

## HIGHLIGHTS OF PRESCRIBING INFORMATION

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

VFEND® (voriconazole) tablets, for oral use

VFEND® (voriconazole) for oral suspension

VFEND® (voriconazole) for injection, for intravenous use

Initial U.S. Approval: 2002

### INDICATIONS AND USAGE

VFEND is an azole antifungal indicated for use in the treatment of:

- Invasive aspergillosis (1.1)
- Candidemia (nonneutropenics) and disseminated candidiasis in skin, abdomen, kidney, bladder wall, and wounds (1.2)
- Esophageal candidiasis (1.3)
- Serious infections caused by *Scedosporium apiospermum* and *Fusarium* species including *Fusarium solani*, in patients intolerant of, or refractory to, other therapy (1.4)

### DOSAGE AND ADMINISTRATION

Recommended Dosage (2.3)			
Infection	Loading dose	Maintenance Dose	
		IV	Oral
Invasive Aspergillosis	6 mg/kg q12h for the first 24 hours	4 mg/kg q12h	200 mg q12h
		3-4 mg/kg q12h	200 mg q12h
Candidemia in nonneutropenics and other deep tissue <i>Candida</i> infections		4 mg/kg q12h	200 mg q12h
Scedosporiosis and Fusariosis		not evaluated	200 mg q12h
Esophageal Candidiasis	Not Evaluated	not evaluated	200 mg q12h

- Adult patients weighing less than 40 kg: oral maintenance dose 100 or 150 mg q12 hours
- See full prescribing information for instructions on reconstitution of lyophilized powder for intravenous use and reconstitution of oral suspension and important administration instructions (2.5, 2.6)

### DOSAGE FORMS AND STRENGTHS

- *Tablets*: 50 mg, 200 mg (3)
- *For Oral Suspension*: 45 grams of powder; after reconstitution 40 mg/mL (3)
- *For Injection*: lyophilized powder containing 200 mg voriconazole and 3200 mg of sulfobutyl ether beta-cyclodextrin sodium (SBECD); after reconstitution 10 mg/mL of voriconazole and 160 mg/mL of SBECD (3)

### CONTRAINDICATIONS

- Hypersensitivity to voriconazole or its excipients (4)
- Coadministration with terfenadine, astemizole, cisapride, pimozide or quinidine, sirolimus due to risk of serious adverse reactions (4, 7)
- Coadministration with rifampin, carbamazepine, long-acting barbiturates, efavirenz, ritonavir, rifabutin, ergot alkaloids, and St. John's Wort due to risk of loss of efficacy (4, 7)

### WARNINGS AND PRECAUTIONS

- *Clinically Significant Drug Interactions*: Review patient's concomitant medications (5.1, 7)

- *Hepatic Toxicity*: Serious hepatic reactions reported. Evaluate liver function tests at start of and during voriconazole therapy (5.2)
- *Visual Disturbances* (including optic neuritis and papilledema): Monitor visual function if treatment continues beyond 28 days (5.3)
- *Embryo-Fetal Toxicity*: Do not administer to pregnant women unless the benefit to the mother outweighs the risk to the fetus. Inform pregnant patient of hazard (5.4, 8.1)
- *Patients with Hereditary Galactose Intolerance Lapp Lactase Deficiency or Glucose-Galactose Malabsorption*: Do not use (5.5)
- *Arrhythmias and QT Prolongation*: Correct potassium, magnesium and calcium prior to use; caution patients with proarrhythmic conditions (5.6)
- *Infusion Related Reactions (including anaphylaxis)*: Stop the infusion (5.7)
- *Dermatological Reactions*: Discontinue for exfoliative cutaneous reactions or phototoxicity. Avoid sunlight due to risk of photosensitivity (5.13)
- *Skeletal Events*: Fluorosis and periostitis with long-term voriconazole therapy. Discontinue if these events occur (5.14)

### ADVERSE REACTIONS

Most common adverse reactions (incidence  $\geq 2\%$ ): visual disturbances, fever, nausea, rash, vomiting, chills, headache, liver function test abnormal, tachycardia, hallucinations (6)

To report SUSPECTED ADVERSE REACTIONS, contact Pfizer Inc at 1-800-438-1985 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch)

### DRUG INTERACTIONS

- CYP3A4, CYP2C9, and CYP2C19 inhibitors and inducers: Adjust VFEND dosage and monitor for adverse reactions or lack of efficacy (4, 7)
- VFEND may increase the concentrations and activity of drugs that are CYP3A4, CYP2C9 and CYP2C19 substrates. Reduce dosage of these other drugs and monitor for adverse reactions (4, 7)
- Phenytoin or Efavirenz: with co-administration, increase maintenance oral and intravenous dosage of VFEND (2.3, 7)

### USE IN SPECIFIC POPULATIONS

- *Pregnancy*: Voriconazole can cause fetal harm when administered to pregnant woman. Inform pregnant women of risk to the fetus (8.1)
- *Pediatrics*: Safety/effectiveness in patients <12 years has not been established (8.4)
- *Hepatic impairment*: Use half the maintenance dose in patients with mild to moderate hepatic impairment (Child-Pugh Class A and B) (2.7)
- *Renal impairment*: Avoid intravenous administration in patients with moderate to severe renal impairment (creatinine clearance <50 mL/min) (2.8)

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

Revised: 6/2017

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## FULL PRESCRIBING INFORMATION

### 1 INDICATIONS AND USAGE

VFEND is indicated for use in patients 12 years of age and older in the treatment of the following fungal infections:

#### 1.1 Invasive Aspergillosis

In clinical trials, the majority of isolates recovered were *Aspergillus fumigatus*. There was a small number of cases of culture-proven disease due to species of *Aspergillus* other than *A. fumigatus* [see *Clinical Studies (14.1)* and *Clinical Pharmacology (12.4)*].

#### 1.2 Candidemia in Non-neutropenic Patients and the Following *Candida* Infections: Disseminated Infections in Skin and Infections in Abdomen, Kidney, Bladder Wall, and Wounds

[see *Clinical Studies (14.2)* and *Clinical Pharmacology (12.4)*]

#### 1.3 Esophageal Candidiasis

[see *Clinical Studies (14.3)* and *Clinical Pharmacology (12.4)*]

#### 1.4 Serious Fungal Infections Caused by *Scedosporium apiospermum* (Asexual Form of *Pseudallescheria boydii*) and *Fusarium* spp. Including *Fusarium solani*, in Patients Intolerant of, or Refractory to, Other Therapy

[see *Clinical Studies (14.4)* and *Clinical Pharmacology (12.4)*]

Specimens for fungal culture and other relevant laboratory studies (including histopathology) should be obtained prior to therapy to isolate and identify causative organism(s). Therapy may be instituted before the results of the cultures and other laboratory studies are known. However, once these results become available, antifungal therapy should be adjusted accordingly.

### 2 DOSAGE AND ADMINISTRATION

#### 2.1 Instructions for Use in All Patients

VFEND Tablets or Oral Suspension should be taken at least one hour before or after a meal.

VFEND I.V. for Injection requires reconstitution to 10 mg/mL and subsequent dilution to 5 mg/mL or less prior to administration as an infusion, at a maximum rate of 3 mg/kg per hour over 1 to 2 hours.

**Do not administer as an IV bolus injection.**

#### 2.2 Use of VFEND I.V. With Other Parenteral Drug Products

Blood products and concentrated electrolytes

**VFEND I.V. must not be infused concomitantly with any blood product or short-term infusion of concentrated electrolytes, even if the two infusions are running in separate intravenous lines (or cannulas).**

Electrolyte disturbances such as hypokalemia, hypomagnesemia and hypocalcemia should be corrected prior to initiation of and during VFEND therapy [see *Warnings and Precautions (5.8)*].

Intravenous solutions containing (non-concentrated) electrolytes

VFEND I.V. can be infused at the same time as other intravenous solutions containing (non-concentrated) electrolytes, but must be infused through a separate line.

Total parenteral nutrition (TPN)

VFEND I.V. can be infused at the same time as total parenteral nutrition, but must be infused in a separate line. If infused through a multiple-lumen catheter, TPN needs to be administered using a different port from the one used for VFEND I.V.

#### 2.3 Recommended Dosing in Adults

Invasive aspergillosis and serious fungal infections due to *Fusarium* spp. and *Scedosporium apiospermum*

See Table 1. Therapy must be initiated with the specified loading dose regimen of intravenous VFEND on Day 1 followed by the recommended maintenance dose (RMD) regimen. Intravenous treatment should be continued for at least 7 days. Once the patient has clinically improved and can tolerate medication given by mouth, the oral tablet form or oral suspension form of VFEND may be utilized. The recommended oral maintenance dose of 200 mg achieves a voriconazole exposure similar to 3 mg/kg IV; a 300 mg oral dose achieves an exposure similar to 4

mg/kg IV. Switching between the intravenous and oral formulations is appropriate because of the high bioavailability of the oral formulation in adults [see *Clinical Pharmacology* (12)].

#### Candidemia in non-neutropenic patients and other deep tissue *Candida* infections

See Table 1. Patients should be treated for at least 14 days following resolution of symptoms or following last positive culture, whichever is longer.

#### Esophageal Candidiasis

See Table 1. Patients should be treated for a minimum of 14 days and for at least 7 days following resolution of symptoms.

**Table 1:  
Recommended Dosing Regimen**

Infection	Loading dose	Maintenance Dose <sup>a,b</sup>	
		IV	Oral <sup>c</sup>
Invasive Aspergillosis <sup>d</sup>	6 mg/kg q12h for the first 24 hours	4 mg/kg q12h	200 mg q12h
Candidemia in nonneutropenic patients and other deep tissue <i>Candida</i> infections	6 mg/kg q12h for the first 24 hours	3-4 mg/kg q12h <sup>e</sup>	200 mg q12h
Esophageal Candidiasis	f	f	200 mg q12h
Scedosporiosis and Fusariosis	6 mg/kg q12h for the first 24 hours	4 mg/kg q12h	200 mg q12h

<sup>a</sup> Increase dose when VFEND is co-administered with phenytoin or efavirenz (7); Decrease dose in patients with hepatic impairment (2.7)

<sup>b</sup> In healthy volunteer studies, the 200 mg oral q12h dose provided an exposure (AUC<sub>0-12h</sub>) similar to a 3 mg/kg IV q12h dose; the 300 mg oral q12h dose provided an exposure (AUC<sub>0-12h</sub>) similar to a 4 mg/kg IV q12h dose [see *Clinical Pharmacology* (12)].

<sup>c</sup> Adult patients who weigh less than 40 kg should receive half of the oral maintenance dose.

<sup>d</sup> In a clinical study of invasive aspergillosis, the median duration of IV VFEND therapy was 10 days (range 2-85 days). The median duration of oral VFEND therapy was 76 days (range 2-232 days) [see *Clinical Studies* (14.1)].

<sup>e</sup> In clinical trials, patients with candidemia received 3 mg/kg IV q12h as primary therapy, while patients with other deep tissue *Candida* infections received 4 mg/kg q12h as salvage therapy. Appropriate dose should be based on the severity and nature of the infection.

<sup>f</sup> Not evaluated in patients with esophageal candidiasis.

## 2.4 Dosage Adjustment

If patient response is inadequate, the oral maintenance dose may be increased from 200 mg every 12 hours (similar to 3 mg/kg IV q12h) to 300 mg every 12 hours (similar to 4 mg/kg IV q12h). For adult patients weighing less than 40 kg, the oral maintenance dose may be increased from 100 mg every 12 hours to 150 mg every 12 hours. If patient is unable to tolerate 300 mg orally every 12 hours, reduce the oral maintenance dose by 50 mg steps to a minimum of 200 mg every 12 hours (or to 100 mg every 12 hours for adult patients weighing less than 40 kg).

If patient is unable to tolerate 4 mg/kg IV q12h, reduce the intravenous maintenance dose to 3 mg/kg q12h.

The maintenance dose of voriconazole should be increased when co-administered with phenytoin or efavirenz [see *Drug Interactions* (7)].

The maintenance dose of voriconazole should be reduced in patients with mild to moderate hepatic impairment, Child-Pugh Class A and B [see *Dosage and Administration* (2.7)]. There are no PK data to allow for dosage adjustment recommendations in patients with severe hepatic impairment (Child-Pugh Class C).

Duration of therapy should be based on the severity of the patient's underlying disease, recovery from immunosuppression, and clinical response.

## 2.5 Intravenous Administration

### Reconstitution

The powder is reconstituted with 19 mL of Water For Injection to obtain an extractable volume of 20 mL of clear concentrate containing 10 mg/mL of voriconazole. It is recommended that a standard 20 mL (non-automated) syringe be used to ensure that the exact amount (19.0 mL) of Water for Injection is dispensed. Discard the vial if a vacuum does not pull the diluent into the vial. Shake the vial until all the powder is dissolved.

#### Dilution

VFEND must be infused over 1-2 hours, at a concentration of 5 mg/mL or less. Therefore, the required volume of the 10 mg/mL VFEND concentrate should be further diluted as follows (appropriate diluents listed below):

1. Calculate the volume of 10 mg/mL VFEND concentrate required based on the patient's weight (see Table 2).
2. In order to allow the required volume of VFEND concentrate to be added, withdraw and discard at least an equal volume of diluent from the infusion bag or bottle to be used. The volume of diluent remaining in the bag or bottle should be such that when the 10 mg/mL VFEND concentrate is added, the final concentration is not less than 0.5 mg/mL nor greater than 5 mg/mL.
3. Using a suitable size syringe and aseptic technique, withdraw the required volume of VFEND concentrate from the appropriate number of vials and add to the infusion bag or bottle. **Discard Partially Used Vials.**

The final VFEND solution must be infused over 1-2 hours at a maximum rate of 3 mg/kg per hour.

