

BOSENTAN- bosentan tablet, for suspension

Zydus Pharmaceuticals USA Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use **BOSENTAN TABLETS FOR ORAL SUSPENSION** safely and effectively. See full prescribing information for **BOSENTAN TABLETS FOR ORAL SUSPENSION**.

BOSENTAN tablets for oral suspension

Initial U.S. Approval: 2001

WARNING: RISKS OF HEPATOTOXICITY and EMBRYO -FETAL TOXICITY

See full prescribing information for complete boxed warning.

Bosentan is available only through a restricted distribution program called the Bosentan Risk Evaluation and Mitigation Strategy (REMS) because of the risks of hepatotoxicity (5.2):

Elevations of liver aminotransferases (ALT, AST) and liver failure have been reported with bosentan (5.1).

- **Measure liver aminotransferases prior to initiation of treatment and then monthly (2.1, 5.1).**
- **Discontinue bosentan if aminotransferase elevations are accompanied by signs or symptoms of liver dysfunction or injury or increases in bilirubin $\geq 2 \times$ ULN (2.4, 5.1).**

Based on animal data, bosentan may cause fetal harm if used during pregnancy (4.1, 5.3, 8.1).

- **Females of reproductive potential: Exclude pregnancy before initiating treatment. Use effective contraception prior to initiation of treatment, during treatment and for one month after stopping bosentan (2.1, 4.1, 5.3, 8.1, 8.3).**
- **When pregnancy is detected, discontinue bosentan as soon as possible (5.3).**

RECENT MAJOR CHANGES

Boxed Warning	11/2025
Dosage and Administration (2.1)	11/2025
Contraindications (4.1)	11/2025
Warnings and Precautions (5.1, 5.2, 5.3)	11/2025

INDICATIONS AND USAGE

Bosentan is an endothelin receptor antagonist indicated for the treatment of pulmonary arterial hypertension (PAH) (WHO Group 1):

- in adults to improve exercise ability and to decrease clinical worsening. Studies establishing effectiveness included predominantly patients with WHO Functional Class II-IV symptoms and etiologies of idiopathic or heritable PAH (60%), PAH associated with connective tissue diseases (21%) and PAH associated with congenital heart disease with left-to-right shunts (18%) (1).
- in pediatric patients aged 3 years and older with idiopathic or congenital PAH to improve pulmonary vascular resistance (PVR), which is expected to result in an improvement in exercise ability (1).

DOSAGE AND ADMINISTRATION

- Patients older than 12 years of age: initiate at 62.5 mg orally twice daily; for patients weighing greater than 40 kg, increase to 125 mg orally twice daily after 4 weeks (2.2).
- Patients 12 years of age and younger: dosage is based on weight, see Table 1 (2.2).
- Reduce the dose and closely monitor patients developing aminotransferase elevations more than 3 X Upper Limit of Normal (ULN) (2.1).

DOSAGE FORMS AND STRENGTHS

- Tablet for oral suspension: 32 mg (3)

CONTRAINDICATIONS

- Pregnancy (4.1)
- Use with Cyclosporine A (4.2)
- Use with Glyburide (4.3)
- Hypersensitivity (4.4)

WARNINGS AND PRECAUTIONS

- Fluid retention: May require intervention (5.4).
- Pulmonary veno-occlusive disease (PVOD): If signs of pulmonary edema occur, consider the diagnosis of associated PVOD and consider discontinuing bosentan (5.5).
- Decreased sperm counts (5.6).
- Decreases in hemoglobin and hematocrit: Monitor hemoglobin levels after 1 and 3 months of treatment, then every 3 months thereafter (5.7).

ADVERSE REACTIONS

- Common adverse reactions ($\geq 3\%$ more than placebo) for the film-coated tablet are respiratory tract infection and anemia (6.1).
- Common adverse reactions ($\geq 15\%$) for the dispersible tablet are upper respiratory tract infections and pyrexia (6.1).

To report SUSPECTED ADVERSE REACTIONS, contact Zydus Pharmaceuticals (USA) Inc. at 1-877-993-8779 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Cytochrome P450: Coadministration of bosentan with drugs metabolized by CYP2C9 and CYP3A can increase exposure to bosentan and/or the coadministered drug (4.2, 4.3, 7.1).
- Hormonal contraceptives: bosentan use decreases contraceptive exposure and reduces effectiveness (7.2).

USE IN SPECIFIC POPULATIONS

- Nursing mothers: Choose breastfeeding or bosentan (8.2).

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.

Revised: 11/2025

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FULL PRESCRIBING INFORMATION

WARNING: RISKS OF HEPATOTOXICITY and EMBRYO-FETAL TOXICITY

Because of the risk of hepatotoxicity, bosentan is available only through a restricted program called the Bosentan Risk Evaluation and Mitigation Strategy (REMS). Under the Bosentan REMS, prescribers, patients and pharmacies must enroll in the program [*see Warnings and Precautions (5.2)*].

Hepatotoxicity

In clinical studies, bosentan caused at least 3-fold upper limit of normal (ULN) elevation of liver aminotransferases (ALT and AST) in about 11% of patients, accompanied by elevated bilirubin in a small number of cases. Because these changes are a marker for potential serious hepatotoxicity, serum aminotransferase levels must be measured prior to initiation of treatment and then monthly [*see Dosage and Administration (2.4), Warnings and Precautions (5.1)*]. In the postmarketing period, in the setting of close monitoring, rare cases of unexplained hepatic cirrhosis were reported after prolonged (> 12 months) therapy with bosentan in patients with multiple comorbidities and drug therapies. There have also been reports of liver failure. The contribution of bosentan in these cases could not be excluded.

In at least one case, the initial presentation (after > 20 months of treatment) included pronounced elevations in aminotransferases and bilirubin levels accompanied by non-specific symptoms, all of which resolved slowly over time after discontinuation of bosentan. This case reinforces the importance of strict adherence to the monthly monitoring schedule for the duration of treatment and the treatment algorithm, which includes stopping bosentan with a rise of aminotransferases accompanied by signs or symptoms of liver dysfunction [*see Dosage and Administration (2.4)*].

Elevations in aminotransferases require close attention [*see Dosage and Administration (2.4)*]. Bosentan should generally be avoided in patients with elevated aminotransferases (> 3 x ULN) at baseline because monitoring for hepatotoxicity may be more difficult. If liver aminotransferase elevations are accompanied by clinical symptoms of hepatotoxicity (such as nausea, vomiting, fever, abdominal pain, jaundice, or unusual lethargy or fatigue) or increases in bilirubin $\geq 2 \times$ ULN, treatment with bosentan should be stopped. There is no experience with the reintroduction of bosentan in these circumstances.

Embryo-Fetal Toxicity

Bosentan is contraindicated for use during pregnancy because it may cause fetal harm if used by pregnant females based on animal data. Therefore, for females of reproductive potential, exclude pregnancy before the start of treatment with bosentan. Advise use of effective contraception before initiation, during treatment and for one month after stopping bosentan. When pregnancy is detected, discontinue bosentan as soon as possible [*see Dosage and Administration (2.1), Contraindications (4.1), Warnings and Precautions (5.3), Drug*

1 INDICATIONS AND USAGE

Bosentan tablets for oral suspension are indicated for the treatment of pulmonary arterial hypertension (PAH) (WHO Group 1):

- in adults to improve exercise ability and to decrease clinical worsening. Studies establishing effectiveness included predominantly patients with WHO Functional Class II-IV symptoms and etiologies of idiopathic or heritable PAH (60%), PAH associated with connective tissue diseases (21%) and PAH associated with congenital heart disease with left-to-right shunts (18%) [see *Clinical Studies (14.1)*].
- in pediatric patients aged 3 years and older with idiopathic or congenital PAH to improve pulmonary vascular resistance (PVR), which is expected to result in an improvement in exercise ability.

2 DOSAGE AND ADMINISTRATION

2.1 Required Monitoring

Healthcare professionals who prescribe bosentan must enroll in the Bosentan REMS and must comply with the required monitoring to minimize the risks associated with bosentan [see *Warnings and Precautions (5.2)*].

Measure liver aminotransferase levels prior to initiation of treatment and then monthly [see *Boxed Warning, Warnings and Precautions (5.1, 5.2)*].

Exclude pregnancy before initiating treatment with bosentan in females of reproductive potential [see *Boxed Warning, Contraindications (4.1), Warnings and Precautions (5.3), Use in Specific Populations (8.1, 8.3)*].

2.2 Recommended Dosage

Administer bosentan orally following the dosing recommendations in Table 1. Doses above 125 mg twice daily did not appear to confer additional benefit sufficient to offset the increased risk of hepatotoxicity.

Table 1 Dosing Recommendations

	Initial 4 weeks	Maintenance (after 4 weeks)
Patients > 12 years of age and > 40 kg	62.5 mg twice daily	125 mg twice daily
Patients > 12 years of age and < 40 kg	62.5 mg twice daily	62.5 mg twice daily
Patients ≤ 12 years of age		
≥ 4 to 8 kg	16 mg twice daily	16 mg twice daily
> 8 to 16 kg	32 mg twice daily	32 mg twice daily
> 16 to 24 kg	48 mg twice daily	48 mg twice daily
> 24 to 40 kg	64 mg twice daily	64 mg twice daily

2.3 Administration

Bosentan tablets for oral suspension (dispersible tablets) should be administered orally twice daily.

Disperse tablets for oral suspension, or dispersible tablet half, in a minimal amount of water immediately before administration.

Store divided dispersible tablet pieces at 20°C to 25°C (68°F to 77°F) in the opened blister for up to 7 days.

2.4 Dosage Adjustments for Aminotransferase Elevations

If aminotransferase levels increase, adjust monitoring and treatment plan according to Table 2.

Discontinue bosentan if liver aminotransferase elevations are accompanied by clinical symptoms of hepatotoxicity (such as nausea, vomiting, fever, abdominal pain, jaundice, or unusual lethargy or fatigue) or bilirubin $\geq 2 \times$ Upper Limit of Normal (ULN). There is no experience with the reintroduction of bosentan in these circumstances.

Table 2 Dosage Adjustment and Monitoring in Patients Developing Aminotransferase Elevations > 3 x ULN

ALT/AST levels	Treatment and monitoring recommendations
> 3 and $\leq 5 \times$ ULN	Confirm by another aminotransferase test; if confirmed, -in <u>adults and pediatric patients > 12 years and > 40 kg</u> , reduce the daily dose to 62.5 mg twice daily or interrupt treatment and monitor aminotransferase levels at least every 2 weeks. If the aminotransferase levels return to pretreatment values, treatment may continue or be reintroduced at 62.5 mg twice daily, with reassessment of aminotransferase levels within 3 days. -in <u>all other pediatric patients</u> , interrupt treatment with no prior dose reduction. If the aminotransferase levels return to pretreatment values, reintroduce at the dose used prior to treatment interruption, with reassessment of aminotransferase levels within 3 days.
> 5 and $\leq 8 \times$ ULN	Confirm by another aminotransferase test; if confirmed, stop treatment and monitor aminotransferase levels at least every 2 weeks. Once the aminotransferase levels return to

	pretreatment values, -in adults and pediatric patients > 12 years and > 40 kg, consider reintroduction of treatment at 62.5 mg twice daily, with reassessment of aminotransferase levels within 3 days. -in all other pediatric patients, consider reintroduction at the dose used prior to treatment interruption, with reassessment of aminotransferase levels within 3 days.
> 8 x ULN	Stop treatment permanently. There is no experience with reintroduction of bosentan in these circumstances.

