value
All-cause mortality	979 (19.9%) /958 (19.5%)	1.001 (0.902, 1.111)	0.98	941 (19.3%) /958 (19.5%)	0.984 (0.886, 1.093)	0.73
CV mortality	827 (16.8%) /830 (16.9%)	0.976 (0.875, 1.090)				
CV mortality, hospitalization for HF, and recurrent non-fatal MI	1,529 (31.1%) /1,567 (31.9%)	0.955 (0.881, 1.035)				

There was no difference in overall mortality among the three treatment groups. There was thus no evidence that combining the ACE inhibitor captopril and the angiotensin II blocker valsartan was of value.

The data were assessed to see whether the effectiveness of valsartan could be demonstrated by showing in a non-inferiority analysis that it preserved a fraction of the effect of captopril, a drug with a demonstrated survival effect in this setting. A conservative estimate of the effect of captopril (based on a pooled analysis of 3 post-infarction studies of captopril and 2 other ACE inhibitors) was a 14% to 16% reduction in mortality compared to placebo. Valsartan would be considered effective if it preserved a meaningful fraction of that effect and unequivocally preserved some of that effect. As shown in the table, the upper bound of the CI for the hazard ratio (valsartan/captopril) for overall or CV mortality is 1.09 to 1.11, a difference of about 9% to 11%, thus making it unlikely that valsartan has less than about half of the estimated effect of captopril and clearly demonstrating an effect of valsartan. The other secondary endpoints were consistent with this conclusion.

Effects on Mortality Amongst Subgroups in VALIANT



There were no clear differences in all-cause mortality based on age, gender, race, or baseline therapies, as shown in the figure above.

16 HOW SUPPLIED/STORAGE AND HANDLING

PREXXARTAN™ (valsartan) Oral Solution contains 4 mg/mL valsartan for oral administration and is supplied as a clear, colorless, grape-flavored solution. PREXXARTAN™ is packaged in bottles containing 473 mL, bottles containing 120 mL and unit dose cups containing 20 mL.

White HDPE bottles of 473 mL: NDC 76281-0155-82

White HDPE bottles of 120 mL: NDC 76281-0155-64

20 mL Unit Dose cups: NDC 76281-0155-56

Unit Dose Tray (10 Unit Dose Cups): NDC 76281-0155-99

Store at 20°C-25°C (68 °F-77 °F); excursions permitted to 15°C -30°C (59 °F -86 °F) [see USP Controlled Room Temperature].

Dispense in tight container (USP).

17 PATIENT COUNSELING INFORMATION

Pregnancy: Advise female patients of childbearing age about the consequences of exposure to PREXXARTAN™ during pregnancy. Discuss treatment options with women planning to become pregnant. Ask patients to report pregnancies to their healthcare provider as soon as possible [*see Warnings and Precautions (5.1), Use in Specific Populations (8.1)*].

Lactation: Advise women not to breastfeed during treatment with PREXXARTAN™ [*see Use in Specific Populations (8.2)*].

Symptomatic Hypotension: Advise patients that lightheadedness can occur, especially during the first days of therapy, and that it should be reported to their healthcare provider. Tell patients that if syncope occurs to discontinue PREXXARTAN™ until the physician has been consulted.

Caution all patients that inadequate fluid intake, excessive perspiration, diarrhea, or vomiting can lead to an excessive fall in blood pressure, with the same consequences of lightheadedness and possible syncope [*see Warnings and Precautions (5.2)*].

Hyperkalemia: Advise patients not to use salt substitutes without consulting their healthcare provider [*see Drug Interactions (7.1)*].

Manufactured by:
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1533 Union Ave,
Pennsauken, NJ 08110

For:
Carmel Biosciences, Inc.
5671 Peachtree Dunwoody Road
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PI-1011

PREXXARTAN™ (prex ar' tan) (valsartan)
Oral Solution 4 mg/mL

Read the Patient Information that comes with **PREXXARTAN™** before you take it and each time you get a refill. There may be new information. This leaflet does not take the place of talking with your doctor about your medical condition or treatment. If you have any questions about **PREXXARTAN™**, ask your doctor or pharmacist.