**Table 2:  
Required Volumes of 10 mg/mL VFEND Concentrate**

Body Weight (kg)	Volume of VFEND Concentrate (10 mg/mL) required for:		
	3 mg/kg dose (number of vials)	4 mg/kg dose (number of vials)	6 mg/kg dose (number of vials)
30	9.0 mL (1)	12 mL (1)	18 mL (1)
35	10.5 mL (1)	14 mL (1)	21 mL (2)
40	12.0 mL (1)	16 mL (1)	24 mL (2)
45	13.5 mL (1)	18 mL (1)	27 mL (2)
50	15.0 mL (1)	20 mL (1)	30 mL (2)
55	16.5 mL (1)	22 mL (2)	33 mL (2)
60	18.0 mL (1)	24 mL (2)	36 mL (2)
65	19.5 mL (1)	26 mL (2)	39 mL (2)
70	21.0 mL (2)	28 mL (2)	42 mL (3)
75	22.5 mL (2)	30 mL (2)	45 mL (3)
80	24.0 mL (2)	32 mL (2)	48 mL (3)
85	25.5 mL (2)	34 mL (2)	51 mL (3)
90	27.0 mL (2)	36 mL (2)	54 mL (3)
95	28.5 mL (2)	38 mL (2)	57 mL (3)
100	30.0 mL (2)	40 mL (2)	60 mL (3)

VFEND I.V. for Injection is a single dose unpreserved sterile lyophile. Therefore, from a microbiological point of view, once reconstituted, the product should be used immediately. If not used immediately, in-use storage times and conditions prior to use are the responsibility of the user and should not be longer than 24 hours at 2° to 8°C (36° to 46°F). This medicinal product is for single use only and any unused solution should be discarded. Only clear solutions without particles should be used.

The reconstituted solution can be diluted with:

- 9 mg/mL (0.9%) Sodium Chloride USP
- Lactated Ringers USP
- 5% Dextrose and Lactated Ringers USP
- 5% Dextrose and 0.45% Sodium Chloride USP
- 5% Dextrose USP
- 5% Dextrose and 20 mEq Potassium Chloride USP
- 0.45% Sodium Chloride USP
- 5% Dextrose and 0.9% Sodium Chloride USP

The compatibility of VFEND I.V. with diluents other than those described above is unknown (see Incompatibilities below).

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

#### Incompatibilities

VFEND I.V. must not be diluted with 4.2% Sodium Bicarbonate Infusion. The mildly alkaline nature of this diluent caused slight degradation of VFEND after 24 hours storage at room temperature. Although refrigerated storage is recommended following reconstitution, use of this diluent is not recommended as a precautionary measure. Compatibility with other concentrations is unknown.

## **2.6 Oral Suspension**

#### Reconstitution

Tap the bottle to release the powder. Add 46 mL of water to the bottle. Shake the closed bottle vigorously for about 1 minute. Remove child-resistant cap and push bottle adaptor into the neck of the bottle. Replace the cap. Write the date of expiration of the reconstituted suspension on the bottle label (the shelf-life of the reconstituted suspension is 14 days at controlled room temperature 15-30°C [59-86°F]).

#### Instructions for use

Shake the closed bottle of reconstituted suspension for approximately 10 seconds before each use. The reconstituted oral suspension should only be administered using the oral dispenser supplied with each pack.

#### Incompatibilities

VFEND for Oral Suspension and the 40 mg/mL reconstituted oral suspension should not be mixed with any other medication or additional flavoring agent. It is not intended that the suspension be further diluted with water or other vehicles.

## **2.7 Use in Patients With Hepatic Impairment**

In the clinical program, patients were included who had baseline liver function tests (ALT, AST) up to 5 times the upper limit of normal. No dose adjustment is necessary in patients with this degree of abnormal liver function, but continued monitoring of liver function tests for further elevations is recommended [*see Warnings and Precautions (5.9)*].

It is recommended that the standard loading dose regimens be used but that the maintenance dose be halved in patients with mild to moderate hepatic cirrhosis (Child-Pugh Class A and B) [*see Clinical Pharmacology (12.3)*].

VFEND has not been studied in patients with severe hepatic cirrhosis (Child-Pugh Class C) or in patients with chronic hepatitis B or chronic hepatitis C disease. VFEND has been associated with elevations in liver function tests and clinical signs of liver damage, such as jaundice, and should only be used in patients with severe hepatic impairment if the benefit outweighs the potential risk. Patients with hepatic impairment must be carefully monitored for drug toxicity.

## **2.8 Use in Patients With Renal Impairment**

The pharmacokinetics of orally administered VFEND are not significantly affected by renal impairment. Therefore, no adjustment is necessary for oral dosing in patients with mild to severe renal impairment [*see Clinical Pharmacology (12.3)*].

In patients with moderate or severe renal impairment (creatinine clearance <50 mL/min), accumulation of the intravenous vehicle, SBECD, occurs. Oral voriconazole should be administered to these patients, unless an assessment of the benefit/risk to the patient justifies the use of intravenous voriconazole. Serum creatinine levels should be closely monitored in these patients, and, if increases occur, consideration should be given to changing to oral voriconazole therapy [*see Warnings and Precautions (5.10)*].

Voriconazole is hemodialyzed with clearance of 121 mL/min. The intravenous vehicle, SBECD, is hemodialyzed with clearance of 55 mL/min. A 4-hour hemodialysis session does not remove a sufficient amount of voriconazole to warrant dose adjustment.

### 3 DOSAGE FORMS AND STRENGTHS

#### Powder for Solution for Injection

VFEND I.V. for Injection is supplied in a single use vial as a sterile lyophilized powder equivalent to 200 mg VFEND and 3200 mg sulfobutyl ether beta-cyclodextrin sodium (SBECD).

#### Tablets

VFEND 50 mg tablets; white, film-coated, round, debossed with “Pfizer” on one side and “VOR50” on the reverse.

VFEND 200 mg tablets; white, film-coated, capsule shaped, debossed with “Pfizer” on one side and “VOR200” on the reverse.

#### Powder for Oral Suspension

VFEND for Oral Suspension is supplied in 100 mL high density polyethylene (HDPE) bottles. Each bottle contains 45 g of powder for oral suspension. Following reconstitution, the volume of the suspension is 75 mL, providing a usable volume of 70 mL (40 mg voriconazole/mL). A 5 mL oral dispenser and a press-in bottle adaptor are also provided.

### 4 CONTRAINDICATIONS

- VFEND is contraindicated in patients with known hypersensitivity to voriconazole or its excipients. There is no information regarding cross-sensitivity between VFEND (voriconazole) and other azole antifungal agents. Caution should be used when prescribing VFEND to patients with hypersensitivity to other azoles.
- Coadministration of terfenadine, astemizole, cisapride, pimozide or quinidine with VFEND is contraindicated because increased plasma concentrations of these drugs can lead to QT prolongation and rare occurrences of *torsade de pointes* [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of VFEND with sirolimus is contraindicated because VFEND significantly increases sirolimus concentrations [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of VFEND with rifampin, carbamazepine and long-acting barbiturates is contraindicated because these drugs are likely to decrease plasma voriconazole concentrations significantly [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of standard doses of voriconazole with efavirenz doses of 400 mg q24h or higher is contraindicated, because efavirenz significantly decreases plasma voriconazole concentrations in healthy subjects at these doses. Voriconazole also significantly increases efavirenz plasma concentrations [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of VFEND with high-dose ritonavir (400 mg q12h) is contraindicated because ritonavir (400 mg q12h) significantly decreases plasma voriconazole concentrations. Coadministration of voriconazole and low-dose ritonavir (100 mg q12h) should be avoided, unless an assessment of the benefit/risk to the patient justifies the use of voriconazole [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of VFEND with rifabutin is contraindicated since VFEND significantly increases rifabutin plasma concentrations and rifabutin also significantly decreases voriconazole plasma concentrations [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].
- Coadministration of VFEND with ergot alkaloids (ergotamine and dihydroergotamine) is contraindicated because VFEND may increase the plasma concentration of ergot alkaloids, which may lead to ergotism [see *Drug Interactions (7) and Clinical Pharmacology (12.3)*].

- Coadministration of VFEND with St. John's Wort is contraindicated because this herbal supplement may decrease voriconazole plasma concentration [*see Drug Interactions (7) and Clinical Pharmacology (12.3)*].

## 5 WARNINGS AND PRECAUTIONS

### 5.1 Drug Interactions

See Table 7 for a listing of drugs that may significantly alter voriconazole concentrations. Also, see Table 8 for a listing of drugs that may interact with voriconazole resulting in altered pharmacokinetics or pharmacodynamics of the other drug [*see Contraindications (4) and Drug Interactions (7)*].

### 5.2 Hepatic Toxicity

**In clinical trials, there have been uncommon cases of serious hepatic reactions during treatment with VFEND (including clinical hepatitis, cholestasis and fulminant hepatic failure, including fatalities).** Instances of hepatic reactions were noted to occur primarily in patients with serious underlying medical conditions (predominantly hematological malignancy). Hepatic reactions, including hepatitis and jaundice, have occurred among patients with no other identifiable risk factors. Liver dysfunction has usually been reversible on discontinuation of therapy [*see Warnings and Precautions (5.9) and Adverse Reactions (6.3)*].

Measure serum transaminase levels and bilirubin at the initiation of VFEND therapy and monitor at least weekly for the first month of treatment. Monitoring frequency can be reduced to monthly during continued use if no clinically significant changes are noted. If liver function tests become markedly elevated compared to baseline, VFEND should be discontinued unless the medical judgment of the benefit-risk of the treatment for the patient justifies continued use [*see Warnings and Precautions (5.9), Dosage and Administration (2.4, 2.7), and Adverse Reactions (6.3)*].

### 5.3 Visual Disturbances

**The effect of VFEND on visual function is not known if treatment continues beyond 28 days.** There have been post-marketing reports of prolonged visual adverse events, including optic neuritis and papilledema. If treatment continues beyond 28 days, visual function including visual acuity, visual field and color perception should be monitored [*see Adverse Reactions (6.2)*].

### 5.4 Embryo-Fetal Toxicity

Voriconazole can cause fetal harm when administered to a pregnant woman.

In animals, voriconazole administration was associated with teratogenicity, embryotoxicity, increased gestational length, dystocia and embryomortality [*see Use in Specific Populations (8.1)*].

If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, inform the patient of the potential hazard to the fetus.

### 5.5 Galactose Intolerance

VFEND tablets contain lactose and should not be given to patients with rare hereditary problems of galactose intolerance, Lapp lactase deficiency or glucose-galactose malabsorption.

### 5.6 Arrhythmias and QT Prolongation

Some azoles, including voriconazole, have been associated with prolongation of the QT interval on the electrocardiogram. During clinical development and post-marketing surveillance, there have been rare cases of arrhythmias, (including ventricular arrhythmias such as *torsade de pointes*), cardiac arrests and sudden deaths in patients taking voriconazole. These cases usually involved seriously ill patients with multiple confounding risk factors, such as history of cardiotoxic chemotherapy, cardiomyopathy, hypokalemia and concomitant medications that may have been contributory.

Voriconazole should be administered with caution to patients with potentially proarrhythmic conditions, such as:

- Congenital or acquired QT-prolongation
- Cardiomyopathy, in particular when heart failure is present
- Sinus bradycardia



- Existing symptomatic arrhythmias
- Concomitant medicinal product that is known to prolong QT interval [*see Contraindications (4), Drug Interactions (7), and Clinical Pharmacology (12.3)*]

Rigorous attempts to correct potassium, magnesium and calcium should be made before starting and during voriconazole therapy [*see Clinical Pharmacology (12.3)*].

### **5.7 Infusion Related Reactions**

During infusion of the intravenous formulation of voriconazole in healthy subjects, anaphylactoid-type reactions, including flushing, fever, sweating, tachycardia, chest tightness, dyspnea, faintness, nausea, pruritus and rash, have occurred uncommonly. Symptoms appeared immediately upon initiating the infusion. Consideration should be given to stopping the infusion should these reactions occur.

### **5.8 Laboratory Tests**

Electrolyte disturbances such as hypokalemia, hypomagnesemia and hypocalcemia should be corrected prior to initiation of and during VFEND therapy.

Patient management should include laboratory evaluation of renal (particularly serum creatinine) and hepatic function (particularly liver function tests and bilirubin).

### **5.9 Patients With Hepatic Impairment**

It is recommended that the standard loading dose regimens be used but that the maintenance dose be halved in patients with mild to moderate hepatic cirrhosis (Child-Pugh Class A and B) receiving VFEND [*see Clinical Pharmacology (12.3) and Dosage and Administration (2.7)*].

VFEND has not been studied in patients with severe cirrhosis (Child-Pugh Class C). VFEND has been associated with elevations in liver function tests and clinical signs of liver damage, such as jaundice, and should only be used in patients with severe hepatic impairment if the benefit outweighs the potential risk. Patients with hepatic impairment must be carefully monitored for drug toxicity.

### **5.10 Patients With Renal Impairment**

In patients with moderate to severe renal dysfunction (creatinine clearance <50 mL/min), accumulation of the intravenous vehicle, SBECD, occurs. Oral voriconazole should be administered to these patients, unless an assessment of the benefit/risk to the patient justifies the use of intravenous voriconazole. Serum creatinine levels should be closely monitored in these patients, and if increases occur, consideration should be given to changing to oral voriconazole therapy [*see Clinical Pharmacology (12.3) and Dosage and Administration (2.8)*].

### **5.11 Monitoring of Renal Function**

Acute renal failure has been observed in patients undergoing treatment with VFEND. Patients being treated with voriconazole are likely to be treated concomitantly with nephrotoxic medications and have concurrent conditions that may result in decreased renal function.

Patients should be monitored for the development of abnormal renal function. This should include laboratory evaluation, particularly serum creatinine.