2.5 Use with Ritonavir

Co-administration of Bosentan in Patients on Ritonavir

In patients who have been receiving ritonavir for at least 10 days, start bosentan at the recommended initial dose once daily or every other day based upon individual tolerability [see *Drug Interactions (7.1)*].

Co-administration of Ritonavir in Patients on Bosentan

Discontinue use of bosentan at least 36 hours prior to initiation of ritonavir. After at least 10 days following the initiation of ritonavir, resume bosentan at the recommended initial dose once daily or every other day based upon individual tolerability [see *Drug Interactions (7.1)*].

2.6 Use in Patients with Pre-existing Hepatic Impairment

Avoid initiation of bosentan in patients with aminotransferases > 3 x ULN. No dose adjustment is required in patients with mildly impaired liver function [see *Warnings and Precautions (5.1)*, *Use in Specific Populations (8.6)*, *Clinical Pharmacology (12.3)*].

3 DOSAGE FORMS AND STRENGTHS

Bosentan tablets for oral suspension, 32 mg are off-white to pale yellow, round, flat, beveled edge tablet debossed with '1508' on one side and bisect on the other.

4 CONTRAINDICATIONS

4.1 Pregnancy

Use of bosentan is contraindicated in females who are pregnant [see *Boxed Warning, Dosage and Administration (2.1)*, *Warnings and Precautions (5.3)*, *Use in Specific Populations (8.1)*].

4.2 Use with Cyclosporine A

Co-administration of cyclosporine A and bosentan resulted in markedly increased plasma concentrations of bosentan. Therefore, concomitant use of bosentan and cyclosporine A is contraindicated [see *Drug Interactions (7.1)*].

4.3 Use with Glyburide

An increased risk of liver enzyme elevations was observed in patients receiving glyburide concomitantly with bosentan. Therefore co-administration of glyburide and bosentan is contraindicated [see *Drug Interactions (7.1)*].

4.4 Hypersensitivity

Bosentan is contraindicated in patients who are hypersensitive to bosentan or any component of the product. Observed reactions include Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS), anaphylaxis, rash and angioedema [see *Adverse Reactions (6.2), Description (11)*].

5 WARNINGS AND PRECAUTIONS

5.1 Hepatotoxicity

ALT or AST $> 3 \times$ ULN were observed in 11% of bosentan-treated patients (n=658) compared to 2% of placebo-treated patients (n=280). Three-fold increases were seen in 12% of 95 pulmonary arterial hypertension (PAH) patients on 125 mg twice daily and 14% of 70 PAH patients on 250 mg twice daily. Eight-fold increases were seen in 2% of PAH patients on 125 mg twice daily and 7% of PAH patients on 250 mg twice daily. Bilirubin increases to $\geq 3 \times$ ULN were associated with aminotransferase increases in 2 of 658 (0.3%) of patients treated with bosentan. In a pooled analysis of four pediatric studies conducted in PAH (n=100), elevations in liver aminotransferases $\geq 3 \times$ ULN were observed in 2% of patients. The combination of hepatocellular injury (increases in aminotransferases of $> 3 \times$ ULN) and increases in total bilirubin ($\geq 2 \times$ ULN) is a marker for potential serious hepatotoxicity.

Elevations of AST or ALT associated with bosentan are dose-dependent, occur both early and late in treatment, usually progress slowly, are typically asymptomatic and usually have been reversible after treatment interruption or cessation. Aminotransferase elevations also may reverse spontaneously while continuing treatment with bosentan.

Liver aminotransferase levels must be measured prior to initiation of treatment and then monthly and therapy adjusted accordingly [see *Dosage and Administration (2.1, 2.4)*]. Discontinue bosentan if liver aminotransferase elevations are accompanied by clinical symptoms of hepatotoxicity (such as nausea, vomiting, fever, abdominal pain, jaundice, or unusual lethargy or fatigue) or increases in bilirubin $\geq 2 \times$ ULN.

Avoid initiation of bosentan in patients with elevated aminotransferases ($> 3 \times$ ULN) prior to drug initiation because monitoring hepatotoxicity in these patients may be more difficult [see *Boxed Warning, Dosage and Administration (2.6), Use in Specific Populations (8.6)*].

In WHO Functional Class II patients, consider whether the benefits of bosentan are sufficient to offset the risk of hepatotoxicity, which may preclude future use as their disease progresses.

Bosentan is only available through a restricted program under REMS [see *Warnings and Precautions (5.2)*].

5.2 Bosentan REMS

Because of the risks of hepatotoxicity, bosentan is available only through a restricted program called the Bosentan REMS. As a component of the Bosentan REMS, prescribers, patients, and pharmacies must enroll in the program [see *Boxed Warning, Warnings and Precautions (5.1)*].

Required components of the Bosentan REMS are:

- Healthcare professionals who prescribe bosentan must review the prescriber educational materials, enroll in the Bosentan REMS and comply with its requirements.
- Healthcare professionals must (1) review serum aminotransferases (ALT/AST) and bilirubin, and agree to order and monitor these tests monthly.
- To receive bosentan, all patients must understand the risks and benefits and complete a patient enrollment form with their prescriber.
- Pharmacies that dispense bosentan must enroll in the program and agree to comply with the Bosentan REMS requirements.

Further information about bosentan and the Bosentan REMS is available at www.BosentanREMSProgram.com or 1-866-359-2612

5.3 Embryo-Fetal Toxicity

Based on data from animal reproduction studies, bosentan may cause fetal harm when administered to a pregnant female and is contraindicated in females who are pregnant. The available human data for endothelin receptor antagonists do not establish the presence or absence of major birth defects related to the use of bosentan. Advise females of reproductive potential about the potential risk to a fetus. Exclude pregnancy prior to bosentan treatment. Advise females of reproductive potential to use effective contraception prior to initiation of treatment with bosentan, during treatment, and for at least one month after the last dose. When pregnancy is detected, discontinue bosentan as soon as possible [see *Dosage and Administration (2.1)*, *Contraindications (4.1)*, *Drug Interactions (7.2)*, *Use in Specific Populations (8.1, 8.3)*].

5.4 Fluid Retention

Peripheral edema is a known clinical consequence of PAH and worsening PAH and is also a known effect of bosentan and other endothelin receptor antagonists. In PAH clinical trials with bosentan, combined adverse events of fluid retention or edema were reported in 1.7% (placebo-corrected) of patients.

In addition, there have been numerous postmarketing reports of fluid retention in patients with pulmonary hypertension occurring within weeks after starting bosentan. Patients required intervention with a diuretic, fluid management, or hospitalization for decompensating heart failure.

If clinically significant fluid retention develops, with or without associated weight gain, further evaluation should be undertaken to determine the cause, such as bosentan or underlying heart failure and the possible need for treatment or discontinuation of

bosentan [see *Adverse Reactions (6.1), Clinical Studies (14.2)*].

5.5 Pulmonary Veno-Occlusive Disease

If signs of pulmonary edema occur, consider the possibility of associated pulmonary veno-occlusive disease and consider whether bosentan should be discontinued.

5.6 Decreased Sperm Counts

Decreased sperm counts have been observed in patients receiving bosentan. Preclinical data also suggest that bosentan, similar to other endothelin receptor antagonists, may have an adverse effect on spermatogenesis [see *Adverse Reactions (6.1), Nonclinical Toxicology (13.1)*].

5.7 Decreases in Hemoglobin and Hematocrit

Treatment with bosentan can cause a dose-related decrease in hemoglobin and hematocrit. There have been postmarketing reports of decreases in hemoglobin concentration and hematocrit that have resulted in anemia requiring transfusion. It is recommended that hemoglobin concentrations be checked after 1 and 3 months and every 3 months thereafter. If a marked decrease in hemoglobin concentration occurs, further evaluation should be undertaken to determine the cause and need for specific treatment [see *Adverse Reactions (6.1)*].

6 ADVERSE REACTIONS

The following important adverse reactions are described elsewhere in the labeling:

- Hepatotoxicity [see *Boxed Warning, Warnings and Precautions (5.1)*]
- Embryo-fetal Toxicity [see *Boxed Warning, Warnings and Precautions (5.3)*]
- Fluid Retention [see *Warnings and Precautions (5.4)*]

6.1 Clinical Trials Experience

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

Safety data on bosentan were obtained from 13 clinical studies (9 placebo-controlled and 4 open-label) in 870 adult patients with PAH and other diseases. Doses up to 8 times the currently recommended clinical dose (125 mg twice daily) were administered for a variety of durations. The exposure to bosentan in these trials ranged from 1 day to 4.1 years (n=94 for 1 year; n=61 for 1.5 years; and n=39 for more than 2 years). Exposure of PAH patients (n=328) to bosentan ranged from 1 day to 1.7 years (n=174 more than 6 months and n=28 more than 12 months).

Treatment discontinuations due to adverse events other than those related to pulmonary hypertension during the clinical trials in adult patients with PAH were more frequent on bosentan (6%; 15/258 patients) than on placebo (3%; 5/172 patients). In this database the only cause of discontinuations > 1% and occurring more often on bosentan was abnormal liver function.

The adverse drug events that occurred in $\geq 3\%$ of the bosentan-treated patients and were more common on bosentan in placebo-controlled trials in PAH at doses of 125 mg

or 250 mg twice daily are shown in Table 3:

Table 3 Adverse events* occurring in \geq 3% of patients treated with Bosentan 125 mg to 250 mg twice daily and more common on Bosentan in placebo-controlled studies in pulmonary arterial hypertension

Adverse Event	Bosentan n = 258		Placebo n = 172	
	No.	%	No.	%
Respiratory Tract Infection**	56	22%	30	17%
Headache	39	15%	25	14%
Edema	28	11%	16	9%
Chest Pain	13	5%	8	5%
Syncope	12	5%	7	4%
Flushing	10	4%	5	3%
Hypotension	10	4%	3	2%
Sinusitis	9	4%	4	2%
Arthralgia	9	4%	3	2%
Serum Aminotransferases, abnormal	9	4%	3	2%
Palpitations	9	4%	3	2%
Anemia	8	3%	-	-

*Note: only AEs with onset from start of treatment to 1 calendar day after end of treatment are included. All reported events (at least 3%) are included except those too general to be informative and those not reasonably associated with the use of the drug because they were associated with the condition being treated or are very common in the treated population.

**Respiratory Tract Infection combines the terms "Nasopharyngitis", "Upper Respiratory Tract Infection" and "Respiratory Tract Infection".

Combined data from Study 351, BREATHE-1 and EARLY

Bosentan was evaluated for safety in 119 pediatric patients in uncontrolled studies. The safety profile was similar to that observed in adult patients with PAH.