What is the most important information I should know about PREXXARTAN™?

PREXXARTAN™ can cause harm or death to an unborn baby. Talk to your doctor about other ways to lower your blood pressure if you plan to become pregnant. If you get pregnant while taking PREXXARTAN™, tell your doctor right away.

What is PREXXARTAN™?

PREXXARTAN™ is a prescription medicine, containing the active substance valsartan, and belongs to a class of drugs called angiotensin II receptor blockers (ARBs). **PREXXARTAN™** is used to:

- Lower high blood pressure (hypertension) in adults and children six years and older.
- Treat heart failure for patients who are unable to swallow valsartan tablets. In these patients, **PREXXARTAN™** may reduce the need for hospitalization related to heart failure.
- Improve the chance of living longer after a heart attack (myocardial infarction) for patients who are unable to swallow valsartan tablets.

PREXXARTAN™ is not recommended for patients under 6 years of age.

High Blood Pressure (Hypertension): Blood pressure is the force in your blood vessels when your heart beats and when your heart rests. You have high blood pressure when the force is too much. **PREXXARTAN™** can help your blood vessels relax so your blood pressure is lower. Medicines that lower your blood pressure lower your chance of having a stroke or heart attack. High blood pressure makes the heart work harder to pump blood throughout the body and causes damage to the blood vessels. If high blood pressure is not treated, it can lead to stroke, heart attack, heart failure, kidney failure and vision problems.

Heart Failure: Heart failure occurs when the heart is weak and cannot pump enough blood to your lungs and the rest of your body. Just walking or moving can make you short of breath, so you may have to rest a lot.

Heart Attack (Myocardial Infarction): A heart attack is caused by a blocked artery that results in damage to the heart muscle.

What should I tell my doctor before taking PREXXARTAN™?

Tell your doctor about all your medical conditions including whether you:

- have any allergies. See the end of this leaflet for a complete list of ingredients in **PREXXARTAN™**.
- have a heart condition
- have liver problems
- have kidney problems
- **are pregnant or planning to become pregnant. See “What is the most important information I should know about PREXXARTAN™?”**
- are breastfeeding. It is not known if **PREXXARTAN™** passes into your breast milk, its effects on the breastfed infant, or the effects on milk production. You and your doctor should decide if you will take **PREXXARTAN™** or breastfeed, but not both. Talk to your doctor about the best way to feed your baby if you take **PREXXARTAN™**.
- have ever had a reaction called angioedema, to another blood pressure medicine. Angioedema causes swelling of the face, lips, tongue, and/or throat, and may cause difficulty breathing.

Tell your doctor about all the medicines you take including prescription and nonprescription medicines, vitamins and herbal supplements. Especially tell your doctor if you take:

- other medicines for high blood pressure or a heart problem
- water pills (also called “diuretics”)
- potassium supplements. Your doctor may check the amount of potassium in your blood periodically.
- a salt substitute. Your doctor may check the amount of potassium in your blood periodically.
- nonsteroidal anti-inflammatory drugs (like ibuprofen or naproxen)
- certain antibiotics (rifamycin group), a drug used to protect against transplant rejection (cyclosporine) or an antiretroviral drug used to treat HIV/AIDS infection (ritonavir). These drugs may increase the effect of valsartan.
- lithium, a medicine used in some types of depression

Know the medicines you take. Keep a list of your medicines with you to show to your doctor and pharmacist when a new medicine is prescribed. Talk to your doctor or pharmacist before you start taking any new medicine. Your doctor or pharmacist will know what medicines are safe to take together.

How should I take PREXXARTAN™?