### **5.12 Monitoring of Pancreatic Function**

Patients with risk factors for acute pancreatitis (e.g., recent chemotherapy, hematopoietic stem cell transplantation [HSCT]) should be monitored for the development of pancreatitis during VFEND treatment.

### **5.13 Dermatological Reactions**

Serious exfoliative cutaneous reactions, such as Stevens-Johnson syndrome, have been reported during treatment with VFEND. If a patient develops an exfoliative cutaneous reaction, VFEND should be discontinued.

VFEND has been associated with photosensitivity skin reaction. Patients, including children, should avoid exposure to direct sunlight during VFEND treatment and should use measures such as protective clothing and sunscreen with high sun protection factor (SPF). If phototoxic reactions occur, the patient should be referred to a dermatologist and

VFEND discontinuation should be considered. If VFEND is continued despite the occurrence of phototoxicity-related lesions, dermatologic evaluation should be performed on a systematic and regular basis to allow early detection and management of premalignant lesions. Squamous cell carcinoma of the skin and melanoma have been reported during long-term VFEND therapy in patients with photosensitivity skin reactions. If a patient develops a skin lesion consistent with premalignant skin lesions, squamous cell carcinoma or melanoma, VFEND should be discontinued.

The frequency of phototoxicity reactions is higher in the pediatric population. Because squamous cell carcinoma has been reported in patients who experience photosensitivity reactions, stringent measures for photoprotection are warranted in children. In children experiencing photoaging injuries such as lentigines or ephelides, sun avoidance and dermatologic follow-up are recommended even after treatment discontinuation.

#### **5.14 Skeletal Adverse Events**

Fluorosis and periostitis have been reported during long-term voriconazole therapy. If a patient develops skeletal pain and radiologic findings compatible with fluorosis or periostitis, voriconazole should be discontinued [*see Adverse Reactions (6.4)*].

### **6 ADVERSE REACTIONS**

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in 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.

#### **6.1 Overview**

The most frequently reported adverse events (all causalities) in the therapeutic trials were visual disturbances (18.7%), fever (5.7%), nausea (5.4%), rash (5.3%), vomiting (4.4%), chills (3.7%), headache (3.0%), liver function test increased (2.7%), tachycardia (2.4%), hallucinations (2.4%). The treatment-related adverse events which most often led to discontinuation of voriconazole therapy were elevated liver function tests, rash, and visual disturbances [*see Warning and Precautions (5.2, 5.3) and Adverse Reactions (6.2, 6.3)*].

#### **6.2 Clinical Trial Experience in Adults**

The data described in Table 3 reflect exposure to voriconazole in 1655 patients in the therapeutic studies. This represents a heterogeneous population, including immunocompromised patients, e.g., patients with hematological malignancy or HIV and non-neutropenic patients. This subgroup does not include healthy subjects and patients treated in the compassionate use and non-therapeutic studies. This patient population was 62% male, had a mean age of 46 years (range 11-90, including 51 patients aged 12-18 years), and was 78% White and 10% Black. Five hundred sixty one patients had a duration of voriconazole therapy of greater than 12 weeks, with 136 patients receiving voriconazole for over six months. Table 3 includes all adverse events which were reported at an incidence of  $\geq 2\%$  during voriconazole therapy in the all therapeutic studies population, studies 307/602 and 608 combined, or study 305, as well as events of concern which occurred at an incidence of  $< 2\%$ .

In study 307/602, 381 patients (196 on voriconazole, 185 on amphotericin B) were treated to compare voriconazole to amphotericin B followed by other licensed antifungal therapy in the primary treatment of patients with acute invasive aspergillosis. The rate of discontinuation from voriconazole study medication due to adverse events was 21.4% (42/196 patients). In study 608, 403 patients with candidemia were treated to compare voriconazole (272 patients) to the regimen of amphotericin B followed by fluconazole (131 patients). The rate of discontinuation from voriconazole study medication due to adverse events was 19.5% out of 272 patients. Study 305 evaluated the effects of oral voriconazole (200 patients) and oral fluconazole (191 patients) in the treatment of esophageal candidiasis. The rate of discontinuation from voriconazole study medication in Study 305 due to adverse events was 7% (14/200 patients). Laboratory test abnormalities for these studies are discussed under Clinical Laboratory Values below.

**Table 3:  
Treatment Emergent Adverse Events**

Rate ≥ 2% on Voriconazole or Adverse Events of Concern in All Therapeutic Studies Population, Studies 307/602-608 Combined, or Study 305. Possibly Related to Therapy or Causality Unknown†

	All Therapeutic Studies	Studies 307/602 and 608 (IV/ oral therapy)			Study 305 (oral therapy)	
	Voriconazole N=1655	Voriconazole N=468	Ampho B* N=185	Ampho B→ Fluconazole N=131	Voriconazole N=200	Fluconazole N=191
	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)
<b>Special Senses**</b>						
Abnormal vision	310 (18.7)	63 (13.5)	1 (0.5)	0	31 (15.5)	8 (4.2)
Photophobia	37 (2.2)	8 (1.7)	0	0	5 (2.5)	2 (1.0)
Chromatopsia	20 (1.2)	2 (0.4)	0	0	2 (1.0)	0
<b>Body as a Whole</b>						
Fever	94 (5.7)	8 (1.7)	25 (13.5)	5 (3.8)	0	0
Chills	61 (3.7)	1 (0.2)	36 (19.5)	8 (6.1)	1 (0.5)	0
Headache	49 (3.0)	9 (1.9)	8 (4.3)	1 (0.8)	0	1 (0.5)
<b>Cardiovascular System</b>						
Tachycardia	39 (2.4)	6 (1.3)	5 (2.7)	0	0	0
<b>Digestive System</b>						
Nausea	89 (5.4)	18 (3.8)	29 (15.7)	2 (1.5)	2 (1.0)	3 (1.6)
Vomiting	72 (4.4)	15 (3.2)	18 (9.7)	1 (0.8)	2 (1.0)	1 (0.5)
Liver function tests abnormal	45 (2.7)	15 (3.2)	4 (2.2)	1 (0.8)	6 (3.0)	2 (1.0)
Cholestatic jaundice	17 (1.0)	8 (1.7)	0	1 (0.8)	3 (1.5)	0
<b>Metabolic and Nutritional Systems</b>						
Alkaline phosphatase increased	59 (3.6)	19 (4.1)	4 (2.2)	3 (2.3)	10 (5.0)	3 (1.6)
Hepatic enzymes increased	30 (1.8)	11 (2.4)	5 (2.7)	1 (0.8)	3 (1.5)	0
SGOT increased	31 (1.9)	9 (1.9)	0	1 (0.8)	8 (4.0)	2 (1.0)
SGPT increased	29 (1.8)	9 (1.9)	1 (0.5)	2 (1.5)	6 (3.0)	2 (1.0)
Hypokalemia	26 (1.6)	3 (0.6)	36 (19.5)	16 (12.2)	0	0
Bilirubinemia	15 (0.9)	5 (1.1)	3 (1.6)	2 (1.5)	1 (0.5)	0
Creatinine increased	4 (0.2)	0	59 (31.9)	10 (7.6)	1 (0.5)	0
<b>Nervous System</b>						
Hallucinations	39 (2.4)	13 (2.8)	1 (0.5)	0	0	0
<b>Skin and Appendages</b>						
Rash	88 (5.3)	20 (4.3)	7 (3.8)	1 (0.8)	3 (1.5)	1 (0.5)
<b>Urogenital</b>						

	All Therapeutic Studies	Studies 307/602 and 608 (IV/ oral therapy)			Study 305 (oral therapy)	
	Voriconazole N=1655	Voriconazole N=468	Ampho B* N=185	Ampho B→ Fluconazole N=131	Voriconazole N=200	Fluconazole N=191
	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)
Kidney function abnormal	10 (0.6)	6 (1.3)	40 (21.6)	9 (6.9)	1 (0.5)	1 (0.5)
Acute kidney failure	7 (0.4)	2 (0.4)	11 (5.9)	7 (5.3)	0	0

<sup>†</sup> Study 307/602: invasive aspergillosis; Study 608: candidemia; Study 305: esophageal candidiasis

\* Amphotericin B followed by other licensed antifungal therapy

\*\* See Warnings and Precautions (5.3)

### Visual Disturbances

Voriconazole treatment-related visual disturbances are common. In therapeutic trials, approximately 21% of patients experienced abnormal vision, color vision change and/or photophobia. Visual disturbances may be associated with higher plasma concentrations and/or doses.

There have been post-marketing reports of prolonged visual adverse events, including optic neuritis and papilledema [*see Warnings and Precautions (5.3)*].

The mechanism of action of the visual disturbance is unknown, although the site of action is most likely to be within the retina. In a study in healthy subjects investigating the effect of 28-day treatment with voriconazole on retinal function, voriconazole caused a decrease in the electroretinogram (ERG) waveform amplitude, a decrease in the visual field, and an alteration in color perception. The ERG measures electrical currents in the retina. The effects were noted early in administration of voriconazole and continued through the course of study drug dosing. Fourteen days after end of dosing, ERG, visual fields and color perception returned to normal [*see Warnings and Precautions (5.7)*].

### Dermatological Reactions

Dermatological reactions were common in the patients treated with voriconazole. The mechanism underlying these dermatologic adverse events remains unknown.

Serious cutaneous reactions, including Stevens-Johnson syndrome, toxic epidermal necrolysis and erythema multiforme have been reported during treatment with VFEND. If a patient develops an exfoliative cutaneous reaction, VFEND should be discontinued.

In addition, VFEND has been associated with photosensitivity skin reactions. Patients should avoid strong, direct sunlight during VFEND therapy. In patients with photosensitivity skin reactions, squamous cell carcinoma of the skin and melanoma have been reported during long-term therapy. If a patient develops a skin lesion consistent with squamous cell carcinoma or melanoma, VFEND should be discontinued [*see Warnings and Precautions (5.13)*].

### Less Common Adverse Events

The following adverse events occurred in <2% of all voriconazole-treated patients in all therapeutic studies (N=1655). This listing includes events where a causal relationship to voriconazole cannot be ruled out or those which may help the physician in managing the risks to the patients. The list does not include events included in Table 5 above and does not include every event reported in the voriconazole clinical program.

*Body as a Whole:* abdominal pain, abdomen enlarged, allergic reaction, anaphylactoid reaction [*see Warnings and Precautions (5.6)*], ascites, asthenia, back pain, chest pain, cellulitis, edema, face edema, flank pain, flu syndrome, graft versus host reaction, granuloma, infection, bacterial infection, fungal infection, injection site pain, injection site infection/inflammation, mucous membrane disorder, multi-organ failure, pain, pelvic pain, peritonitis, sepsis, substernal chest pain.

*Cardiovascular:* atrial arrhythmia, atrial fibrillation, AV block complete, bigeminy, bradycardia, bundle branch block, cardiomegaly, cardiomyopathy, cerebral hemorrhage, cerebral ischemia, cerebrovascular accident, congestive

heart failure, deep thrombophlebitis, endocarditis, extrasystoles, heart arrest, hypertension, hypotension, myocardial infarction, nodal arrhythmia, palpitation, phlebitis, postural hypotension, pulmonary embolus, QT interval prolonged, supraventricular extrasystoles, supraventricular tachycardia, syncope, thrombophlebitis, vasodilatation, ventricular arrhythmia, ventricular fibrillation, ventricular tachycardia (including torsade de pointes) [see *Warnings and Precautions* (5.6)].

*Digestive:* anorexia, cheilitis, cholecystitis, cholelithiasis, constipation, diarrhea, duodenal ulcer perforation, duodenitis, dyspepsia, dysphagia, dry mouth, esophageal ulcer, esophagitis, flatulence, gastroenteritis, gastrointestinal hemorrhage, GGT/LDH elevated, gingivitis, glossitis, gum hemorrhage, gum hyperplasia, hematemesis, hepatic coma, hepatic failure, hepatitis, intestinal perforation, intestinal ulcer, jaundice, enlarged liver, melena, mouth ulceration, pancreatitis, parotid gland enlargement, periodontitis, proctitis, pseudomembranous colitis, rectal disorder, rectal hemorrhage, stomach ulcer, stomatitis, tongue edema.

*Endocrine:* adrenal cortex insufficiency, diabetes insipidus, hyperthyroidism, hypothyroidism.

*Hemic and Lymphatic:* agranulocytosis, anemia (macrocytic, megaloblastic, microcytic, normocytic), aplastic anemia, hemolytic anemia, bleeding time increased, cyanosis, DIC, ecchymosis, eosinophilia, hypervolemia, leukopenia, lymphadenopathy, lymphangitis, marrow depression, pancytopenia, petechia, purpura, enlarged spleen, thrombocytopenia, thrombotic thrombocytopenic purpura.

*Metabolic and Nutritional:* albuminuria, BUN increased, creatine phosphokinase increased, edema, glucose tolerance decreased, hypercalcemia, hypercholesteremia, hyperglycemia, hyperkalemia, hypermagnesemia, hypernatremia, hyperuricemia, hypocalcemia, hypoglycemia, hypomagnesemia, hyponatremia, hypophosphatemia, peripheral edema, uremia.

*Musculoskeletal:* arthralgia, arthritis, bone necrosis, bone pain, leg cramps, myalgia, myasthenia, myopathy, osteomalacia, osteoporosis.

*Nervous System:* abnormal dreams, acute brain syndrome, agitation, akathisia, amnesia, anxiety, ataxia, brain edema, coma, confusion, convulsion, delirium, dementia, depersonalization, depression, diplopia, dizziness, encephalitis, encephalopathy, euphoria, Extrapyramidal Syndrome, grand mal convulsion, Guillain-Barré syndrome, hypertonia, hypesthesia, insomnia, intracranial hypertension, libido decreased, neuralgia, neuropathy, nystagmus, oculogyric crisis, paresthesia, psychosis, somnolence, suicidal ideation, tremor, vertigo.