Decreased Sperm Counts

An open-label, single-arm, multicenter, safety study evaluated the effect on testicular function of bosentan 62.5 mg twice daily for 4 weeks, followed by 125 mg twice daily for 5 months. Twenty-five male patients with WHO functional class III and IV PAH and normal baseline sperm count were enrolled. Twenty-three completed the study and 2 discontinued due to adverse events not related to testicular function. There was a decline in sperm count of at least 50% in 25% of the patients after 3 or 6 months of treatment with bosentan. Sperm count remained within the normal range in all 22 patients with data after 6 months and no changes in sperm morphology, sperm motility, or hormone levels were observed. One patient developed marked oligospermia at 3 months and the sperm count remained low with 2 follow-up measurements over the subsequent 6 weeks. Bosentan was discontinued and after 2 months the sperm count had returned to baseline levels. Based on these findings and preclinical data from endothelin receptor antagonists, it cannot be excluded that endothelin receptor

antagonists such as bosentan have an adverse effect on spermatogenesis.

Decreases in Hemoglobin and Hematocrit

Treatment with bosentan can cause a dose-related decrease in hemoglobin and hematocrit. It is recommended that hemoglobin concentrations be checked after 1 and 3 months and every 3 months thereafter. If a marked decrease in hemoglobin concentration occurs, further evaluation should be undertaken to determine the cause and need for specific treatment.

The overall mean decrease in hemoglobin concentration for adult bosentan-treated patients was 0.9 g/dL (change to end of treatment). Most of this decrease of hemoglobin concentration was detected during the first few weeks of bosentan treatment and hemoglobin levels stabilized by 4 weeks to 12 weeks of bosentan treatment. In placebo-controlled studies of all uses of bosentan, marked decreases in hemoglobin (> 15% decrease from baseline resulting in values < 11 g/dL) were observed in 6% of bosentan-treated patients and 3% of placebo-treated patients. In patients with PAH treated with doses of 125 mg and 250 mg twice daily, marked decreases in hemoglobin occurred in 3% compared to 1% in placebo-treated patients.

A decrease in hemoglobin concentration by at least 1 g/dL was observed in 57% of bosentan-treated patients as compared to 29% of placebo-treated patients. In 80% of those patients whose hemoglobin decreased by at least 1 g/dL, the decrease occurred during the first 6 weeks of bosentan treatment.

During the course of treatment, the hemoglobin concentration remained within normal limits in 68% of bosentan-treated patients compared to 76% of placebo patients. The explanation for the change in hemoglobin is not known, but it does not appear to be hemorrhage or hemolysis.

In a pooled analysis of pediatric patients (N=100) with PAH treated with bosentan, a decrease in hemoglobin levels to < 10 g/dL from baseline was reported in 11% of patients. There was no decrease to < 8 g/dL.

6.2 Postmarketing Experience

There have been several postmarketing reports of angioedema associated with the use of bosentan. The onset of the reported cases occurred within a range of 8 hours to 21 days after starting therapy. Some patients were treated with an antihistamine and their signs of angioedema resolved without discontinuing bosentan.

The following additional adverse reactions have been reported during the postapproval use of bosentan. Because these adverse reactions are reported from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to bosentan exposure:

- Unexplained hepatic cirrhosis [*see Boxed Warning*]
- Liver failure [*see Boxed Warning*]
- Hypersensitivity, DRESS and anaphylaxis [*see Contraindications (4.4)*]
- Thrombocytopenia
- Rash
- Jaundice
- Anemia requiring transfusion
- Neutropenia and leukopenia

- Nasal congestion
- Autoimmune hepatitis

7 DRUG INTERACTIONS

7.1 Cytochrome P450 Drug Interactions

Bosentan is metabolized by CYP2C9 and CYP3A. Inhibition of these enzymes may increase the plasma concentration of bosentan [see *Clinical Pharmacology (12.3)*]. Concomitant administration of both a CYP2C9 inhibitor (such as fluconazole or amiodarone) and a strong CYP3A inhibitor (e.g., ketoconazole, itraconazole) or a moderate CYP3A inhibitor (e.g., amprenavir, erythromycin, fluconazole, diltiazem) with bosentan will likely lead to large increases in plasma concentrations of bosentan. Co-administration of such combinations of a CYP2C9 inhibitor plus a strong or moderate CYP3A inhibitor with bosentan is not recommended.

Bosentan is an inducer of CYP3A and CYP2C9. Consequently plasma concentrations of drugs metabolized by these two isozymes will be decreased when bosentan is co-administered. Bosentan had no relevant inhibitory effect on any CYP isozyme *in vitro* (CYP1A2, CYP2C9, CYP2C19, CYP2D6, CYP3A). Consequently, bosentan is not expected to increase the plasma concentrations of drugs metabolized by these enzymes.

Figure 1

CYP3A induction-mediated effect of bosentan on other drugs

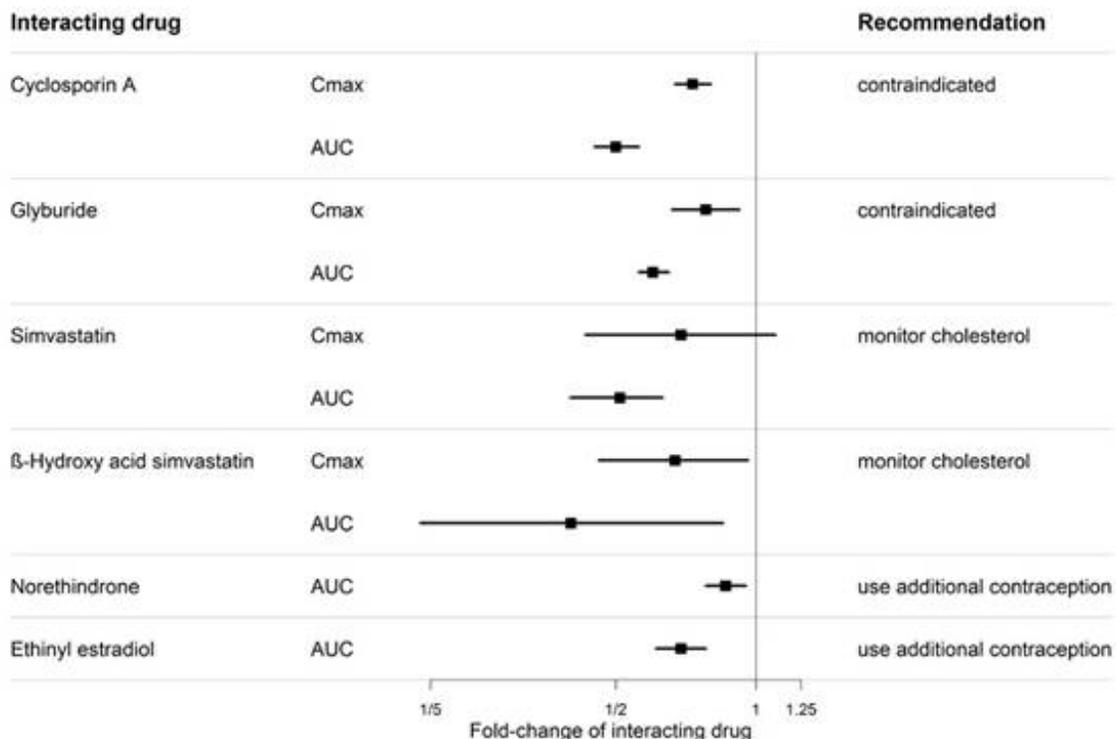
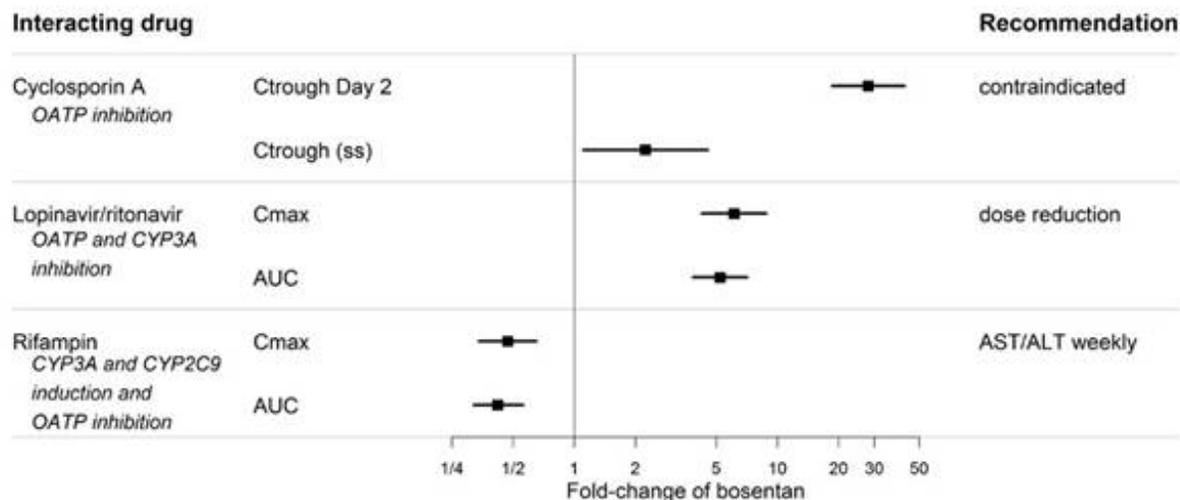


Figure 2

Effect of other drugs on bosentan



7.2 Hormonal Contraceptives

Hormonal contraceptives, including oral, injectable, transdermal and implantable forms, may not be reliable when bosentan is co-administered. Females should practice additional methods of contraception and not rely on hormonal contraception alone when taking bosentan [see *Use in Specific Populations (8.3)*].

An interaction study demonstrated that co-administration of bosentan and a combination oral hormonal contraceptive produced average decreases of norethindrone and ethinyl estradiol levels of 14% and 31%, respectively. However, decreases in exposure were as much as 56% and 66%, respectively, in individual subjects.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on data from animal reproduction studies, bosentan may cause fetal harm, including birth defects and fetal death, when administered to a pregnant female and is contraindicated during pregnancy [see *Contraindications (4.1)*]. Available data from postmarketing reports and published literature over decades of use with endothelin receptor antagonists (ERA) in the same class as bosentan have not identified an increased risk of major birth defects; however, these data are limited. Methodological limitations of these postmarketing reports and published literature include lack of a control group; limited information regarding dose, duration, and timing of drug exposure; and missing data. These limitations preclude establishing a reliable estimate of the risk of adverse fetal and neonatal outcomes with maternal ERA use. In animal reproduction studies, oral administration of bosentan to pregnant rats at 2- times the maximum recommended human dose (MRHD) on a mg/m² basis caused teratogenic effects in rats, including malformations of the head, mouth, face and large blood vessels [see *Animal Data*]. Advise pregnant women of the potential risk to a fetus.

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.

Data

Animal Data

Bosentan was teratogenic in rats given oral doses two times the MRHD (on a mg/m² basis). In an embryo-fetal toxicity study in rats, bosentan showed dose-dependent teratogenic effects, including malformations of the head, mouth, face and large blood vessels. Bosentan increased stillbirths and pup mortality at oral doses 2 and 10 times the MRHD (on a mg/m² basis). Although birth defects were not observed in rabbits given oral doses of up to the equivalent of 10.5 g/day in a 70 kg person, plasma concentrations of bosentan in rabbits were lower than those reached in the rat. The similarity of malformations induced by bosentan and those observed in endothelin-1 knockout mice and in animals treated with other endothelin receptor antagonists indicates that embryo-fetal toxicity is a class effect of these drugs.