- The recommended dosing with PREXXARTAN™ oral solution is not the same as the recommended dosing with valsartan tablets. It is important that you take **PREXXARTAN™** exactly as prescribed by your doctor.
- For treatment of high blood pressure, take **PREXXARTAN™** two times each day, at the same time each day.
- For adult patients with heart failure or who have had a heart attack, take **PREXXARTAN™** two times each day, at the same time each day. Your doctor may start you on a low dose of **PREXXARTAN™** and may increase the dose during your treatment.
- **PREXXARTAN™** can be taken with or without food.
- If you miss a dose, take it as soon as you remember. If it is close to your next dose, do not take the missed dose. Take the next dose at your regular time.
- If you take too much **PREXXARTAN™**, call your doctor or Poison Control Center, or go to the nearest hospital emergency room.

What are the possible side effects of PREXXARTAN™?

PREXXARTAN™ may cause the following serious side effects: Injury or death to an unborn baby. See “What is the most important information I should know about PREXXARTAN™?”

Low Blood Pressure (Hypotension)

Low blood pressure is most likely to happen if you also take water pills, are on a low-salt diet, get dialysis treatments, have heart problems, or get sick with vomiting or diarrhea. If you have heart failure or have had a heart attack, **PREXXARTAN™** should only be taken if you are unable to swallow valsartan tablets. Lie down if you feel faint or dizzy. Call your doctor right away.

Kidney Problems

Kidney problems may get worse if you already have kidney disease. Some patients will have changes in blood tests for kidney function and may need a lower dose of **PREXXARTAN™**. Call your doctor if you get swelling in your feet, ankles, or hands, or unexplained weight gain. If you have heart failure, your doctor should check your kidney function before prescribing **PREXXARTAN™**.

Increased Potassium

If you have heart failure, increases in potassium may occur, especially if you have kidney problems.

The most common side effects of PREXXARTAN™ used to treat people with high blood pressure include:

- headache
- dizziness
- flu symptoms
- tiredness
- stomach (abdominal) pain

Side effects were generally mild and brief. They generally have not caused patients to stop taking **PREXXARTAN™**.

The most common side effects of PREXXARTAN™ used to treat people with heart failure include:

- dizziness
- low blood pressure
- diarrhea
- joint and back pain
- tiredness
- high blood potassium

Common side effects of PREXXARTAN™ used to treat people after a heart attack which caused them to stop taking the drug include:

- low blood pressure
- cough
- high blood creatinine (decreased kidney function)
- rash

Tell your doctor if you get any side effects that bothers you or that does not go away.

These are not all the possible side effects of **PREXXARTAN™**. For a complete list, ask your doctor or pharmacist.

How do I store PREXXARTAN™?

- Store **PREXXARTAN™** oral solution at room temperature between 20°C to 25°C (68°F to 77°F). Once opened, the bottle can be stored for up to 90 days.
- Keep **PREXXARTAN™** oral solution in a closed container in a dry place.
- Keep **PREXXARTAN™** and all medicines out of the reach of children.

General information about PREXXARTAN™

Medicines are sometimes prescribed for conditions that are not mentioned in patient information leaflets. Do not **use PREXXARTAN™** for a condition for which it was not prescribed. Do not give **PREXXARTAN™** to other people even if they have the same symptoms you have. It may harm them.

This leaflet summarizes the most important information about **PREXXARTAN™**. If you would like more information, talk to your doctor. You can ask your doctor or pharmacist for information about **PREXXARTAN™** that is written for health professionals.

For more information about **PREXXARTAN™**, ask your pharmacist or doctor, visit the web www.PREXXARTAN.com, or call [\[1-732-314-4550\]](tel:1-732-314-4550).

What are the ingredients in PREXXARTAN™?

Active ingredient: valsartan

Inactive ingredients: grape flavor, methylparaben NF, poloxamer 188, potassium sorbate, propylene glycol NF, purified water USP, sodium citrate dihydrate USP, and sucralose NF.