*Respiratory System:* cough increased, dyspnea, epistaxis, hemoptysis, hypoxia, lung edema, pharyngitis, pleural effusion, pneumonia, respiratory disorder, respiratory distress syndrome, respiratory tract infection, rhinitis, sinusitis, voice alteration.

*Skin and Appendages:* alopecia, angioedema, contact dermatitis, discoid lupus erythematosus, eczema, erythema multiforme, exfoliative dermatitis, fixed drug eruption, furunculosis, herpes simplex, maculopapular rash, melanoma, melanosis, photosensitivity skin reaction, pruritus, pseudoporphyria, psoriasis, skin discoloration, skin disorder, skin dry, Stevens-Johnson syndrome, squamous cell carcinoma, sweating, toxic epidermal necrolysis, urticaria.

*Special Senses:* abnormality of accommodation, blepharitis, color blindness, conjunctivitis, corneal opacity, deafness, ear pain, eye pain, eye hemorrhage, dry eyes, hypoacusis, keratitis, keratoconjunctivitis, mydriasis, night blindness, optic atrophy, optic neuritis, otitis externa, papilledema, retinal hemorrhage, retinitis, scleritis, taste loss, taste perversion, tinnitus, uveitis, visual field defect.

*Urogenital:* anuria, blighted ovum, creatinine clearance decreased, dysmenorrhea, dysuria, epididymitis, glycosuria, hemorrhagic cystitis, hematuria, hydronephrosis, impotence, kidney pain, kidney tubular necrosis, metrorrhagia, nephritis, nephrosis, oliguria, scrotal edema, urinary incontinence, urinary retention, urinary tract infection, uterine hemorrhage, vaginal hemorrhage.

### **6.3 Clinical Laboratory Values**

The overall incidence of clinically significant transaminase abnormalities in all therapeutic studies was 12.4% (206/1655) of patients treated with voriconazole. Increased incidence of liver function test abnormalities may be associated with higher plasma concentrations and/or doses. The majority of abnormal liver function tests either

resolved during treatment without dose adjustment or following dose adjustment, including discontinuation of therapy.

Voriconazole has been infrequently associated with cases of serious hepatic toxicity including cases of jaundice and rare cases of hepatitis and hepatic failure leading to death. Most of these patients had other serious underlying conditions.

Liver function tests should be evaluated at the start of and during the course of VFEND therapy. Patients who develop abnormal liver function tests during VFEND therapy should be monitored for the development of more severe hepatic injury. Patient management should include laboratory evaluation of hepatic function (particularly liver function tests and bilirubin). Discontinuation of VFEND must be considered if clinical signs and symptoms consistent with liver disease develop that may be attributable to VFEND [see *Warnings and Precautions (5.2)*].

Acute renal failure has been observed in severely ill patients undergoing treatment with VFEND. Patients being treated with voriconazole are likely to be treated concomitantly with nephrotoxic medications and have concurrent conditions that may result in decreased renal function. It is recommended that patients are monitored for the development of abnormal renal function. This should include laboratory evaluation, particularly serum creatinine.

Tables 4 to 6 show the number of patients with hypokalemia and clinically significant changes in renal and liver function tests in three randomized, comparative multicenter studies. In study 305, patients with esophageal candidiasis were randomized to either oral voriconazole or oral fluconazole. In study 307/602, patients with definite or probable invasive aspergillosis were randomized to either voriconazole or amphotericin B therapy. In study 608, patients with candidemia were randomized to either voriconazole or the regimen of amphotericin B followed by fluconazole.

**Table 4:**  
**Protocol 305 – Patients with Esophageal Candidiasis**  
**Clinically Significant Laboratory Test Abnormalities**

	<b>Criteria*</b>	<b>Voriconazole</b>	<b>Fluconazole</b>
		<b>n/N (%)</b>	<b>n /N (%)</b>
T. Bilirubin	>1.5x ULN	8/185 (4.3)	7/186 (3.8)
AST	>3.0x ULN	38/187 (20.3)	15/186 (8.1)
ALT	>3.0x ULN	20/187 (10.7)	12/186 (6.5)
Alk phos	>3.0x ULN	19/187 (10.2)	14/186 (7.5)

\* Without regard to baseline value

n = number of patients with a clinically significant abnormality while on study therapy

N = total number of patients with at least one observation of the given lab test while on study therapy

ULN = upper limit of normal

**Table 5:**  
**Protocol 307/602 – Primary Treatment of Invasive Aspergillosis**  
**Clinically Significant Laboratory Test Abnormalities**

	<b>Criteria*</b>	<b>Voriconazole</b>	<b>Amphotericin B**</b>
		<b>n/N (%)</b>	<b>n/N (%)</b>
T. Bilirubin	>1.5x ULN	35/180 (19.4)	46/173 (26.6)
AST	>3.0x ULN	21/180 (11.7)	18/174 (10.3)
ALT	>3.0x ULN	34/180 (18.9)	40/173 (23.1)
Alk phos	>3.0x ULN	29/181 (16.0)	38/173 (22.0)
Creatinine	>1.3x ULN	39/182 (21.4)	102/177 (57.6)
Potassium	<0.9x LLN	30/181 (16.6)	70/178 (39.3)

\* Without regard to baseline value

\*\* Amphotericin B followed by other licensed antifungal therapy

n = number of patients with a clinically significant abnormality while on study therapy

N = total number of patients with at least one observation of the given lab test while on study therapy

ULN = upper limit of normal  
 LLN = lower limit of normal

**Table 6:**  
**Protocol 608 – Treatment of Candidemia**  
**Clinically Significant Laboratory Test Abnormalities**

	Criteria*	Voriconazole	Amphotericin B followed by Fluconazole
		n/N (%)	n/N (%)
T. Bilirubin	>1.5x ULN	50/261 (19.2)	31/115 (27.0)
AST	>3.0x ULN	40/261 (15.3)	16/116 (13.8)
ALT	>3.0x ULN	22/261 (8.4)	15/116 (12.9)
Alk phos	>3.0x ULN	59/261 (22.6)	26/115 (22.6)
Creatinine	>1.3x ULN	39/260 (15.0)	32/118 (27.1)
Potassium	<0.9x LLN	43/258 (16.7)	35/118 (29.7)

\* Without regard to baseline value

n = number of patients with a clinically significant abnormality while on study therapy

N = total number of patients with at least one observation of the given lab test while on study therapy

ULN = upper limit of normal

LLN = lower limit of normal

#### 6.4 Postmarketing Experience

The following adverse reactions have been identified during post approval use of voriconazole. 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.

*Skeletal*: fluorosis and periostitis have been reported during long-term voriconazole therapy [see *Warnings and Precautions (5.14)*].

#### 7 DRUG INTERACTIONS

**Table 7:**  
**Effect of Other Drugs on Voriconazole Pharmacokinetics [see *Clinical Pharmacology (12.3)*]**

Drug/Drug Class (Mechanism of Interaction by the Drug)	Voriconazole Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> after 200 mg q12h)	Recommendations for Voriconazole Dosage Adjustment/Comments
Rifampin* and Rifabutin* (CYP450 Induction)	Significantly Reduced	<b>Contraindicated</b>
Efavirenz (400 mg q24h)** (CYP450 Induction)	Significantly Reduced	<b>Contraindicated</b>
Efavirenz (300 mg q24h)** (CYP450 Induction)	Slight Decrease in AUC <sub>τ</sub>	When voriconazole is coadministered with efavirenz, voriconazole oral maintenance dose should be increased to 400 mg q12h and efavirenz should be decreased to 300 mg q24h
High-dose Ritonavir (400 mg q12h)** (CYP450 Induction)	Significantly Reduced	<b>Contraindicated</b>
Low-dose Ritonavir (100 mg q12h)** (CYP450 Induction)	Reduced	Coadministration of voriconazole and low-dose ritonavir (100 mg q12h) should be avoided, unless an assessment of the benefit/risk to the patient justifies the use of

Drug/Drug Class (Mechanism of Interaction by the Drug)	Voriconazole Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> after 200 mg q12h)	Recommendations for Voriconazole Dosage Adjustment/Comments
		voriconazole
Carbamazepine (CYP450 Induction)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Likely to Result in Significant Reduction	<b>Contraindicated</b>
Long Acting Barbiturates (CYP450 Induction)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Likely to Result in Significant Reduction	<b>Contraindicated</b>
Phenytoin* (CYP450 Induction)	Significantly Reduced	Increase voriconazole maintenance dose from 4 mg/kg to 5 mg/kg IV q12h or from 200 mg to 400 mg orally q12h (100 mg to 200 mg orally q12h in patients weighing less than 40 kg)
St. John's Wort (CYP450 inducer; P-gp inducer)	Significantly Reduced	<b>Contraindicated</b>
Oral Contraceptives** containing ethinyl estradiol and norethindrone (CYP2C19 Inhibition)	Increased	Monitoring for adverse events and toxicity related to voriconazole is recommended when coadministered with oral contraceptives
Fluconazole** (CYP2C9, CYP2C19 and CYP3A4 Inhibition)	Significantly Increased	Avoid concomitant administration of voriconazole and fluconazole. Monitoring for adverse events and toxicity related to voriconazole is started within 24 h after the last dose of fluconazole.
Other HIV Protease Inhibitors (CYP3A4 Inhibition)	<i>In Vivo</i> Studies Showed No Significant Effects of Indinavir on Voriconazole Exposure  <i>In Vitro</i> Studies Demonstrated Potential for Inhibition of Voriconazole Metabolism (Increased Plasma Exposure)	No dosage adjustment in the voriconazole dosage needed when coadministered with indinavir  Frequent monitoring for adverse events and toxicity related to voriconazole when coadministered with other HIV protease inhibitors
Other NNRTIs*** (CYP3A4 Inhibition or CYP450 Induction)	<i>In Vitro</i> Studies Demonstrated Potential for Inhibition of Voriconazole Metabolism by Delavirdine and Other NNRTIs (Increased Plasma Exposure)  A Voriconazole-Efavirenz Drug Interaction Study Demonstrated the Potential for the Metabolism of Voriconazole to be Induced by Efavirenz and Other NNRTIs (Decreased Plasma Exposure)	Frequent monitoring for adverse events and toxicity related to voriconazole  Careful assessment of voriconazole effectiveness

\* Results based on *in vivo* clinical studies generally following repeat oral dosing with 200 mg q12h voriconazole to healthy subjects

\*\* Results based on *in vivo* clinical study following repeat oral dosing with 400 mg q12h for 1 day, then 200 mg q12h for at least 2 days voriconazole to healthy subjects

\*\*\* Non-Nucleoside Reverse Transcriptase Inhibitors

**Table 8:**  
**Effect of Voriconazole on Pharmacokinetics of Other Drugs [see *Clinical Pharmacology* (12.3)]**

Drug/Drug Class (Mechanism of Interaction by Voriconazole)	Drug Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> )	Recommendations for Drug Dosage Adjustment/Comments
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Drug/Drug Class (Mechanism of Interaction by Voriconazole)	Drug Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> )	Recommendations for Drug Dosage Adjustment/Comments
Sirolimus* (CYP3A4 Inhibition)	Significantly Increased	<b>Contraindicated</b>
Rifabutin* (CYP3A4 Inhibition)	Significantly Increased	<b>Contraindicated</b>
Efavirenz (400 mg q24h)** (CYP3A4 Inhibition)  Efavirenz (300 mg q24h)** (CYP3A4 Inhibition)	Significantly Increased  Slight Increase in AUC <sub>τ</sub>	<b>Contraindicated</b>  When voriconazole is coadministered with efavirenz, voriconazole oral maintenance dose should be increased to 400 mg q12h and efavirenz should be decreased to 300 mg q24h
High-dose Ritonavir (400 mg q12h)**(CYP3A4 Inhibition)  Low-dose Ritonavir (100 mg q12h)**	No Significant Effect of Voriconazole on Ritonavir C <sub>max</sub> or AUC <sub>τ</sub>  Slight Decrease in Ritonavir C <sub>max</sub> and AUC <sub>τ</sub>	<b>Contraindicated</b> because of significant reduction of voriconazole C <sub>max</sub> and AUC <sub>τ</sub>  Coadministration of voriconazole and low-dose ritonavir (100 mg q12h) should be avoided (due to the reduction in voriconazole C <sub>max</sub> and AUC <sub>τ</sub> ) unless an assessment of the benefit/risk to the patient justifies the use of voriconazole
Terfenadine, Astemizole, Cisapride, Pimozide, Quinidine (CYP3A4 Inhibition)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Drug Plasma Exposure Likely to be Increased	<b>Contraindicated</b> because of potential for QT prolongation and rare occurrence of <i>torsade de pointes</i>
Ergot Alkaloids (CYP450 Inhibition)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Drug Plasma Exposure Likely to be Increased	<b>Contraindicated</b>
Cyclosporine* (CYP3A4 Inhibition)	AUC <sub>τ</sub> Significantly Increased; No Significant Effect on C <sub>max</sub>	When initiating therapy with VFEND in patients already receiving cyclosporine, reduce the cyclosporine dose to one-half of the starting dose and follow with frequent monitoring of cyclosporine blood levels. Increased cyclosporine levels have been associated with nephrotoxicity. When VFEND is discontinued, cyclosporine concentrations must be frequently monitored and the dose increased as necessary.
Methadone*** (CYP3A4 Inhibition)	Increased	Increased plasma concentrations of methadone have been associated with toxicity including QT prolongation. Frequent monitoring for adverse events and toxicity related to methadone is recommended during coadministration. Dose reduction of methadone may be needed
Fentanyl (CYP3A4 Inhibition)	Increased	Reduction in the dose of fentanyl and other long-acting opiates metabolized by CYP3A4 should be considered when