8.2 Lactation

Risk Summary

Data from a case report describe the presence of bosentan in human milk. There is insufficient information about the effects of bosentan on the breastfed infant and no information on the effects of bosentan on milk production. Because of the potential for serious adverse reactions, such as fluid retention and hepatotoxicity, in breastfed infants from bosentan, advise women not to breastfeed during treatment with bosentan.

8.3 Females and Males of Reproductive Potential

Based on data from animal reproductive toxicity studies, bosentan may cause fetal harm, including birth defects and fetal death, when administered to a pregnant patient and is contraindicated during pregnancy [*see Contraindications (4.1), Use in Specific Populations (8.1)*].

Pregnancy Testing

Verify that females of reproductive potential are not pregnant prior to initiating bosentan. The patient should contact her physician immediately for pregnancy testing if onset of menses is delayed or pregnancy is suspected. If the pregnancy test is positive, the physician and patient must discuss the risks to her, the pregnancy and the fetus [*see Dosage and Administration (2.1), Contraindications (4.1), Warnings and Precautions (5.3)*].

Contraception

Drug interaction studies show that bosentan reduces serum levels of the estrogen and progestin in oral contraceptives. Based on these findings, hormonal contraceptives (including oral, injectable, transdermal and implantable contraceptives) may be less effective for preventing pregnancy in patients using bosentan and should not be used as a patient's only contraceptive method [*see Drug Interactions (7.2)*]. Females of

reproductive potential using bosentan should use effective methods of contraception prior to initiation of treatment, during treatment and for 1 month after treatment with bosentan [see *Warnings and Precautions (5.3)*]. Patients may choose one highly effective form of contraception (intrauterine devices (IUD) or tubal sterilization) or a combination of methods (hormone method with a barrier method or two barrier methods). If a partner's vasectomy is the chosen method of contraception, a hormone or barrier method must be used along with this method. Counsel patients on pregnancy planning and prevention, including emergency contraception, or designate counseling by another healthcare provider trained in contraceptive counselling.

Infertility

Males

Decreased sperm counts have been observed in patients receiving bosentan. Based on these findings and findings in animals, bosentan may impair fertility in males of reproductive potential. It is not known whether effects on fertility would be reversible [see *Warnings and Precautions (5.6)*, *Adverse Reactions (6.1)*, *Nonclinical Toxicology (13.1)*].

8.4 Pediatric Use

The efficacy of bosentan in patients < 18 years is supported by data from an uncontrolled trial in which 19 pediatric patients were treated with bosentan. In this study, cardiopulmonary hemodynamic improvements were similar to those seen in adults treated with bosentan [see *Clinical Studies (14.1)*]. Safety in pediatric patients is supported by data from 100 pediatric patients treated with bosentan for a median of 17 months [see *Clinical Studies (6.1)*, *Adverse Reactions (14.1)*].

Juvenile Animal Toxicity Data

In a juvenile rat toxicity study, rats were treated from Day 4 postpartum to adulthood (Day 69 postpartum). Decreased body weights, absolute weights of testes and epididymides and reduced number of sperm in epididymides were observed after weaning. No effect on testis histology or sperm morphology and function was seen. The NOAEL was 4 times (at Day 4 postpartum) and 2 times (Day 69 postpartum) the human therapeutic exposure, respectively.

No effects on general development, sensory, cognitive function and reproductive performance were detected at the highest dose tested in juvenile rats, 7 times the therapeutic exposure in children with PAH.

8.5 Geriatric Use

Clinical studies of bosentan did not include sufficient numbers of subjects aged 65 and older to determine whether they respond differently from younger subjects.

8.6 Hepatic Impairment

Because there is *in vitro* and *in vivo* evidence that the main route of excretion of bosentan is biliary, liver impairment could be expected to increase exposure (C_{max} and AUC) of bosentan. The pharmacokinetics of bosentan have not been evaluated in patients with severe liver impairment (Child-Pugh Class C). In patients with moderate hepatic impairment (Child-Pugh Class B), the systemic exposures to bosentan and its

active metabolite increased significantly. Bosentan should generally be avoided in patients with moderate or severe liver impairment. Pharmacokinetics of bosentan were not altered in patients with mild impairment of hepatic function (Child-Pugh Class A) [see *Dosage and Administration (2.6)*, *Warnings and Precautions (5.1)*, *Clinical Pharmacology (12.3)*].

8.7 Renal Impairment

The effect of renal impairment on the pharmacokinetics of bosentan is small and does not require dosing adjustment [see *Clinical Pharmacology (12.3)*].

10 OVERDOSAGE

Bosentan has been given as a single dose of up to 2,400 mg in normal volunteers, or up to 2,000 mg/day for 2 months in patients, without any major clinical consequences. The most common side effect was headache of mild to moderate intensity. In the cyclosporine A interaction study, in which doses of 500 and 1,000 mg twice daily of bosentan were given concomitantly with cyclosporine A, trough plasma concentrations of bosentan increased 30-fold, resulting in severe headache, nausea and vomiting, but no serious adverse events. Mild decreases in blood pressure and increases in heart rate were observed.

In the postmarketing period, there was one reported overdose of 10,000 mg of bosentan taken by an adolescent male patient. He had symptoms of nausea, vomiting, hypotension, dizziness, sweating and blurred vision. He recovered within 24 hours with blood pressure support.

Bosentan is unlikely to be effectively removed by dialysis due to the high molecular weight and extensive plasma protein binding.

11 DESCRIPTION

Bosentan tablets for oral suspension is an endothelin receptor antagonist that belongs to a class of highly substituted pyrimidine derivatives, with no chiral centers. It is designated chemically as 4-tert-butyl-N-[6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-[2,2']-bipyrimidin-4-yl]-benzenesulfonamide monohydrate and has the following structural formula:

Bosentan has a molecular weight of 569.63 and a molecular formula of $C_{27}H_{29}N_5O_6S \cdot H_2O$. Bosentan is a white to yellowish powder. It is soluble in acetonitrile,

slightly soluble in methanol and practically insoluble in water. In the solid state, bosentan is very stable, is not hygroscopic and is not light sensitive.

Bosentan is available as a 32 mg tablet for oral suspension and contains the following excipients: acesulfame potassium, aspartame, basic butylated methacrylate copolymer, colloidal silicon dioxide, corn starch, croscarmellose sodium, magnesium stearate, mannitol, sodium lauryl sulfate, stearic acid, talcum and tutti frutti flavor. Each dispersible tablet contains 2.08 mg of phenylalanine. Each dispersible tablet contains 33.045 mg of bosentan monohydrate, equivalent to 32 mg anhydrous bosentan.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Bosentan is a specific and competitive antagonist at endothelin receptor types ET_A and ET_B. Bosentan has a slightly higher affinity for ET_A receptors than for ET_B receptors. The clinical impact of dual endothelin blockage is unknown.

Endothelin-1 (ET-1) is a neurohormone, the effects of which are mediated by binding to ET_A and ET_B receptors in the endothelium and vascular smooth muscle. ET-1 concentrations are elevated in plasma and lung tissue of patients with PAH, suggesting a pathogenic role for ET-1 in this disease.

12.3 Pharmacokinetics

General

After oral administration, maximum plasma concentrations of bosentan are attained within 3 hours to 5 hours and the terminal elimination half-life is about 5 hours in healthy adult subjects. The exposure to bosentan after intravenous and oral administration is about twice as high in adult patients with PAH as it is in healthy adult subjects.

In a relative bioavailability study in healthy adults, the peak plasma concentration and area under the plasma concentration-time curve for bosentan are on an average 14% and 11%, respectively, lower following administration of the oral dispersible tablet compared to the film-coated tablet.

Absorption

The absolute bioavailability of bosentan in normal volunteers is about 50% and is unaffected by food.

Bosentan is highly bound (> 98%) to plasma proteins, mainly albumin. Bosentan does not penetrate into erythrocytes. The volume of distribution is about 18 L.

Elimination

Metabolism

Bosentan has three metabolites, one of which is pharmacologically active and may contribute 10% to 20% of the effect of bosentan. Bosentan is an inducer of CYP2C9 and CYP3A and possibly also of CYP2C19. Upon multiple oral dosing, plasma concentrations in healthy adults decrease gradually to 50% to 65% of those seen after single dose administration, probably the effect of auto-induction of the metabolizing liver enzymes. Steady-state is reached within 3 days to 5 days.

Excretion

Bosentan is eliminated by biliary excretion following metabolism in the liver. Less than 3% of an administered oral dose is recovered in urine. Total clearance after a single intravenous dose is about 4 L/h in patients with PAH.

Specific Populations

Pediatrics

The average plasma exposure to bosentan at steady state (AUC_{SS}) in pediatric patients with PAH aged 3 years to 15 years treated with 31.25 mg, 62.5 mg or 125 mg (approximately 2 mg/kg) film-coated tablet twice daily is 37% lower than that observed in adult patients with PAH receiving 125 mg film-coated tablet twice daily. Following administration of 4 mg/kg twice daily doses of dispersible tablet in patients with PAH aged 2 years to 11 years, the average systemic exposure to bosentan at steady state is similar to that observed with 2 mg/kg. The average exposure to bosentan in these pediatric patients was approximately half the exposure in adult patients treated with 125 mg film-coated tablets twice daily.

The exposure to bosentan at 2 mg/kg three times daily dosing of dispersible tablet is similar to that of 2 mg/kg twice daily dosing in patients with PAH aged 3 months to 12 years.

Based on these findings, exposure to bosentan reaches a plateau at lower doses in pediatric patients than in adults and doses higher than 2 mg/kg twice daily do not increase the exposure to bosentan in pediatric patients.

Hepatic Impairment

In vitro and *in vivo* evidence showing extensive hepatic metabolism of bosentan suggests that liver impairment could significantly increase exposure to bosentan. In a study comparing 8 patients with mild liver impairment (Child-Pugh Class A) to 8 controls, the single- and multiple- dose pharmacokinetics of bosentan were not altered in patients with mild hepatic impairment.

In another small (N=8) pharmacokinetic study, the steady-state AUC of bosentan was on average 4.7 times higher and the active metabolite Ro 48-5033 was 12.4 times higher in 5 patients with moderately impaired liver function (Child-Pugh Class B) and PAH associated with portal hypertension than in 3 patients with normal liver function and PAH of other etiologies.

The pharmacokinetics of bosentan have not been evaluated in patients with severe liver impairment (Child-Pugh Class C) [see *Dosage and Administration (2.2)*, *Warnings and Precautions (5.1)*, *Use in Specific Populations (8.6)*].

Renal Impairment

In patients with severe renal impairment (creatinine clearance 15 mL/min to 30 mL/min), plasma concentrations of bosentan were essentially unchanged and plasma concentrations of the three metabolites were increased about 2-fold compared to subjects with normal renal function. These differences do not appear to be clinically important.

Drug Interactions

Ketoconazole

Co-administration of bosentan 125 mg twice daily and ketoconazole, a potent CYP3A inhibitor, increased the plasma concentrations of bosentan by approximately 100% in normal volunteers. No dose adjustment of bosentan is necessary, but increased effects of bosentan should be considered.