Manufactured by:

AptaPharma, Inc.
Pennsauken, NJ 08110 USA

For:

Carmel Biosciences, Inc.
Atlanta, GA 30342 USA

PI-1011

2.1770"

4.75"

Recommended Dosage: See Prescribing Information.

NDC: 76281-0155-64 120 mL

Rx Only

Pharmacist: Dispense in a tight container.

Store at 20°C to 25°C (68°F to 77°F). Excursions permitted to 15°C - 30°C (59°F - 86°F). [See USP Controlled Room Temperature]

Inactive Ingredients: Grape Flavor, Methylparaben, Poloxamer, Potassium Sorbate, Propylene Glycol, Purified water, Sodium Citrate Dihydrate and Sucralose

Each 1 mL contains 4 mg Valsartan, USP

Manufactured by:
AptaPharma, Inc.
for Carmel Biosciences, Inc.
Atlanta, GA 30342

Prexxartan™
(valsartan) **Oral Solution**
20 mg/5 mL
(4 mg/mL)



LR-1015

No Varnish Area

GTIN: XXXXXXXXXX
SR: XXXXXXXXXX
LOT: XXXXXXX
EXP: YYYY-MM



7.00"

3.00"

Recommended Dosage: See Prescribing Information.

Pharmacist: Dispense in a tight container.

Store at 20°C to 25°C (68°F to 77°F). Excursions permitted to 15°C - 30°C (59°F - 86°F). [See USP Controlled Room Temperature]

Inactive Ingredients: Grape Flavor, Methylparaben, Poloxamer, Potassium Sorbate, Propylene Glycol, Purified water, Sodium Citrate Dihydrate and Sucralose

Each 1 mL contains 4 mg Valsartan, USP

NDC: 76281-0155-82 473 mL

Rx Only

Prexxartan™
(valsartan)

20 mg/5 mL
(4 mg/mL)

Oral Solution

Manufactured by:
AptaPharma, Inc.
for Carmel Biosciences, Inc.
Atlanta, GA 30342



LR-1014

No Varnish Area

GTIN: XXXXXXXXXXXXX
SN: XXXXXXXXXXXXX
LOT: XXXXXX
EXP: YYYY-MM

10.0"

4.0"

Recommended Dosage: See Prescribing Information.

Pharmacist: Dispense in a tight container.

Store at 20°C to 25°C (68°F to 77°F). Excursions permitted to 15°C - 30°C (59°F - 86°F). [See USP Controlled Room Temperature]

Inactive Ingredients: Grape Flavor, Methylparaben, Poloxamer, Potassium Sorbate, Propylene Glycol, Purified water, Sodium Citrate Dihydrate and Sucralose

Each 20 mL cup contains 80 mg Valsartan, USP

NDC: 76281-0155-99 20 mL

10 Unit Dose Cups

Prexxartan™

(valsartan)

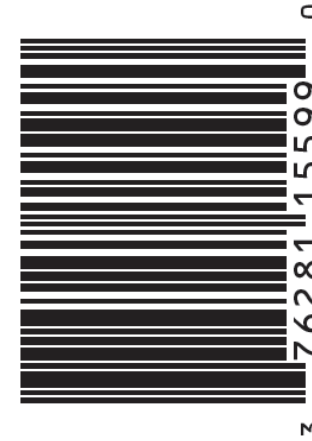
**80 mg/20 mL
(4 mg/mL)**

**Oral
Solution**

Manufactured by:
AptaPharma, Inc.
for Carmel Biosciences, Inc.
Atlanta, GA 30342

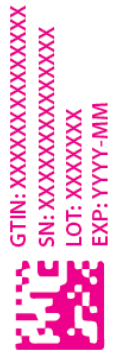


Rx Only



LR-1017

No Varnish Area



(b) (4)



Joyce
Crich

Digitally signed by Joyce Crich

Date: 9/22/2026 02:24:53PM

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