<b>Drug/Drug Class (Mechanism of Interaction by Voriconazole)</b>	<b>Drug Plasma Exposure (C<sub>max</sub> and AUC<sub>τ</sub>)</b>	<b>Recommendations for Drug Dosage Adjustment/Comments</b>
		coadministered with VFEND. Extended and frequent monitoring for opiate-associated adverse events may be necessary [ <i>see Drug Interactions (7)</i> ]
Alfentanil (CYP3A4 Inhibition)	Significantly Increased	Reduction in the dose of alfentanil and other opiates metabolized by CYP3A4 (e.g., sufentanil) should be considered when coadministered with VFEND. A longer period for monitoring respiratory and other opiate-associated adverse events may be necessary [ <i>see Drug Interactions (7)</i> ].
Oxycodone (CYP3A4 Inhibition)	Significantly Increased	Reduction in the dose of oxycodone and other long-acting opiates metabolized by CYP3A4 should be considered when coadministered with VFEND. Extended and frequent monitoring for opiate-associated adverse events may be necessary [ <i>see Drug Interactions (7)</i> ].
NSAIDs**** including ibuprofen and diclofenac (CYP2C9 Inhibition)	Increased	Frequent monitoring for adverse events and toxicity related to NSAIDs. Dose reduction of NSAIDs may be needed [ <i>see Drug Interactions (7)</i> ].
Tacrolimus* (CYP3A4 Inhibition)	Significantly Increased	When initiating therapy with VFEND in patients already receiving tacrolimus, reduce the tacrolimus dose to one-third of the starting dose and follow with frequent monitoring of tacrolimus blood levels. Increased tacrolimus levels have been associated with nephrotoxicity. When VFEND is discontinued, tacrolimus concentrations must be frequently monitored and the dose increased as necessary.
Phenytoin* (CYP2C9 Inhibition)	Significantly Increased	Frequent monitoring of phenytoin plasma concentrations and frequent monitoring of adverse effects related to phenytoin.
Oral Contraceptives containing ethinyl estradiol and norethindrone (CYP3A4 Inhibition)**	Increased	Monitoring for adverse events related to oral contraceptives is recommended during coadministration.
Warfarin* (CYP2C9 Inhibition)	Prothrombin Time Significantly Increased	Monitor PT or other suitable anti-coagulation tests. Adjustment of warfarin dosage may be needed.
Omeprazole* (CYP2C19/3A4 Inhibition)	Significantly Increased	When initiating therapy with VFEND in patients already receiving omeprazole doses of 40

Drug/Drug Class (Mechanism of Interaction by Voriconazole)	Drug Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> )	Recommendations for Drug Dosage Adjustment/Comments
		mg or greater, reduce the omeprazole dose by one-half. The metabolism of other proton pump inhibitors that are CYP2C19 substrates may also be inhibited by voriconazole and may result in increased plasma concentrations of other proton pump inhibitors.
Other HIV Protease Inhibitors (CYP3A4 Inhibition)	<i>In Vivo</i> Studies Showed No Significant Effects on Indinavir Exposure  <i>In Vitro</i> Studies Demonstrated Potential for Voriconazole to Inhibit Metabolism (Increased Plasma Exposure)	No dosage adjustment for indinavir when coadministered with VFEND  Frequent monitoring for adverse events and toxicity related to other HIV protease inhibitors
Other NNRTIs***** (CYP3A4 Inhibition)	A Voriconazole-Efavirenz Drug Interaction Study Demonstrated the Potential for Voriconazole to Inhibit Metabolism of Other NNRTIs (Increased Plasma Exposure)	Frequent monitoring for adverse events and toxicity related to NNRTI
Benzodiazepines (CYP3A4 Inhibition)	<i>In Vitro</i> Studies Demonstrated Potential for Voriconazole to Inhibit Metabolism (Increased Plasma Exposure)	Frequent monitoring for adverse events and toxicity (i.e., prolonged sedation) related to benzodiazepines metabolized by CYP3A4 (e.g., midazolam, triazolam, alprazolam). Adjustment of benzodiazepine dosage may be needed.
HMG-CoA Reductase Inhibitors (Statins) (CYP3A4 Inhibition)	<i>In Vitro</i> Studies Demonstrated Potential for Voriconazole to Inhibit Metabolism (Increased Plasma Exposure)	Frequent monitoring for adverse events and toxicity related to statins. Increased statin concentrations in plasma have been associated with rhabdomyolysis. Adjustment of the statin dosage may be needed.
Dihydropyridine Calcium Channel Blockers (CYP3A4 Inhibition)	<i>In Vitro</i> Studies Demonstrated Potential for Voriconazole to Inhibit Metabolism (Increased Plasma Exposure)	Frequent monitoring for adverse events and toxicity related to calcium channel blockers. Adjustment of calcium channel blocker dosage may be needed.
Sulfonylurea Oral Hypoglycemics (CYP2C9 Inhibition)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Drug Plasma Exposure Likely to be Increased	Frequent monitoring of blood glucose and for signs and symptoms of hypoglycemia. Adjustment of oral hypoglycemic drug dosage may be needed.
Vinca Alkaloids (CYP3A4 Inhibition)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Drug Plasma Exposure Likely to be Increased	Frequent monitoring for adverse events and toxicity (i.e., neurotoxicity) related to vinca alkaloids. Reserveazole antifungals, including voriconazole, for patients receiving a vinca alkaloid who have no alternative antifungal treatment options.
Everolimus (CYP3A4 Inhibition)	Not Studied <i>In Vivo</i> or <i>In Vitro</i> , but Drug Plasma Exposure Likely to be Increased	Concomitant administration of voriconazole and everolimus is not recommended.

Drug/Drug Class (Mechanism of Interaction by Voriconazole)	Drug Plasma Exposure (C <sub>max</sub> and AUC <sub>τ</sub> )	Recommendations for Drug Dosage Adjustment/Comments

<sup>†</sup> Results based on *in vivo* clinical studies generally following repeat oral dosing with 200 mg BID voriconazole to healthy subjects

<sup>\*\*</sup> Results based on *in vivo* clinical study following repeat oral dosing with 400 mg q12h for 1 day, then 200 mg q12h for at least 2 days voriconazole to healthy subjects

<sup>\*\*\*</sup> Results based on *in vivo* clinical study following repeat oral dosing with 400 mg q12h for 1 day, then 200 mg q12h for 4 days voriconazole to subjects receiving a methadone maintenance dose (30-100 mg q24h)

<sup>\*\*\*\*</sup> Non-Steroidal Anti-Inflammatory Drug

<sup>\*\*\*\*\*</sup> Non-Nucleoside Reverse Transcriptase Inhibitors

## 8 USE IN SPECIFIC POPULATIONS

### 8.1 Pregnancy

#### Risk Summary

Voriconazole can cause fetal harm when administered to a pregnant woman. There are no available data on the use of VFEND in pregnant women. In animal reproduction studies, oral voriconazole was teratogenic in rats and embryotoxic in rabbits. Cleft palates and hydronephrosis/hydroureter were observed in rat pups exposed to voriconazole during organogenesis at and above 10 mg/kg (0.3 times the recommended maintenance dose of 200 mg every 12 hours based on body surface area comparisons). In rabbits, embryomortality, reduced fetal weight and increased incidence of skeletal variations, cervical ribs and extrasternal ossification sites were observed in pups when pregnant rabbits were orally dosed at 100 mg/kg (6 times the RMD based on body surface area comparisons) during organogenesis. Rats exposed to voriconazole from implantation to weaning experienced increased gestational length and dystocia, which were associated with increased perinatal pup mortality at the 10 mg/kg dose. [see Data]. If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, inform the patient of the potential hazard to the fetus [see Warnings and Precautions (5.4)].

The background risk of major birth defects and miscarriage for the indicated populations is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20% respectively.

#### Data

##### *Animal Data*

Voriconazole was administered orally to pregnant rats during organogenesis (gestation days 6-17) at 10, 30, and 60 mg/kg/day. Voriconazole was teratogenic with increased incidences in hydroureter and hydronephrosis at 10 mg/kg/day or greater, approximately 0.3 times the recommended human dose (RMD) based on mg/m<sup>2</sup>, and cleft palate at 60 mg/kg, approximately 2 times the recommended human dose (RMD) based on mg/m<sup>2</sup>. Reduced ossification of sacral and caudal vertebrae, skull, pubic, and hyoid bone, supernumerary ribs, anomalies of the sternbrae, and dilatation of the ureter/renal pelvis were also observed at doses of 10 mg/kg or greater. There was no evidence of maternal toxicity at any dose.

Voriconazole was administered orally to pregnant rabbits during the period of organogenesis (gestation days 7-19) at 10, 40, and 100 mg/kg/day. Voriconazole produced embryofetal toxicity (increased post-implantation loss, decreased fetal body weight) in association with maternal toxicity (decreased body weight gain and food consumption) at 100 mg/kg/day (6 times the RMD based on mg/m<sup>2</sup>). Fetal skeletal variations (increases in the incidence of cervical rib and extra sternbral ossification sites) were observed at 100 mg/kg/day.

In a peri- and postnatal toxicity study in rats, voriconazole was administered orally to female rats from implantation through the end of lactation at 1, 3, and 10 mg/kg/day. Voriconazole prolonged the duration of gestation and labor and produced dystocia with related increases in maternal mortality and decreases in perinatal survival of F1 pups at 10 mg/kg/day, approximately 0.3 times the RMD.

## 8.2 Lactation

### Risk Summary

No data are available regarding the presence of voriconazole in human milk, the effects of voriconazole on the breastfed infant, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for VFEND and any potential adverse effects on the breastfed child from VFEND or from the underlying maternal condition.

## 8.3 Females and Males of Reproductive Potential

### Contraception

Advise females of reproductive potential to use effective contraception during treatment with VFEND. The coadministration of voriconazole with the oral contraceptive, Ortho-Novum<sup>®</sup> (35 mcg ethinyl estradiol and 1 mg norethindrone), results in an interaction between these two drugs, but is unlikely to reduce the contraceptive effect. Monitoring for adverse reactions associated with oral contraceptives and voriconazole is recommended [*see Drug Interactions (7) and Clinical Pharmacology (12.3)*].

## 8.4 Pediatric Use

Safety and effectiveness in pediatric patients below the age of 12 years have not been established.

A total of 22 patients aged 12 to 18 years with invasive aspergillosis were included in the therapeutic studies. Twelve out of 22 (55%) patients had successful response after treatment with a maintenance dose of voriconazole 4 mg/kg q12h.

Sparse plasma sampling for pharmacokinetics in adolescents was conducted in the therapeutic studies [*see Clinical Pharmacology (12.3)*]. A population pharmacokinetic analysis was conducted on pooled data from 35 immunocompromised pediatric patients aged 2 to <12 years old who were included in two pharmacokinetic studies of intravenous voriconazole (single dose and multiple dose). Twenty-four of these patients received multiple intravenous maintenance doses of 3 mg/kg and 4 mg/kg. A comparison of the pediatric and adult population pharmacokinetic data revealed that the predicted average steady state plasma concentrations were similar at the maintenance dose of 4 mg/kg every 12 hours in children and 3 mg/kg every 12 hours in adults (medians of 1.19 µg/mL and 1.16 µg/mL in children and adults, respectively).

There have been postmarketing reports of pancreatitis in pediatric patients.

## 8.5 Geriatric Use

In multiple dose therapeutic trials of voriconazole, 9.2% of patients were ≥65 years of age and 1.8% of patients were ≥75 years of age. In a study in healthy subjects, the systemic exposure (AUC) and peak plasma concentrations ( $C_{max}$ ) were increased in elderly males compared to young males. Pharmacokinetic data obtained from 552 patients from 10 voriconazole therapeutic trials showed that voriconazole plasma concentrations in the elderly patients were approximately 80% to 90% higher than those in younger patients after either IV or oral administration. However, the overall safety profile of the elderly patients was similar to that of the young so no dosage adjustment is recommended [*see Clinical Pharmacology (12.3)*].

## 10 OVERDOSAGE

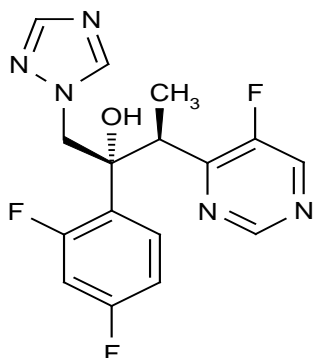
In clinical trials, there were three cases of accidental overdose. All occurred in pediatric patients who received up to five times the recommended intravenous dose of voriconazole. A single adverse event of photophobia of 10 minutes duration was reported.

There is no known antidote to voriconazole.

Voriconazole is hemodialyzed with clearance of 121 mL/min. The intravenous vehicle, SBECD, is hemodialyzed with clearance of 55 mL/min. In an overdose, hemodialysis may assist in the removal of voriconazole and SBECD from the body.

## 11 DESCRIPTION

VFEND<sup>®</sup> (voriconazole), an azole antifungal agent is available as a lyophilized powder for solution for intravenous infusion, film-coated tablets for oral administration, and as a powder for oral suspension. The structural formula is:



Voriconazole is designated chemically as (2R,3S)-2-(2, 4-difluorophenyl)-3-(5-fluoro-4-pyrimidinyl)-1-(1H-1,2,4-triazol-1-yl)-2-butanol with an empirical formula of C<sub>16</sub>H<sub>14</sub>F<sub>3</sub>N<sub>5</sub>O and a molecular weight of 349.3.

Voriconazole drug substance is a white to light-colored powder.

VFEND I.V. is a white lyophilized powder containing nominally 200 mg voriconazole and 3200 mg sulfobutyl ether beta-cyclodextrin sodium in a 30 mL Type I clear glass vial.