Warfarin

Co-administration of bosentan 500 mg twice daily for 6 days in normal volunteers decreased the plasma concentrations of both S-warfarin (a CYP2C9 substrate) and R-warfarin (a CYP3A substrate) by 29% and 38%, respectively. Clinical experience with concomitant administration of bosentan and warfarin in patients with PAH did not show clinically relevant changes in INR or warfarin dose (baseline vs. end of the clinical studies) and the need to change the warfarin dose during the trials due to changes in INR or due to adverse events was similar among bosentan- and placebo-treated patients.

Digoxin, Nimodipine and Losartan

Bosentan has no significant pharmacokinetic interactions with digoxin and nimodipine and losartan has no significant effect on plasma levels of bosentan.

Sildenafil

In normal volunteers, co-administration of multiple doses of 125 mg twice daily bosentan and 80 mg three times daily sildenafil resulted in a reduction of sildenafil plasma concentrations by 63% and increased bosentan plasma concentrations by 50%. The changes in plasma concentrations were not considered clinically relevant and dose adjustments are not necessary. This recommendation holds true when sildenafil is used for the treatment of PAH or erectile dysfunction.

Tadalafil

Bosentan (125 mg twice daily) reduced tadalafil (40 mg once per day) systemic exposure (AUC) by 42% and C_{max} by 27% following multiple dose co-administration. Tadalafil did not affect the exposure (AUC and C_{max}) of bosentan or its metabolites.

Figure 3

CYP Induction-mediated effect of bosentan on other drugs

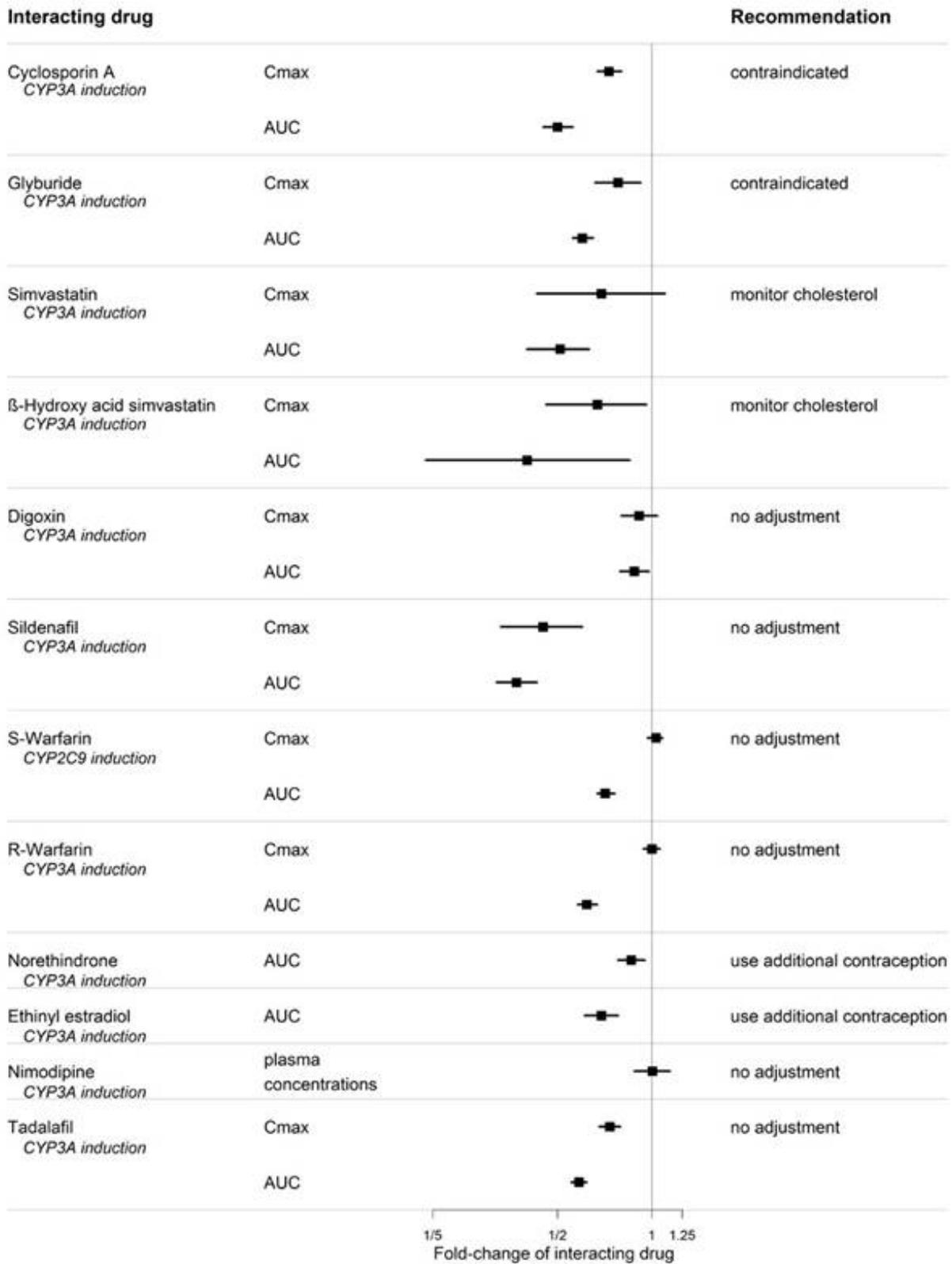
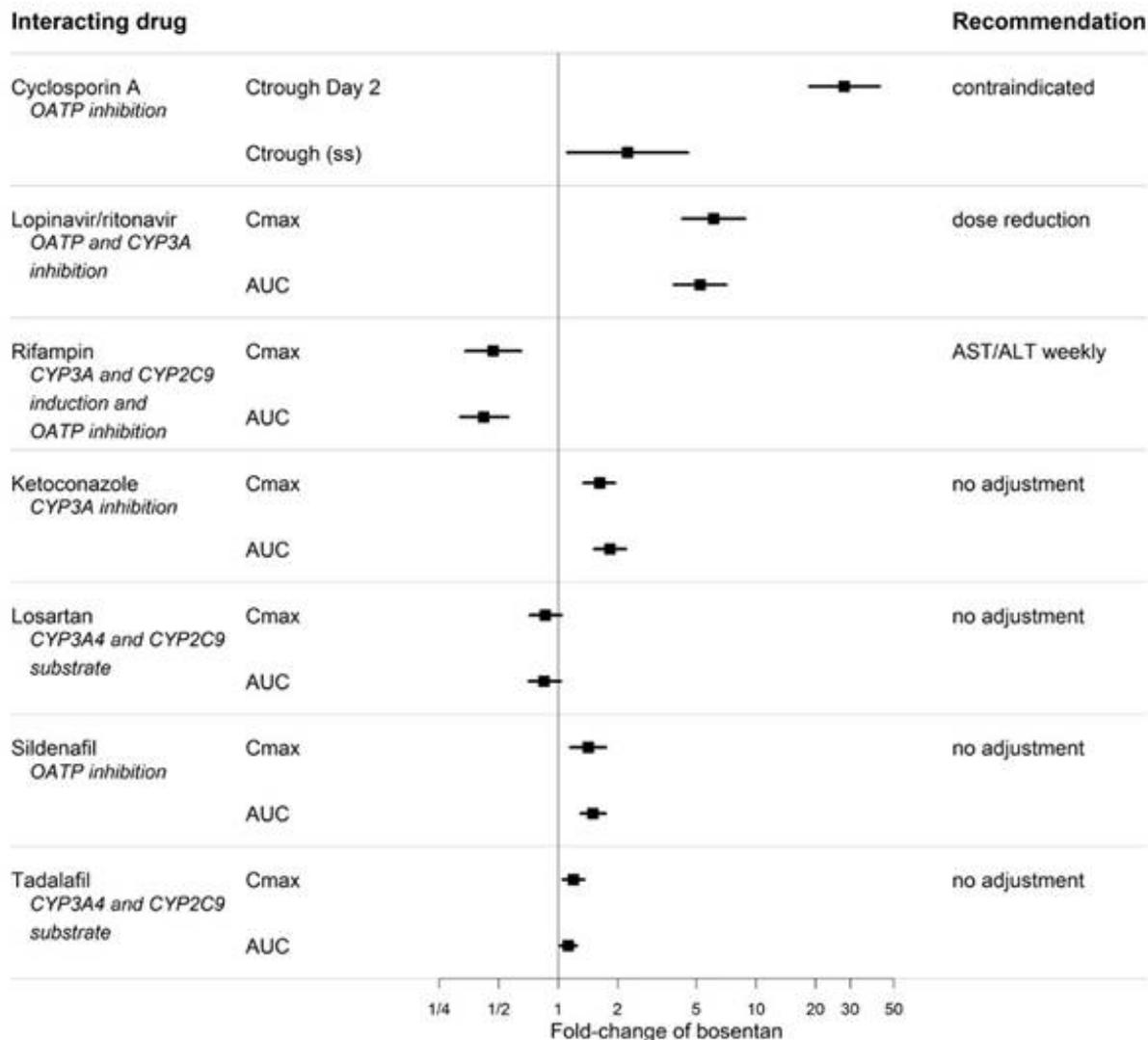


Figure 4
Effects of other drugs on bosentan



13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis and Mutagenesis

Two years of dietary administration of bosentan to mice produced an increased incidence of hepatocellular adenomas and carcinomas in males at doses as low as 450 mg/kg/day (about 8 times the maximum recommended human dose [MRHD] of 125 mg twice daily, on a mg/m² basis). In the same study, doses greater than 2,000 mg/kg/day (about 32 times the MRHD) were associated with an increased incidence of colon adenomas in both males and females. In rats, dietary administration of bosentan for two years was associated with an increased incidence of brain astrocytomas in males at doses as low as 500 mg/kg/day (about 16 times the MRHD). In a comprehensive battery of *in vitro* tests (the microbial mutagenesis assay, the unscheduled DNA synthesis assay, the V-79 mammalian cell mutagenesis assay and human lymphocyte assay) and an *in vivo* mouse micronucleus assay, there was no evidence for any mutagenic or clastogenic activity of bosentan.

Impairment of Fertility/Testicular Function

The development of testicular tubular atrophy and impaired fertility has been linked with the chronic administration of certain endothelin receptor antagonists in rodents.

Treatment with bosentan at oral doses of up to 1,500 mg/kg/day (50 times the MRHD on a mg/m² basis) or intravenous doses up to 40 mg/kg/day had no effects on sperm count, sperm motility, mating performance or fertility in male and female rats. An increased incidence of testicular tubular atrophy was observed in rats given bosentan orally at doses as low as 125 mg/kg/day (about 4 times the MRHD and the lowest doses tested) for two years but not at doses as high as 1,500 mg/kg/day (about 50 times the MRHD) for 6 months. Effects on sperm count and motility were evaluated only in the much shorter duration fertility studies in which males had been exposed to the drug for 4 weeks to 6 weeks. An increased incidence of tubular atrophy was not observed in mice treated for 2 years at doses up to 4,500 mg/kg/day (about 75 times the MRHD) or in dogs treated up to 12 months at doses up to 500 mg/kg/day (about 50 times the MRHD).

14 CLINICAL STUDIES

14.1 Pulmonary Arterial Hypertension

WHO Functional Class III-IV

Two randomized, double-blind, multi-center, placebo-controlled trials were conducted in 32 and 213 patients. The larger study (BREATHE-1) compared 2 doses (125 mg twice daily and 250 mg twice daily) of bosentan with placebo. The smaller study (Study 351) compared 125 mg twice daily with placebo. Patients had severe (WHO functional Class III-IV) PAH: idiopathic or heritable PAH (72%) or PAH associated with scleroderma or other connective tissue diseases (21%), or to autoimmune diseases (7%). There were no patients with PAH associated with other conditions such as HIV disease or recurrent pulmonary emboli.

In both studies, bosentan or placebo was added to patients' current therapy, which could have included a combination of digoxin, anticoagulants, diuretics and vasodilators (e.g., calcium channel blockers, ACE inhibitors), but not epoprostenol. Bosentan was given at a dose of 62.5 mg twice daily for 4 weeks and then at 125 mg twice daily or 250 mg twice daily for either 12 (BREATHE-1) or 8 (Study 351) additional weeks. The primary study endpoint was 6-minute walk distance. In addition, symptoms and functional status were assessed. Hemodynamic measurements were made at 12 weeks in Study 351.

The mean age was about 49 years. About 80% of patients were female and about 80% were Caucasian. Patients had been diagnosed with pulmonary hypertension for a mean of 2.4 years.

Submaximal Exercise Ability

Results of the 6-minute walk distance at 3 months (Study 351) or 4 months (BREATHE-1) are shown in Table 4.

Table 4 Effects of Bosentan on 6-minute walk distance

	BREATHE-1		Study 351	
	Bosentan	Bosentan	Bosentan	

	125 mg twice daily (n=74)	250 mg twice daily (n=70)	Placebo (n=69)	125 mg twice daily (n=21)	Placebo (n=11)
Baseline	326 ± 73	333 ± 75	344 ± 76	360 ± 86	355 ± 82
End point	353 ± 115	379 ± 101	336 ± 129	431 ± 66	350 ± 147
Change from baseline	27 ± 75	46 ± 62	-8 ± 96	70 ± 56	-6 ± 121
Placebo-subtracted	35 ^(a)	54 ^(b)		76 ^(c)	

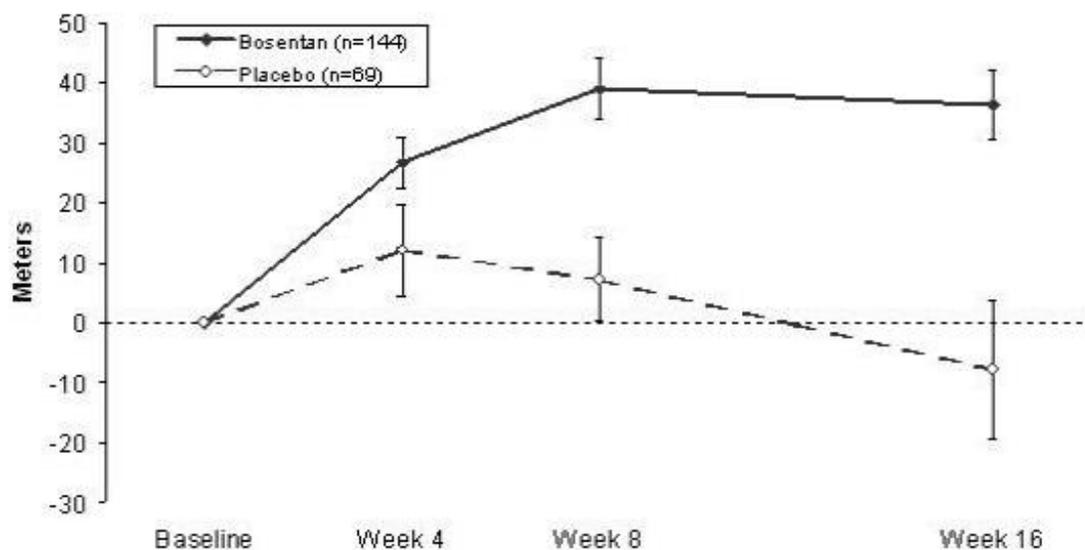
Distance in meters; mean ± standard deviation. Changes are to week 16 for BREATHE-1 and to week 12 for Study 351.

(a) $p=0.01$; by Wilcoxon; (b) $p=0.0001$; by Wilcoxon; (c) $p=0.02$; by Student's t-test.

In both trials, treatment with bosentan resulted in a significant increase in exercise ability. The improvement in walk distance was apparent after 1 month of treatment (with 62.5 mg twice daily) and fully developed by about 2 months of treatment (Figure 5). It was maintained for up to 7 months of double-blind treatment. Walking distance was somewhat greater with 250 mg twice daily, but the potential for increased hepatotoxicity causes this dose not to be recommended [see *Dosage and Administration (2.2)*]. There were no apparent differences in treatment effects on walk distance among subgroups analyzed by demographic factors, baseline disease severity, or disease etiology, but the studies had little power to detect such differences.

Figure 5

Mean Change in 6-min Walk Distance (BREATHE-1)



Change from baseline in 6-minute walking distance from start of therapy to week 16 in the placebo and combined bosentan (125 mg twice daily and 250 mg twice daily) groups. Values are expressed as mean ± standard error of the mean.

Hemodynamic Changes

Invasive hemodynamic parameters were assessed in Study 351. Treatment with

bosentan led to a significant increase in cardiac index (CI) associated with a significant reduction in pulmonary artery pressure (PAP), pulmonary vascular resistance (PVR) and mean right atrial pressure (RAP) (Table 5).

The relationship between hemodynamic effects and improvements in 6-minute walk distance is unknown.

Table 5 Change from Baseline to Week 12: Hemodynamic Parameters

	Bosentan 125 mg twice daily		Placebo
CI (L/min/m²)	n=20		n=10
Baseline	2.35±0.73		2.48±1.03
Absolute Change	0.50±0.46		-0.52±0.48
Treatment Effect			1.02 ^(a)
Mean PAP (mmHg)	n=20		n=10
Baseline	53.7±13.4		55.7±10.5
Absolute Change	-1.6±5.1		5.1±8.8
Treatment Effect			-6.7 ^(b)
PVR (dyn•sec•cm⁻⁵)	n=19		n=10
Baseline	896±425		942±430
Absolute Change	-223±245		191±235
Treatment Effect			-415 ^(a)
Mean RAP (mmHg)	n=19		n=10
Baseline	9.7±5.6		9.9±4.1
Absolute Change	-1.3±4.1		4.9±4.6
Treatment Effect			-6.2 ^(a)

Values shown are means ± SD

(a) $p \leq 0.001$; (b) $p < 0.02$

Symptoms and Functional Status

Symptoms of PAH were assessed by Borg dyspnea score, WHO functional class and rate of "clinical worsening." Clinical worsening was assessed as the sum of death, hospitalizations for PAH, discontinuation of therapy because of PAH and need for epoprostenol. There was a significant reduction in dyspnea during walk tests (Borg dyspnea score) and significant improvement in WHO functional class in bosentan-treated patients. There was a significant reduction in the rate of clinical worsening (Table 6 and Figure 6). Figure 6 shows the log-rank test reflecting clinical worsening over 28 weeks.

Table 6 Incidence of Clinical Worsening, Intent To Treat Population

	BREATHE-1		Study 351	
	Bosentan 125/250 mg twice daily	Placebo (n=69)	Bosentan 125 mg twice daily	Placebo (n=11)

	(n=144)		(n=21)	
Patients with clinical worsening [n (%)]	9 (6%) ^(a)	14 (20%)	0 (0%) ^(b)	3 (27%)
Death	1 (1%)	2 (3%)	0 (0%)	0 (0%)
Hospitalization for PAH	6 (4%)	9 (13%)	0 (0%)	3 (27%)
Discontinuation due to worsening of PAH	5 (3%)	6 (9%)	0 (0%)	3 (27%)
Receipt of epoprostenol ^(c)	4 (3%)	3 (4%)	0 (0%)	3 (27%)

Note: Patients may have had more than one reason for clinical worsening.

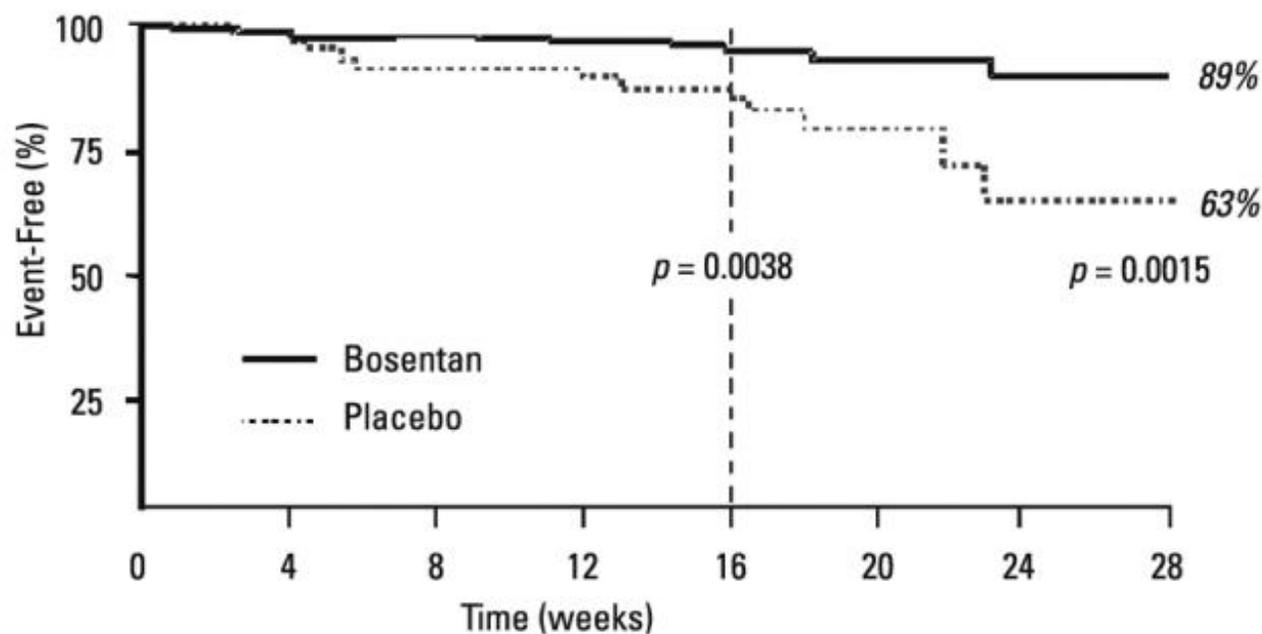
(a) $p=0.0015$ vs. placebo by log-rank test. There was no relevant difference between the 125 mg and 250 mg twice daily groups.

(b) $p=0.033$ vs. placebo by Fisher's exact test.

(c) Receipt of epoprostenol was always a consequence of clinical worsening.

Figure 6

Time to Clinical Worsening (BREATHE-1)



Time from randomization to clinical worsening with Kaplan-Meier estimate of the proportions of failures in BREATHE-1. All patients (n=144 in the bosentan group and n=69 in the placebo group) participated in the first 16 weeks of the study. A subset of this population (n=35 in the bosentan group and 13 in the placebo group) continued double-blind therapy for up to 28 weeks.