VFEND I.V. is intended for administration by intravenous infusion. It is a single-dose, unpreserved product. Vials containing 200 mg lyophilized voriconazole are intended for reconstitution with Water for Injection to produce a solution containing 10 mg/mL VFEND and 160 mg/mL of sulfobutyl ether beta-cyclodextrin sodium. The resultant solution is further diluted prior to administration as an intravenous infusion [*see Dosage and Administration (2)*].

VFEND Tablets contain 50 mg or 200 mg of voriconazole. The inactive ingredients include lactose monohydrate, pregelatinized starch, croscarmellose sodium, povidone, magnesium stearate and a coating containing hypromellose, titanium dioxide, lactose monohydrate and triacetin.

VFEND for Oral Suspension is a white to off-white powder providing a white to off-white orange-flavored suspension when reconstituted. Bottles containing 45 g powder for oral suspension are intended for reconstitution with water to produce a suspension containing 40 mg/mL voriconazole. The inactive ingredients include colloidal silicon dioxide, titanium dioxide, xanthan gum, sodium citrate dihydrate, sodium benzoate, anhydrous citric acid, natural orange flavor, and sucrose.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Voriconazole is an antifungal drug [*see Microbiology (12.4)*]

### 12.2 Pharmacodynamics

#### Exposure-Response Relationship For Efficacy and Safety

In 10 clinical trials (N=1121), the median values for the average and maximum voriconazole plasma concentrations in individual patients across these studies was 2.51 µg/mL (inter-quartile range 1.21 to 4.44 µg/mL) and 3.79 µg/mL (inter-quartile range 2.06 to 6.31 µg/mL), respectively. A pharmacokinetic-pharmacodynamic analysis of patient data from 6 of these 10 clinical trials (N=280) could not detect a positive association between mean, maximum or minimum plasma voriconazole concentration and efficacy. However, pharmacokinetic/pharmacodynamic analyses of the data from all 10 clinical trials identified positive associations between plasma voriconazole concentrations and rate of both liver function test abnormalities and visual disturbances [*see Adverse Reactions (6)*].

#### Cardiac Electrophysiology

A placebo-controlled, randomized, crossover study to evaluate the effect on the QT interval of healthy male and female subjects was conducted with three single oral doses of voriconazole and ketoconazole. Serial ECGs and plasma samples were obtained at specified intervals over a 24-hour post dose observation period. The placebo-adjusted mean maximum increases in QTc from baseline after 800, 1200, and 1600 mg of voriconazole and after ketoconazole 800 mg were all <10 msec. Females exhibited a greater increase in QTc than males, although all mean changes were <10 msec. Age was not found to affect the magnitude of increase in QTc. No subject in any group had an increase in QTc of  $\geq 60$  msec from baseline. No subject experienced an interval exceeding the potentially clinically relevant threshold of 500 msec. However, the QT effect of voriconazole combined with drugs known to prolong the QT interval is unknown [see *Contraindications (4) and Drug Interactions (7)*].

### 12.3 Pharmacokinetics

The pharmacokinetics of voriconazole have been characterized in healthy subjects, special populations and patients.

The pharmacokinetics of voriconazole are non-linear due to saturation of its metabolism. The interindividual variability of voriconazole pharmacokinetics is high. Greater than proportional increase in exposure is observed with increasing dose. It is estimated that, on average, increasing the oral dose from 200 mg q12h to 300 mg q12h leads to an approximately 2.5-fold increase in exposure ( $AUC_{\tau}$ ); similarly, increasing the intravenous dose from 3 mg/kg q12h to 4 mg/kg q12h produces an approximately 2.5-fold increase in exposure (Table 9).

**Table 9:**  
**Geometric Mean (%CV) Plasma Voriconazole Pharmacokinetic Parameters in Adults Receiving Different Dosing Regimens**

	<b>6 mg/kg IV (loading dose)</b>	<b>3 mg/kg IV q12h</b>	<b>4 mg/kg IV q12h</b>	<b>400 mg Oral (loading dose)</b>	<b>200 mg Oral q12h</b>	<b>300 mg Oral q12h</b>
N	35	23	40	17	48	16
$AUC_{12}$ ( $\mu\text{g}\cdot\text{h/mL}$ )	13.9 (32)	13.7 (53)	33.9 (54)	9.31 (38)	12.4 (78)	34.0 (53)
$C_{\text{max}}$ ( $\mu\text{g/mL}$ )	3.13 (20)	3.03 (25)	4.77 (36)	2.30 (19)	2.31 (48)	4.74 (35)
$C_{\text{min}}$ ( $\mu\text{g/mL}$ )	--	0.46 (97)	1.73 (74)	--	0.46 (120)	1.63 (79)

Note: Parameters were estimated based on non-compartmental analysis from 5 pharmacokinetic studies.  $AUC_{12}$  = area under the curve over 12 hour dosing interval,  $C_{\text{max}}$  = maximum plasma concentration,  $C_{\text{min}}$  = minimum plasma concentration. CV = coefficient of variation.

When the recommended intravenous loading dose regimen is administered to healthy subjects, plasma concentrations close to steady state are achieved within the first 24 hours of dosing (eg, 6 mg/kg IV q12h on day 1 followed by 3 mg/kg IV q12h). Without the loading dose, accumulation occurs during twice-daily multiple dosing with steady-state plasma voriconazole concentrations being achieved by day 6 in the majority of subjects.

#### Absorption

The pharmacokinetic properties of voriconazole are similar following administration by the intravenous and oral routes. Based on a population pharmacokinetic analysis of pooled data in healthy subjects (N=207), the oral bioavailability of voriconazole is estimated to be 96% (CV 13%). Bioequivalence was established between the 200 mg tablet and the 40 mg/mL oral suspension when administered as a 400 mg q12h loading dose followed by a 200 mg q12h maintenance dose.

Maximum plasma concentrations ( $C_{\text{max}}$ ) are achieved 1-2 hours after dosing. When multiple doses of voriconazole are administered with high-fat meals, the mean  $C_{\text{max}}$  and  $AUC_{\tau}$  are reduced by 34% and 24%, respectively when administered as a tablet and by 58% and 37% respectively when administered as the oral suspension [see *Dosage and Administration (2)*].

In healthy subjects, the absorption of voriconazole is not affected by coadministration of oral ranitidine, cimetidine, or omeprazole, drugs that are known to increase gastric pH.

#### Distribution









itraconazole. *In vitro* studies also show that the major metabolite of voriconazole, voriconazole N-oxide, inhibits the metabolic activity of CYP2C9 and CYP3A4 to a greater extent than that of CYP2C19. Therefore, there is potential for voriconazole and its major metabolite to increase the systemic exposure (plasma concentrations) of other drugs metabolized by these CYP450 enzymes.

***The systemic exposure of the following drugs is significantly increased or is expected to be significantly increased by coadministration of voriconazole and their use is contraindicated:***

***Sirolimus (CYP3A4 substrate)***—Repeat dose administration of oral voriconazole (400 mg q12h for 1 day, then 200 mg q12h for 8 days) increased the  $C_{max}$  and AUC of sirolimus (2 mg single dose) an average of 7-fold (90% CI: 5.7, 7.5) and 11-fold (90% CI: 9.9, 12.6), respectively, in healthy male subjects. ***Coadministration of voriconazole and sirolimus is contraindicated*** [see *Contraindications (4) and Warnings and Precautions (5.1)*].

***Terfenadine, astemizole, cisapride, pimozone and quinidine (CYP3A4 substrates)***—Although not studied *in vitro* or *in vivo*, concomitant administration of voriconazole with terfenadine, astemizole, cisapride, pimozone or quinidine may result in inhibition of the metabolism of these drugs. Increased plasma concentrations of these drugs can lead to QT prolongation and rare occurrences of torsade de pointes. ***Coadministration of voriconazole and terfenadine, astemizole, cisapride, pimozone and quinidine is contraindicated*** [see *Contraindications (4) and Warnings and Precautions (5.1)*].

***Ergot alkaloids***—Although not studied *in vitro* or *in vivo*, voriconazole may increase the plasma concentration of ergot alkaloids (ergotamine and dihydroergotamine) and lead to ergotism. ***Coadministration of voriconazole with ergot alkaloids is contraindicated*** [see *Contraindications (4) and Warnings and Precautions (5.1)*].

***Everolimus (CYP3A4 substrate, P-gp substrate)***—Although not studied *in vitro* or *in vivo*, voriconazole may increase plasma concentrations of everolimus, which could potentially lead to exacerbation of everolimus toxicity. Currently there are insufficient data to allow dosing recommendations in this situation. Therefore, co-administration of voriconazole with everolimus is not recommended [see *Drug Interactions (7)*].

***Coadministration of voriconazole with the following agents results in increased exposure or is expected to result in increased exposure to these drugs. Therefore, careful monitoring and/or dosage adjustment of these drugs is needed:***

***Alfentanil (CYP3A4 substrate)***—Coadministration of multiple doses of oral voriconazole (400 mg q12h on day 1, 200 mg q12h on day 2) with a single 20 mcg/kg intravenous dose of alfentanil with concomitant naloxone resulted in a 6-fold increase in mean alfentanil  $AUC_{0-\infty}$  and a 4-fold prolongation of mean alfentanil elimination half-life, compared to when alfentanil was given alone. An increase in the incidence of delayed and persistent alfentanil-associated nausea and vomiting during co-administration of voriconazole and alfentanil was also observed. Reduction in the dose of alfentanil or other opiates that are also metabolized by CYP3A4 (e.g., sufentanil), and extended close monitoring of patients for respiratory and other opiate-associated adverse events, may be necessary when any of these opiates is coadministered with voriconazole [see *Warnings and Precautions (5.1)*].

***Fentanyl (CYP3A4 substrate)***: In an independent published study, concomitant use of voriconazole (400 mg q12h on Day 1, then 200 mg q12h on Day 2) with a single intravenous dose of fentanyl (5  $\mu$ g/kg) resulted in an increase in the mean  $AUC_{0-\infty}$  of fentanyl by 1.4-fold (range 0.81- to 2.04-fold). When voriconazole is co-administered with fentanyl IV, oral or transdermal dosage forms, extended and frequent monitoring of patients for respiratory depression and other fentanyl-associated adverse events is recommended, and fentanyl dosage should be reduced if warranted [see *Warnings and Precautions (5.1)*].

***Oxycodone (CYP3A4 substrate)***: In an independent published study, coadministration of multiple doses of oral voriconazole (400 mg q12h, on Day 1 followed by five doses of 200 mg q12h on Days 2 to 4) with a single 10 mg oral dose of oxycodone on Day 3 resulted in an increase in the mean  $C_{max}$  and  $AUC_{0-\infty}$  of oxycodone by 1.7-fold (range 1.4- to 2.2-fold) and 3.6-fold (range 2.7- to 5.6-fold), respectively. The mean elimination half-life of oxycodone was also increased by 2.0-fold (range 1.4- to 2.5-fold). Voriconazole also increased the visual effects (heterophoria and miosis) of oxycodone. A reduction in oxycodone dosage may be needed during voriconazole treatment to avoid opioid related adverse effects. Extended and frequent monitoring for adverse effects associated with oxycodone and other long-acting opiates metabolized by CYP3A4 is recommended [see *Warnings and Precautions (5.1)*].

**Cyclosporine (CYP3A4 substrate)**—In stable renal transplant recipients receiving chronic cyclosporine therapy, concomitant administration of oral voriconazole (200 mg q12h for 8 days) increased cyclosporine  $C_{max}$  and  $AUC_{\tau}$  an average of 1.1 times (90% CI: 0.9, 1.41) and 1.7 times (90% CI: 1.5, 2.0), respectively, as compared to when cyclosporine was administered without voriconazole. When initiating therapy with voriconazole in patients already receiving cyclosporine, it is recommended that the cyclosporine dose be reduced to one-half of the original dose and followed with frequent monitoring of the cyclosporine blood levels. Increased cyclosporine levels have been associated with nephrotoxicity. When voriconazole is discontinued, cyclosporine levels should be frequently monitored and the dose increased as necessary [see Warnings and Precautions (5.1)].

**Methadone (CYP3A4, CYP2C19, CYP2C9 substrate)**—Repeat dose administration of oral voriconazole (400 mg q12h for 1 day, then 200 mg q12h for 4 days) increased the  $C_{max}$  and  $AUC_{\tau}$  of pharmacologically active Rmethadone by 31% (90% CI: 22%, 40%) and 47% (90% CI: 38%, 57%), respectively, in subjects receiving a methadone maintenance dose (30-100 mg q24h). The  $C_{max}$  and  $AUC$  of (S)-methadone increased by 65% (90% CI: 53%, 79%) and 103% (90% CI: 85%, 124%), respectively. Increased plasma concentrations of methadone have been associated with toxicity including QT prolongation. Frequent monitoring for adverse events and toxicity related to methadone is recommended during coadministration. Dose reduction of methadone may be needed [see Warnings and Precautions (5.1)].

**Tacrolimus (CYP3A4 substrate)**—Repeat oral dose administration of voriconazole (400 mg q12h x 1 day, then 200 mg q12h x 6 days) increased tacrolimus (0.1 mg/kg single dose)  $C_{max}$  and  $AUC_{\tau}$  in healthy subjects by an average of 2-fold (90% CI: 1.9, 2.5) and 3-fold (90% CI: 2.7, 3.8), respectively. When initiating therapy with voriconazole in patients already receiving tacrolimus, it is recommended that the tacrolimus dose be reduced to one-third of the original dose and followed with frequent monitoring of the tacrolimus blood levels. Increased tacrolimus levels have been associated with nephrotoxicity. When voriconazole is discontinued, tacrolimus levels should be carefully monitored and the dose increased as necessary [see Warnings and Precautions (5.1)].