WHO Functional Class II

In a randomized, double-blind, multicenter, placebo-controlled trial, 185 mildly symptomatic PAH patients with WHO Functional Class II (mean baseline 6-minute walk distance of 443 meters) received bosentan 62.5 mg twice daily for 4 weeks followed by 125 mg twice daily (n=93), or placebo (n=92) for 6 months. Enrolled patients were treatment-naïve (n=156) or on a stable dose of sildenafil (n=29). The coprimary endpoints were change from baseline to month 6 in PVR and 6-minute walk distance.

Time to clinical worsening (assessed as the sum of death, hospitalization due to PAH complications, or symptomatic progression of PAH), Borg dyspnea index, change in WHO functional class and hemodynamics were assessed as secondary endpoints.

Compared with placebo, bosentan treatment was associated with a reduced incidence of worsening of at least one functional class (3% bosentan vs. 13% placebo, $p = 0.03$) and improvement in hemodynamic variables (PVR, mPAP, TPR, cardiac index and SVO₂; $p < 0.05$). The +19 m mean (+14 m median) increase in 6-minute walk distance with bosentan vs. placebo was not significant ($p = 0.08$). There was a significant delay in time to clinical worsening (first seen primarily as symptomatic progression of PAH) with bosentan compared with placebo (hazard ratio 0.2, $p = 0.01$). Findings were consistent in strata with or without treatment with sildenafil at baseline.

Long-term Treatment of PAH

Long-term follow-up of patients with Class III and IV PAH who were treated with bosentan in open-label extensions of trials (N=235) showed that 93% and 84% of patients were still alive at 1 year and 2 years, respectively, after the start of treatment.

These uncontrolled observations do not allow comparison with a group not given bosentan and cannot be used to determine the long-term effect of bosentan on mortality.

Pediatric Studies

The efficacy of bosentan was evaluated in an open-label, uncontrolled study in 19 pediatric patients with PAH aged 3 years to 15 years. Patients had primary pulmonary hypertension (n=10) or PAH related to congenital heart diseases (9 patients) and were WHO functional class II (n=15, 79%) or class III (n=4; 21%) at baseline. Patients were dosed with bosentan at approximately 2 mg/kg twice daily (body weight adjusted dose, corresponding to the recommended adult dose) [see *Dosage and Administration (2.1)*] for 12 weeks. Half of the patients in each group were already being treated with intravenous epoprostenol and the dose of epoprostenol remained constant for the duration of the study.

Hemodynamics were measured in 17 patients (Table 7). The mean decrease in PVR was 389 dyn·sec·cm⁻⁵, which was similar to the effect seen in adults. Hemodynamic improvements from baseline were similar with or without co-administration of epoprostenol.

Table 7 Change from Baseline to Week 12: Hemodynamic Parameters

	Bosentan 2 mg/kg twice daily
CI (L/min/m²)	n=17
Baseline	4±1.5
Absolute Change	0.5±1.4
Mean PAP (mmHg)	n=18
Baseline	60±18
Absolute Change	-8±9
PVR (dyn•sec•cm⁻⁵)	n=17
Baseline	1,195±755

Absolute Change	-389±616
Mean RAP (mmHg)	n=18
Baseline	6.1±2.7
Absolute Change	-0.5±2.3

Values shown are means ± SD

Pulmonary Arterial Hypertension in Adults related to Congenital Heart Disease with Left-to-Right Shunts

A small study (N=54) and its open-label extension (N=37) of up to 40 weeks in adult patients with Eisenmenger physiology demonstrated effects of bosentan on exercise and safety that were similar to those seen in other trials in patients with PAH (WHO Group 1).

14.2 Lack of Benefit in Congestive Heart Failure

Bosentan is not effective in the treatment of congestive heart failure with left ventricular dysfunction. In a pair of studies, 1,613 subjects with NYHA Class III-IV heart failure, left ventricular ejection fraction < 35%, on diuretics, ACE inhibitor and other therapies, were randomized to placebo or bosentan (62.5 mg twice daily titrated as tolerated to 125 mg twice daily) and followed for up to 70 weeks. Use of bosentan was associated with no benefit on patient global assessment (the primary end point) or mortality. However, hospitalizations for heart failure were more common during the first 4 weeks to 8 weeks after bosentan was initiated. In a placebo-controlled trial of patients with severe chronic heart failure, there was an increased incidence of hospitalization for CHF associated with weight gain and increased leg edema during the first 4 weeks to 8 weeks of treatment with bosentan. Patients required intervention with a diuretic, fluid management, or hospitalization for decompensating heart failure.

16 HOW SUPPLIED/STORAGE AND HANDLING

Bosentan tablets for oral suspension, 32 mg are off-white to pale yellow, round, flat, beveled edge tablet debossed with '1508' on one side and bisect on the other and are supplied as follows:

NDC 70710-1508-8 in unit-dose blister cartons of 56 tablets (4 x 14 unit-dose)

This package is child-resistant.

Store at 20°C to 25°C (68°F to 77°F). Excursions are permitted between 15°C and 30°C (59°F and 86°F). [See USP Controlled Room Temperature]. Divided dispersible tablets should be stored under the same conditions and used within 7 days. Tablet pieces may be returned to the opened blister and stored there out of reach of children for up to 7 days.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Medication Guide)

Restricted Access

Advise the patient that bosentan is only available through a restricted access program called the Bosentan REMS and that bosentan is available only from certified pharmacies enrolled in the Bosentan REMS.

As a component of the Bosentan REMS, prescribers must review the contents of the bosentan Medication Guide with the patient before initiating bosentan and patients must sign the Patient Enrollment Form to confirm that they understand the risks of bosentan.

Instruct patients that the risks associated with bosentan include hepatotoxicity. Discuss with the patient the requirement to measure serum aminotransferases monthly [see *Dosage and Administration (2.1, 2.4), Warnings and Precautions (5.1, 5.2)*].

Embryo-Fetal toxicity

Counsel female patients of reproductive potential about the need to use effective contraception prior to treatment, during treatment with bosentan and for one month after treatment discontinuation. Females of reproductive potential should have a negative pregnancy test prior to treatment [see *Dosage and Administration (2.1), Warnings and Precautions (5.3), Drug Interactions (7.2), Use in Specific Populations (8.1, 8.3)*].

Patients should be instructed to immediately contact their physician if they suspect they may be pregnant. Patients should seek additional contraceptive advice from a gynecologist or similar expert as needed.

Educate and counsel females of reproductive potential on the use of emergency contraception in the event of unprotected sex or contraceptive failure.

Advise pre-pubertal females to report any changes in their reproductive status immediately to her prescriber.

Lactation

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

Infertility

Advise males of reproductive potential that bosentan may impair fertility [see *Warnings and Precautions (5.6), Adverse Reactions (6.1), Use in Specific Populations (8.3) and Nonclinical Toxicology (13.1)*].

Phenylketonurics

Bosentan dispersible tablets contain phenylalanine, a component of aspartame. Each dispersible tablet contains 2.08 mg of phenylalanine.

Other Risks Associated with Bosentan

Instruct patients that the risks associated with bosentan also include the following:

- Decreases in hemoglobin and hematocrit - advise patients of the importance of hemoglobin testing
- Decreases in sperm count
- Fluid retention

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Zydus Lifesciences Ltd.

Ahmedabad-382213, India

Distributed by:

Zydus Pharmaceuticals (USA) Inc.

Pennington, NJ 08534

Rev.: 11/25

Medication Guide

Bosentan (boe sen' tan) Tablets for Oral Suspension

Read the Medication Guide that comes with bosentan tablets for oral suspension before you start taking it and each time you get a refill. There may be new information. This Medication Guide does not take the place of talking with your healthcare provider about your medical condition or your treatment.

What is the most important information I should know about bosentan tablets for oral suspension?

Bosentan tablets for oral suspension can cause serious side effects including:

Liver damage.

- Liver damage may not cause symptoms at first. Only a blood test can show if you have early liver damage. You must have your blood tested to check your liver function before you start bosentan tablets for oral suspension and each month after that. Your healthcare provider will order these tests. Regular blood tests are important because they will help your healthcare provider adjust or stop your treatment before there is permanent damage.
- Tell your healthcare provider if you have had liver problems, including liver problems while taking other medicines. Call your healthcare provider right away if you have any of these symptoms of liver problems while taking bosentan tablets for oral suspension:
 - nausea
 - vomiting
 - fever
 - unusual tiredness
 - stomach area (abdominal) pain
 - yellowing of the skin or the whites of your eyes (jaundice)

Because of the risk for liver damage, bosentan is only available through a restricted program called the Bosentan REMS. Before you begin taking bosentan, you must read and agree to all of the requirements of the Bosentan REMS.

Serious birth defects.

- Bosentan tablets for oral suspension can cause serious birth defects if taken during pregnancy. You should not be pregnant when you start taking bosentan tablets for oral suspension or during bosentan treatment. Serious birth defects from bosentan tablets for oral suspension may happen early in pregnancy. Females who are able to

get pregnant should have a negative pregnancy test before starting treatment with bosentan tablets for oral suspension.

- Talk to your healthcare provider about your menstrual cycle. Your healthcare provider will decide when to do a pregnancy test and will order a pregnancy test for you depending on your menstrual cycle.
- Females who **are able** to get pregnant are females who:
 - have entered puberty, even if they have not started their menstrual period **and**
 - have a uterus **and**
 - have not gone through menopause. Menopause means that you have not had a menstrual period for at least 12 months for natural reasons, or that you have had your ovaries removed.
- Females who **are not able** to get pregnant are females who:
 - have not yet entered puberty, **or**
 - do not have a uterus, **or**
 - have gone through menopause. Menopause means that you have not had a menstrual period for at least 12 months for natural reasons, or that you have had your ovaries removed **or**
 - are infertile for other medical reasons and this infertility is permanent and cannot be reversed.
- **Females who are able to get pregnant should use effective birth control prior to starting treatment with bosentan tablets for oral suspension, during treatment with bosentan tablets for oral suspension and for one month after stopping bosentan tablets for oral suspension because the medicine may still be in the body.**
- Talk with your healthcare provider or gynecologist (a doctor who specializes in female reproduction) to find out about options for effective birth control that you may use to prevent pregnancy during treatment with bosentan tablets for oral suspension.
- If you decide that you want to change the form of birth control that you use, talk with your healthcare provider or gynecologist to be sure that you choose another acceptable form of birth control.
- **Do not have unprotected sex. Talk to your healthcare provider or pharmacist right away if you have unprotected sex or if you think your birth control has failed. Your healthcare provider may talk with you about using emergency birth control.**
- **Tell your healthcare provider right away if you miss a menstrual period or think you may be pregnant.**

If you are the parent or caregiver of a female child who started taking bosentan tablets for oral suspension before reaching puberty, you should check your child regularly to see if she is developing signs of puberty. Tell your healthcare provider right away if you notice that she is developing

breast buds or any pubic hair. Your healthcare provider should decide if your child has reached puberty. **Your child may reach puberty before having her first menstrual period.**

Bosentan tablets for oral suspension 32 mg dispersible tablets contain Aspartame. Phenylketonurics: Contains Phenylalanine 2.08 mg per 32 mg dispersible tablet.

See " **What are the possible side effects of bosentan tablets for oral suspension?** " for more information about side effects.

What are bosentan tablets for oral suspension?

Bosentan tablets for oral suspension are a prescription medicine used to treat people with certain types of pulmonary arterial hypertension (PAH), which is high blood pressure in the vessels of the lungs.

Bosentan tablets for oral suspension can improve your ability to exercise and can slow the worsening of your physical condition and symptoms.

Bosentan tablets for oral suspension lowers high blood pressure in your lungs and lets your heart pump blood more efficiently.

Bosentan tablets for oral suspension is only:

Prescribed by healthcare providers who are enrolled in the Bosentan REMS. Available to people who understand the risks and requirements and agree to enroll in the Bosentan REMS.

Who should not take bosentan tablets for oral suspension?

Do not take bosentan tablets for oral suspension if you:

- **are pregnant, plan to become pregnant, or become pregnant during bosentan tablets for oral suspension treatment. Bosentan tablets for oral suspension can cause serious birth defects.** All females should read the **birth defects** section of " **What is the most important information I should know about bosentan tablets for oral suspension?** "
- take any of these medicines:
 - cyclosporine A used to treat psoriasis and rheumatoid arthritis and to prevent rejection of heart, liver and kidney transplants
 - glyburide used to treat diabetes
- are allergic to bosentan or any of the ingredients in bosentan tablets for oral suspension. See the end of this Medication Guide for a complete list of the ingredients in bosentan tablets for oral suspension. If you have a rash, hives or your lips swell after taking bosentan tablets for oral suspension, it may be a sign of allergy. You should stop taking your bosentan tablets for oral suspension and talk to your healthcare provider.

What should I tell my healthcare provider before taking bosentan tablets for oral suspension?

Bosentan tablets for oral suspension may not be right for you. **Tell your healthcare provider about all your medical conditions, including if you:**

- **have liver problems.**
- **are pregnant or plan to become pregnant. See "What is the**

most important information I should know about bosentan tablets for oral suspension?"

- **are breast-feeding or plan to breast feed.** Bosentan passes into your breast milk. It is not known if bosentan in breast milk can harm your baby. You and your healthcare provider should decide if you will take bosentan tablets for oral suspension or breastfeed.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins and herbal supplements. Bosentan tablets for oral suspension and other medicines may affect how each other works and cause side effects. Especially tell your healthcare provider if you take:

- hormone-based birth control, such as pills, shots, patches and implants. These birth control methods may not work as well when taken with bosentan tablets for oral suspension.
- simvastatin or other "-statin" medicines used to lower cholesterol
- rifampin used for tuberculosis
- ketoconazole, fluconazole, itraconazole, or voriconazole used for fungal infections
- warfarin sodium used to prevent blood clots
- ritonavir used to treat HIV

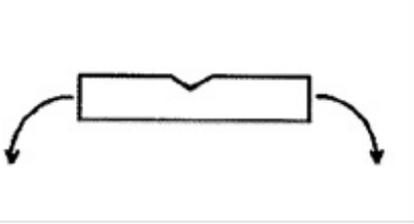
There may be more than one brand name medicine. Ask your healthcare provider if you are not sure if your medicine is one that is listed above. Know the medicines you take. Keep a list of them and show it to your healthcare provider or pharmacist when you get a new medicine.

How should I take bosentan tablets for oral suspension?

Your healthcare provider will give you detailed information about the Bosentan REMS Program.

- Bosentan tablets for oral suspension are only available through certified pharmacies. You will only receive a 30-day supply of bosentan tablets for oral suspension at one time.
- Take bosentan tablets for oral suspension exactly as prescribed.
- Your healthcare provider will tell you how much bosentan tablets for oral suspension to take and when to take it.
- In most cases, you will take 1 tablet in the morning and 1 in the evening.
- You can take bosentan tablets for oral suspension orally with or without food.
- If you take more than the prescribed dose of bosentan tablets for oral suspension, call your healthcare provider right away.
- If you miss a dose of bosentan tablets for oral suspension, take your tablet as soon as you remember. Do not take 2 doses at the same time. If it is almost time for your next dose, skip the missed dose. Just take the next dose at your regular time.
- Do not stop taking bosentan tablets for oral suspension unless your healthcare provider tells you to. Suddenly stopping your treatment may cause your symptoms to get worse. If you need to stop taking bosentan tablets for oral suspension, speak with your healthcare provider about the right way to stop.

- If necessary, the 32 mg dispersible tablet can be divided into halves by breaking along a line cut into the surface. Hold the tablet between the thumb and index finger on either side of the line, with the line facing upwards and break the tablet along the line (see figure below).



- Each 32 mg dispersible tablet, or tablet part, can be dispersed in a minimal amount of water to make a liquid medicine immediately before administration. When the tablet has fully dispersed, the liquid should be administered to the patient.

What are the possible side effects of bosentan tablets for oral suspension?

Bosentan tablets for oral suspension can cause serious side effects, including:

- **See "What is the most important information I should know about bosentan tablets for oral suspension?"**
- **Fluid retention and swelling of your ankles and legs. Bosentan tablets for oral suspension can cause your body to hold too much water and you may get swelling of your ankles and legs. Tell your healthcare provider if you have swelling of your ankles and legs that happens either with or without weight gain, or if you have more trouble with your breathing than normal. Your healthcare provider will look for the cause of this.**
- **Lower Sperm Count. Some men who take bosentan tablets for oral suspension may have lower sperm counts. This may affect your ability to father a child. Tell your healthcare provider if fertility is a concern for you.**
- **Low red blood cell levels (anemia). Your healthcare provider will do blood tests to check your red blood cells during treatment with bosentan tablets for oral suspension.**

The most common side effects of bosentan tablets for oral suspension include:

- respiratory tract infection
- headache
- fainting
- flushing
- low blood pressure
- inflamed nose passages (sinusitis)
- joint pain
- irregular heart beats

Tell your doctor if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of bosentan tablets for oral suspension. For more information, ask your doctor or pharmacist. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store bosentan tablets for oral suspension?

- Store bosentan tablets for oral suspension at room temperature between 68°F to 77°F (20°C to 25°C).
- Dispersible tablets that have been broken to adjust the dose of medicine can be stored at room temperature, between 68°F to 77°F (20°C to 25°C) and should be used within 7 days of having been broken. Tablet pieces may be returned to the opened blister and stored there out of reach of children for up to 7 days.
- Drug bosentan tablets for oral suspension come in child-resistant unit-dose blister pack of 56 tablets.

Keep bosentan tablets for oral suspension and all medicines out of the reach of children.

General information about bosentan tablets for oral suspension

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use bosentan tablets for oral suspension for a condition for which it was not prescribed. Do not give bosentan tablets for oral suspension to other people, even if they have the same symptoms that you have. It may harm them.

This Medication Guide summarizes the most important information about bosentan tablets for oral suspension. If you would like more information, talk with your healthcare provider. You can ask your pharmacist or healthcare provider for information about bosentan tablets for oral suspension that is written for health professionals. For more information, go to MedicalAffairs@zydususa.com or Tel.: 1-877-993-8779.

What are the ingredients in bosentan tablets for oral suspension?

Active ingredient: bosentan

Inactive ingredients: acesulfame potassium, aspartame, basic butylated methacrylate copolymer, colloidal silicon dioxide, corn starch, croscarmellose sodium, magnesium stearate, mannitol, sodium lauryl sulfate, stearic acid, talcum and tutti frutti flavor.

Manufactured by:
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Ahmedabad-382213, India

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This Medication Guide has been approved by the U.S. Food and Drug Administration.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70710-1508-8 in unit-dose blister cartons of 56 tablets (4 x 14 unit-dose)

Bosentan tablets for oral suspension, 32 mg

Rx only

Zydus

**START
HERE**



**Bosentan Tablet for
Oral Suspension 32 mg**

Distributed by:
Zydus Pharmaceuticals (USA) Inc.,
Pennington, NJ 08534

Lot: XXXXXXXX
Exp: 00/0000

Rev.: 06/22



**Bosentan Tablet for
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Exp: 00/0000

Rev.: 06/22



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Exp: 00/0000

Rev.: 06/22

PEEL

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Rev.: 06/22



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Rev.: 06/22

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Exp: 00/0000

Rev.: 06/22

Phenylketonurics:
Contains Phenylalanine
2.08 mg per tablet
for oral suspension.

Made in INDIA

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Bosentan Tablet for Oral Suspension 32 mg

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Bosentan Tablet for Oral Suspension 32 mg

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Exp: 00/0000

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Lot: XXXXXXXX
Exp: 00/0000

Rev: 0622

NDC 70710-1508-8

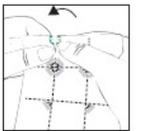
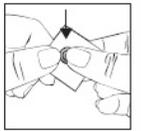
Each tablet contains 32 mg of anhydrous bosentan.

Usual Dosage:
Disperse tablets for oral suspension, or tablet half, in a minimal amount of water immediately before administration.
See enclosed full prescribing information.

Important: Read accompanying prescribing information carefully.
Store at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature].
Divided tablets for oral suspension should be stored at 20°C to 25°C (68°F to 77°F) and used within 7 days. Tablet pieces may be returned to the opened blister and stored there for up to 7 days.

This package is child-resistant.
Keep this and all drugs out of the reach of children.

Directions for opening:

		
Step 1 : To separate the cavity, initiate tearing from the "Start here" arrow along the perforated lines.	Step 2 : Peel the printed paper backing off from arrow to expose foil.	Step 3 : Gently Push the tablet through foil.

Manufactured by:
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Ahmedabad-382213, India

Distributed by:
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Rev: 0624

Bosentan Tablets for Oral Suspension 32 mg

Bosentan Tablets for Oral Suspension

32 mg

PHARMACIST: Dispense the accompanying Medication Guide to each patient.

Phenylketonurics:
Contains Phenylalanine
2.08 mg per tablet for oral suspension.

56 Tablets
(4 x 14 unit-dose)
Rx only

NDC 70710-1508-8

Bosentan Tablets for Oral Suspension 32 mg

BOSENTAN

bosentan tablet, for suspension

Product Information

Product Type

HUMAN PRESCRIPTION DRUG

Item Code (Source)

NDC:70710-1508

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOSENTAN (UNII: Q326023R30) (BOSENTAN ANHYDROUS - UNII:XUL93R30K2)	BOSENTAN ANHYDROUS	32 mg

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ASPARTAME (UNII: Z0H242BBR1)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIMETHYLAMINOETHYL METHACRYLATE - BUTYL METHACRYLATE - METHYL METHACRYLATE COPOLYMER (UNII: 905HNO1SIH)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	WHITE (off-white to pale yellow)	Score	2 pieces
Shape	ROUND	Size	10mm
Flavor	TUTTI FRUTTI	Imprint Code	1508
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70710-1508-8	4 in 1 CARTON	02/17/2026	
1	NDC:70710-1508-7	14 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213981	02/17/2026	

Labeler - Zydus Pharmaceuticals USA Inc. (156861945)

Registrant - Zydus Pharmaceuticals USA Inc. (156861945)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		863362789	ANALYSIS(70710-1508) , MANUFACTURE(70710-1508)

Revised: 11/2025

Zydus Pharmaceuticals USA Inc.