**Warfarin (CYP2C9 substrate)**—Coadministration of voriconazole (300 mg q12h x 12 days) with warfarin (30 mg single dose) significantly increased maximum prothrombin time by approximately 2 times that of placebo in healthy subjects. Close monitoring of prothrombin time or other suitable anticoagulation tests is recommended if warfarin and voriconazole are coadministered and the warfarin dose adjusted accordingly [see Warnings and Precautions (5.1)].

**Oral Coumarin Anticoagulants (CYP2C9, CYP3A4 substrates)**—Although not studied *in vitro* or *in vivo*, voriconazole may increase the plasma concentrations of coumarin anticoagulants and therefore may cause an increase in prothrombin time. If patients receiving coumarin preparations are treated simultaneously with voriconazole, the prothrombin time or other suitable anti-coagulation tests should be monitored at close intervals and the dosage of anticoagulants adjusted accordingly [see Warnings and Precautions (5.1)].

**Statins (CYP3A4 substrates)**—Although not studied clinically, voriconazole has been shown to inhibit lovastatin metabolism *in vitro* (human liver microsomes). Therefore, voriconazole is likely to increase the plasma concentrations of statins that are metabolized by CYP3A4. It is recommended that dose adjustment of the statin be considered during coadministration. Increased statin concentrations in plasma have been associated with rhabdomyolysis [see Warnings and Precautions (5.1)].

**Benzodiazepines (CYP3A4 substrates)**—Although not studied clinically, voriconazole has been shown to inhibit midazolam metabolism *in vitro* (human liver microsomes). Therefore, voriconazole is likely to increase the plasma concentrations of benzodiazepines that are metabolized by CYP3A4 (e.g., midazolam, triazolam, and alprazolam) and lead to a prolonged sedative effect. It is recommended that dose adjustment of the benzodiazepine be considered during coadministration [see Warnings and Precautions (5.1)].

**Calcium Channel Blockers (CYP3A4 substrates)**—Although not studied clinically, voriconazole has been shown to inhibit felodipine metabolism *in vitro* (human liver microsomes). Therefore, voriconazole may increase the plasma concentrations of calcium channel blockers that are metabolized by CYP3A4. Frequent monitoring for adverse events and toxicity related to calcium channel blockers is recommended during coadministration. Dose adjustment of the calcium channel blocker may be needed [see Warnings and Precautions (5.1)].

**Sulfonylureas (CYP2C9 substrates)**—Although not studied *in vitro* or *in vivo*, voriconazole may increase plasma concentrations of sulfonylureas (e.g., tolbutamide, glipizide, and glyburide) and therefore cause hypoglycemia.

Frequent monitoring of blood glucose and appropriate adjustment (i.e., reduction) of the sulfonyleurea dosage is recommended during coadministration [see *Warnings and Precautions* (5.1)].

**Vinca Alkaloids (CYP3A4 substrates)**—Although not studied *in vitro* or *in vivo*, voriconazole may increase the plasma concentrations of the vinca alkaloids (e.g., vincristine and vinblastine) and lead to neurotoxicity. Therefore, reserve azole antifungals, including voriconazole, for patients receiving a vinca alkaloid, including vincristine, who have no alternative antifungal treatment options [see *Warnings and Precautions* (5.1)].

**Non-Steroidal Anti-Inflammatory Drugs (NSAIDs; CYP2C9 substrates):** In two independent published studies, single doses of ibuprofen (400 mg) and diclofenac (50 mg) were coadministered with the last dose of voriconazole (400 mg q12h on Day 1, followed by 200 mg q12h on Day 2). Voriconazole increased the mean  $C_{max}$  and AUC of the pharmacologically active isomer, S (+)-ibuprofen by 20% and 100%, respectively. Voriconazole increased the mean  $C_{max}$  and AUC of diclofenac by 114% and 78%, respectively.

A reduction in ibuprofen and diclofenac dosage may be needed during concomitant administration with voriconazole. Patients receiving voriconazole concomitantly with other NSAIDs (e.g., celecoxib, naproxen, lornoxicam, meloxicam) that are also metabolized by CYP2C9 should be carefully monitored for NSAID-related adverse events and toxicity, and dosage reduction should be made if warranted [see *Warnings and Precautions* (5.1)].

**No significant pharmacokinetic interactions were observed when voriconazole was coadministered with the following agents. Therefore, no dosage adjustment for these agents is recommended:**

**Prednisolone (CYP3A4 substrate)**—Voriconazole (200 mg q12h x 30 days) increased  $C_{max}$  and AUC of prednisolone (60 mg single dose) by an average of 11% and 34%, respectively, in healthy subjects.

**Digoxin (P-glycoprotein mediated transport)**—Voriconazole (200 mg q12h x 12 days) had no significant effect on steady state  $C_{max}$  and  $AUC_{\tau}$  of digoxin (0.25 mg once daily for 10 days) in healthy subjects.

**Mycophenolic acid (UDP-glucuronyl transferase substrate)**—Voriconazole (200 mg q12h x 5 days) had no significant effect on the  $C_{max}$  and  $AUC_{\tau}$  of mycophenolic acid and its major metabolite, mycophenolic acid glucuronide after administration of a 1 g single oral dose of mycophenolate mofetil.

## Two-Way Interactions

**Concomitant use of the following agents with voriconazole is contraindicated:**

**Rifabutin (potent CYP450 inducer)**—Rifabutin (300 mg once daily) decreased the  $C_{max}$  and  $AUC_{\tau}$  of voriconazole at 200 mg twice daily by an average of 67% (90% CI: 58%, 73%) and 79% (90% CI: 71%, 84%), respectively, in healthy subjects. During coadministration with rifabutin (300 mg once daily), the steady state  $C_{max}$  and  $AUC_{\tau}$  of voriconazole following an increased dose of 400 mg twice daily were on average approximately 2 times higher, compared with voriconazole alone at 200 mg twice daily. Coadministration of voriconazole at 400 mg twice daily with rifabutin 300 mg twice daily increased the  $C_{max}$  and  $AUC_{\tau}$  of rifabutin by an average of 3-times (90% CI: 2.2, 4.0) and 4 times (90% CI: 3.5, 5.4), respectively, compared to rifabutin given alone. **Coadministration of voriconazole and rifabutin is contraindicated** [see *Contraindications* (4)].

**Significant drug interactions that may require dosage adjustment, frequent monitoring of drug levels and/or frequent monitoring of drug-related adverse events/toxicity:**

**Efavirenz, a non-nucleoside reverse transcriptase inhibitor (CYP450 inducer; CYP3A4 inhibitor and substrate)**—Standard doses of voriconazole and efavirenz (400 mg q24h or higher) must not be coadministered [see *Drug Interactions* (7)]. Steady state efavirenz (400 mg PO q24h) decreased the steady state  $C_{max}$  and  $AUC_{\tau}$  of voriconazole (400 mg PO q12h for 1 day, then 200 mg PO q12h for 8 days) by an average of 61% and 77%, respectively, in healthy male subjects. Voriconazole at steady state (400 mg PO q12h for 1 day, then 200 mg q12h for 8 days) increased the steady state  $C_{max}$  and  $AUC_{\tau}$  of efavirenz (400 mg PO q24h for 9 days) by an average of 38% and 44%, respectively, in healthy subjects.

The pharmacokinetics of adjusted doses of voriconazole and efavirenz were studied in healthy male subjects following administration of voriconazole (400 mg PO q12h on Days 2 to 7) with efavirenz (300 mg PO q24h on Days 1-7), relative to steady-state administration of voriconazole (400 mg for 1 day, then 200 mg PO q12h for 2

days) or efavirenz (600 mg q24h for 9 days). Coadministration of voriconazole 400 mg q12h with efavirenz 300 mg q24h, decreased voriconazole  $AUC_{\tau}$  by 7% (90% CI: -23%, 13%) and increased  $C_{max}$  by 23% (90% CI: -1%, 53%); efavirenz  $AUC_{\tau}$  was increased by 17% (90% CI: 6%, 29%) and  $C_{max}$  was equivalent.

**Coadministration of standard doses of voriconazole and efavirenz (400 mg q24h or higher) is contraindicated.**

Voriconazole may be coadministered with efavirenz if the voriconazole maintenance dose is increased to 400 mg q12h and the efavirenz dose is decreased to 300 mg q24h. When treatment with voriconazole is stopped, the initial dosage of efavirenz should be restored [see *Dosage and Administration (2.4)*, *Contraindications (4)*, and *Drug Interactions (7)*].

**Phenytoin (CYP2C9 substrate and potent CYP450 inducer)**—Repeat dose administration of phenytoin (300 mg once daily) decreased the steady state  $C_{max}$  and  $AUC_{\tau}$  of orally administered voriconazole (200 mg q12h x 14 days) by an average of 50% and 70%, respectively, in healthy subjects. Administration of a higher voriconazole dose (400 mg q12h x 7 days) with phenytoin (300 mg once daily) resulted in comparable steady state voriconazole  $C_{max}$  and  $AUC_{\tau}$  estimates as compared to when voriconazole was given at 200 mg q12h without phenytoin.

Phenytoin may be coadministered with voriconazole if the maintenance dose of voriconazole is increased from 4 mg/kg to 5 mg/kg intravenously every 12 hours or from 200 mg to 400 mg orally, every 12 hours (100 mg to 200 mg orally, every 12 hours in patients less than 40 kg) [see *Dosage and Administration (2.4)* and *Drug Interactions (7)*].

Repeat dose administration of voriconazole (400 mg q12h x 10 days) increased the steady state  $C_{max}$  and  $AUC_{\tau}$  of phenytoin (300 mg once daily) by an average of 70% and 80%, respectively, in healthy subjects. The increase in phenytoin  $C_{max}$  and AUC when coadministered with voriconazole may be expected to be as high as 2 times the  $C_{max}$  and AUC estimates when phenytoin is given without voriconazole. Therefore, frequent monitoring of plasma phenytoin concentrations and phenytoin-related adverse effects is recommended when phenytoin is coadministered with voriconazole [see *Warnings and Precautions (5.1)*].

**Omeprazole (CYP2C19 inhibitor; CYP2C19 and CYP3A4 substrate)**—Coadministration of omeprazole (40 mg once daily x 10 days) with oral voriconazole (400 mg q12h x 1 day, then 200 mg q12h x 9 days) increased the steady state  $C_{max}$  and  $AUC_{\tau}$  of voriconazole by an average of 15% (90% CI: 5%, 25%) and 40% (90% CI: 29%, 55%), respectively, in healthy subjects. No dosage adjustment of voriconazole is recommended.

Coadministration of voriconazole (400 mg q12h x 1 day, then 200 mg x 6 days) with omeprazole (40 mg once daily x 7 days) to healthy subjects significantly increased the steady state  $C_{max}$  and  $AUC_{\tau}$  of omeprazole an average of 2 times (90% CI: 1.8, 2.6) and 4 times (90% CI: 3.3, 4.4), respectively, as compared to when omeprazole is given without voriconazole. When initiating voriconazole in patients already receiving omeprazole doses of 40 mg or greater, it is recommended that the omeprazole dose be reduced by one-half [see *Warnings and Precautions (5.1)*].

The metabolism of other proton pump inhibitors that are CYP2C19 substrates may also be inhibited by voriconazole and may result in increased plasma concentrations of these drugs.

**Oral Contraceptives (CYP3A4 substrate; CYP2C19 inhibitor)**—Coadministration of oral voriconazole (400 mg q12h for 1 day, then 200 mg q12h for 3 days) and oral contraceptive (Ortho-Novum1/35<sup>®</sup> consisting of 35 mcg ethinyl estradiol and 1 mg norethindrone, q24h) to healthy female subjects at steady state increased the  $C_{max}$  and  $AUC_{\tau}$  of ethinyl estradiol by an average of 36% (90% CI: 28%, 45%) and 61% (90% CI: 50%, 72%), respectively, and that of norethindrone by 15% (90% CI: 3%, 28%) and 53% (90% CI: 44%, 63%), respectively in healthy subjects. Voriconazole  $C_{max}$  and  $AUC_{\tau}$  increased by an average of 14% (90% CI: 3%, 27%) and 46% (90% CI: 32%, 61%), respectively. Monitoring for adverse events related to oral contraceptives, in addition to those for voriconazole, is recommended during coadministration [see *Warnings and Precautions (5.1)*].

**No significant pharmacokinetic interaction was seen and no dosage adjustment of these drugs is recommended:**

**Indinavir (CYP3A4 inhibitor and substrate)**—Repeat dose administration of indinavir (800 mg TID for 10 days) had no significant effect on voriconazole  $C_{max}$  and AUC following repeat dose administration (200 mg q12h for 17 days) in healthy subjects.

Repeat dose administration of voriconazole (200 mg q12h for 7 days) did not have a significant effect on steady state  $C_{max}$  and  $AUC_{\tau}$  of indinavir following repeat dose administration (800 mg TID for 7 days) in healthy subjects.

### Other Two-Way Interactions Expected to be Significant Based on *In Vitro* and *In Vivo* Findings:

**Other HIV Protease Inhibitors (CYP3A4 substrates and inhibitors)**—*In vitro* studies (human liver microsomes) suggest that voriconazole may inhibit the metabolism of HIV protease inhibitors (e.g., saquinavir, amprenavir and nelfinavir). *In vitro* studies (human liver microsomes) also show that the metabolism of voriconazole may be inhibited by HIV protease inhibitors (e.g., saquinavir and amprenavir). Patients should be frequently monitored for drug toxicity during the coadministration of voriconazole and HIV protease inhibitors [see *Warnings and Precautions* (5.1)].

**Other Non-Nucleoside Reverse Transcriptase Inhibitors (NNRTIs) (CYP3A4 substrates, inhibitors or CYP450 inducers)**—*In vitro* studies (human liver microsomes) show that the metabolism of voriconazole may be inhibited by a NNRTI (e.g., delavirdine). The findings of a clinical voriconazole-efavirenz drug interaction study in healthy male subjects suggest that the metabolism of voriconazole may be induced by a NNRTI. This *in vivo* study also showed that voriconazole may inhibit the metabolism of a NNRTI [see *Drug Interactions* (7) and *Warnings and Precautions* (5.9)]. Patients should be frequently monitored for drug toxicity during the coadministration of voriconazole and other NNRTIs (e.g., nevirapine and delavirdine) [see *Warnings and Precautions* (5.1)]. Dose adjustments are required when voriconazole is co-administered with efavirenz [see *Drug Interactions* (7) and *Warnings and Precautions* (5.1)].

## 12.4 Microbiology

### Mechanism of Action

Voriconazole is an azole antifungal drug. The primary mode of action of voriconazole is the inhibition of fungal cytochrome P-450-mediated 14 alpha-lanosterol demethylation, an essential step in fungal ergosterol biosynthesis. The accumulation of 14 alpha-methyl sterols correlates with the subsequent loss of ergosterol in the fungal cell wall and may be responsible for the antifungal activity of voriconazole.

### Drug Resistance

Voriconazole drug resistance development has not been adequately studied *in vitro* against *Candida*, *Aspergillus*, *Scedosporium* and *Fusarium* species. The frequency of drug resistance development for the various fungi for which this drug is indicated is not known.

Fungal isolates exhibiting reduced susceptibility to fluconazole or itraconazole may also show reduced susceptibility to voriconazole, suggesting cross-resistance can occur among these azoles. The relevance of cross-resistance and clinical outcome has not been fully characterized. Clinical cases where azole cross-resistance is demonstrated may require alternative antifungal therapy.

### Activity *In Vitro* and *In Vivo*

Voriconazole has been shown to be active against most strains of the following microorganisms, **both *in vitro* and in clinical infections**.

*Aspergillus fumigatus*

*Aspergillus flavus*

*Aspergillus niger*

*Aspergillus terreus*

*Candida albicans*

*Candida glabrata* (In clinical studies, the voriconazole MIC<sub>90</sub> was 4 µg/mL)\*

*Candida krusei*

*Candida parapsilosis*

*Candida tropicalis*

*Fusarium* spp. including *Fusarium solani*

*Scedosporium apiospermum*

\* In clinical studies, voriconazole MIC<sub>90</sub> for *C. glabrata* baseline isolates was 4 µg/mL; 13/50 (26%) *C. glabrata* baseline isolates were resistant (MIC ≥4 µg/mL) to voriconazole. However, based on 1054 isolates tested in surveillance studies the MIC<sub>90</sub> was 1 µg/mL (see Table 12).

The following data are available, **but their clinical significance is unknown**. At least 90 percent of the following fungi exhibit an *in vitro* minimum inhibitory concentration (MIC) less than or equal to the susceptible breakpoint for voriconazole against isolates of similar genus or organism group. However, the effectiveness of voriconazole in treating clinical infections due to these fungi has not been established in adequate and well-controlled clinical trials:

*Candida lusitanae*  
*Candida guilliermondii*

Susceptibility Testing Methods<sup>1,2</sup>

***Aspergillus* species and other filamentous fungi**

No interpretive criteria have been established for *Aspergillus* species and other filamentous fungi.

***Candida* species**

The interpretive standards for voriconazole against *Candida* species are applicable only to tests performed using Clinical Laboratory and Standards Institute (CLSI) microbroth dilution reference method M27 for MIC read at 48 hours or disk diffusion reference method M44 for zone diameter read at 24 hours.<sup>1,2</sup>

**Broth Microdilution Techniques**—Quantitative methods are used to determine antifungal MICs. These MICs provide estimates of the susceptibility of *Candida* spp. to antifungal agents. MICs should be determined using a standardized procedure at 48 hours.<sup>1</sup> Standardized procedures are based on a microdilution method (broth) with standardized inoculum concentrations and standardized concentrations of voriconazole powder. The MIC values should be interpreted according to the criteria provided in Table 10.

**Diffusion Techniques**—Qualitative methods that require measurement of zone diameters also provide reproducible estimates of the susceptibility of *Candida* spp. to an antifungal agent. One such standardized procedure requires the use of standardized inoculum concentrations.<sup>2</sup> This procedure uses paper disks impregnated with 1 µg of voriconazole to test the susceptibility of yeasts to voriconazole at 24 hours. Disk diffusion interpretive criteria are also provided in Table 10.

**Table 10:  
 Susceptibility Interpretive Criteria for Voriconazole<sup>1,2</sup>**

	Broth Microdilution at 48 hours (MIC in µg/mL)			Disk Diffusion at 24 hours (Zone diameters in mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)	Susceptible (S)	Intermediate (I)	Resistant (R)
Voriconazole	≤1.0	2.0	≥4.0	≥17	14-16	≤13

**NOTE:** Shown are the breakpoints (µg/mL) for voriconazole against *Candida* species.

A report of *Susceptible* (S) indicates that the antimicrobial drug is likely to inhibit growth of the microorganism if the antimicrobial drug reaches the concentration usually achievable at the site of infection. A report of *intermediate* (I) indicates that the result should be considered equivocal, and, if the microorganism is not fully susceptible to alternative, clinically feasible drugs, the test should be repeated. This category implies possible clinical applicability in body sites where the drugs are physiologically concentrated or when a high dosage of drug is used. This category also provides a buffer zone that prevents small uncontrolled technical factors from causing major discrepancies in interpretation. A report of *Resistant* (R) indicates that the antimicrobial is not likely to inhibit growth of the pathogen if the antimicrobial drug reaches the concentrations usually achievable at the infection site; other therapy should be selected.

Quality Control

Standardized susceptibility test procedures require the use of quality control organisms to ensure the accuracy of the technical aspects of the test procedures. Standard voriconazole powder and 1 µg disks should provide the following range of values noted in Table 11.



**NOTE:** Quality control microorganisms are specific strains of organisms with intrinsic biological properties relating to resistance mechanisms and their genetic expression within fungi; the specific strains used for microbiological control are not clinically significant.

**Table 11:**  
**Acceptable Quality Control Ranges for Voriconazole to be used in Validation of Susceptibility Test Results**

QC Strain	Broth Microdilution (MIC in $\mu\text{g/mL}$ ) at 48-hour	Disk Diffusion (Zone diameter in mm) at 24-hour
<i>Candida parapsilosis</i> ATCC 22019	0.03-0.25	28-37
<i>Candida krusei</i> ATCC 6258	0.12-1.0	16-25
<i>Candida albicans</i> ATCC 90028	*	31-42

\* Quality control ranges have not been established for this strain/antifungal agent combination due to their extensive interlaboratory variation during initial quality control studies.  
ATCC is a registered trademark of the American Type Culture Collection.

## 12.5 Pharmacogenomics

CYP2C19, significantly involved in the metabolism of voriconazole, exhibits genetic polymorphism. Approximately 15-20% of Asian populations may be expected to be poor metabolizers. For Caucasians and Blacks, the prevalence of poor metabolizers is 3-5%. Studies conducted in Caucasian and Japanese healthy subjects have shown that poor metabolizers have, on average, 4-fold higher voriconazole exposure ( $AUC_t$ ) than their homozygous extensive metabolizer counterparts. Subjects who are heterozygous extensive metabolizers have, on average, 2-fold higher voriconazole exposure than their homozygous extensive metabolizer counterparts [see *Clinical Pharmacology* (12.3)].

## 13 NONCLINICAL TOXICOLOGY

### 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Two-year carcinogenicity studies were conducted in rats and mice. Rats were given oral doses of 6, 18 or 50 mg/kg voriconazole, or 0.2, 0.6, or 1.6 times the recommended maintenance dose on a  $\text{mg/m}^2$  basis. Hepatocellular adenomas were detected in females at 50 mg/kg and hepatocellular carcinomas were found in males at 6 and 50 mg/kg. Mice were given oral doses of 10, 30 or 100 mg/kg voriconazole, or 0.1, 0.4, or 1.4 times the RMD on a  $\text{mg/m}^2$  basis. In mice, hepatocellular adenomas were detected in males and females and hepatocellular carcinomas were detected in males at 1.4 times the RMD of voriconazole.

Voriconazole demonstrated clastogenic activity (mostly chromosome breaks) in human lymphocyte cultures *in vitro*. Voriconazole was not genotoxic in the Ames assay, CHO HGPRT assay, the mouse micronucleus assay or the *in vivo* DNA repair test (Unscheduled DNA Synthesis assay).

Voriconazole administration induced no impairment of male or female fertility in rats dosed at 50 mg/kg, or 1.6 times the RMD (recommended maintenance dose).

## 14 CLINICAL STUDIES

Voriconazole, administered orally or parenterally, has been evaluated as primary or salvage therapy in 520 patients aged 12 years and older with infections caused by *Aspergillus* spp., *Fusarium* spp., and *Scedosporium* spp.

### 14.1 Invasive Aspergillosis

Voriconazole was studied in patients for primary therapy of invasive aspergillosis (randomized, controlled study 307/602), for primary and salvage therapy of aspergillosis (non-comparative study 304) and for treatment of patients with invasive aspergillosis who were refractory to, or intolerant of, other antifungal therapy (non-comparative study 309/604).

#### Study 307/602 – Primary Therapy of Invasive Aspergillosis











VFEND Tablets should be stored at 15° - 30°C (59° - 86°F) [see USP Controlled Room Temperature].

VFEND Powder for Oral Suspension should be stored at 2° - 8°C (36° - 46° F) (in a refrigerator) before reconstitution. The shelf-life of the powder for oral suspension is 18 months.

The reconstituted suspension should be stored at 15° - 30°C (59° - 86°F) [see USP Controlled Room Temperature]. Do not refrigerate or freeze. Keep the container tightly closed. The shelf-life of the reconstituted suspension is 14 days. Any remaining suspension should be discarded 14 days after reconstitution.

## **17 PATIENT COUNSELING INFORMATION**

Advise the Patient to read the FDA-Approved Patient Labeling

### Embryo-Fetal Toxicity

- Advise female patients of the potential risks to a fetus.
- Advise females of reproductive potential to use effective contraception during treatment with VFEND.

This product's label may have been updated. For current full prescribing information, please visit [www.pfizer.com](http://www.pfizer.com).



LAB-0271-32.2a2

**VFEND<sup>®</sup> /VEE-fend/**

**(voriconazole) IV injection, tablets, liquid**

Read the Patient Information that comes with VFEND before you start taking it and each time you get a refill. There may be new information. This information does not take the place of talking with your healthcare provider about your condition or treatment.

**What is VFEND?**

VFEND is a prescription medicine used to treat certain serious fungal infections in your blood and body. These infections are called “aspergillosis,” “esophageal candidiasis,” “*Scedosporium*,” “*Fusarium*,” and “candidemia”.

It is not known if VFEND is safe and effective in children younger than 12 years old.

**Who should not take VFEND?**

**Do not take VFEND if you:**

- **are allergic to voriconazole or any of the ingredients in VFEND.** See the end of this leaflet for a complete list of ingredients in VFEND.
- **are taking any of the following medicines:**
  - cisapride (Propulsid<sup>®</sup>)
  - pimozone (Orap<sup>®</sup>)
  - quinidine (like Quinaglute<sup>®</sup>)
  - sirolimus (Rapamune<sup>®</sup>)
  - rifampin (Rifadin<sup>®</sup>)
  - carbamazepine (Tegretol<sup>®</sup>)
  - long-acting barbiturates like phenobarbital (Luminal<sup>®</sup>)
  - efavirenz (Sustiva<sup>®</sup>)
  - ritonavir (Norvir<sup>®</sup>)
  - rifabutin (Mycobutin<sup>®</sup>)
  - ergotamine, dihydroergotamine (ergot alkaloids)
  - St. John’s Wort (herbal supplement)

Ask your healthcare provider or pharmacist if you are not sure if you are taking any of the medicines listed above.

Do not start taking a new medicine without talking to your healthcare provider or pharmacist.

**What should I tell my healthcare provider before taking VFEND?**

Before you take VFEND, tell your healthcare provider if you:

- have or ever had heart disease, or an abnormal heart rate or rhythm. Your healthcare provider may order a test to check your heart (EKG) before starting VFEND.
- have liver or kidney problems. Your healthcare provider may do blood tests to make sure you can take VFEND.
- have trouble digesting dairy products, lactose (milk sugar), or regular table sugar. VFEND tablets contain lactose. VFEND liquid contains sucrose (table sugar).
- are pregnant or plan to become pregnant. VFEND can harm your unborn baby. Talk to your healthcare provider if you are pregnant or plan to become pregnant. Women who can become pregnant should use effective birth control while taking VFEND.



- are breast-feeding or plan to breast-feed. It is not known if VFEND passes into breast milk. Talk to your healthcare provider about the best way to feed your baby if you take VFEND.

**Tell your healthcare provider about all the medicines you take**, including prescription and non-prescription medicines, vitamins and herbal supplements.

VFEND may affect the way other medicines work, and other medicines may affect how VFEND works.

Know what medicines you take. Keep a list of them to show your healthcare provider or pharmacist when you get a new medicine.

### **How should I take VFEND?**

- **VFEND may be prescribed to you as:**

- VFEND I.V. (intravenous infusion) or
- VFEND tablets or
- VFEND oral suspension

- VFEND I.V. will be given to you by a healthcare provider over 1 to 2 hours.
- Take VFEND tablets or oral suspension exactly as your healthcare provider tells you to.
- Take VFEND tablets or oral suspension at least 1 hour before or at least 1 hour after meals.

VFEND oral suspension will be mixed for you by your pharmacist. **Do not** mix VFEND oral suspension with any other medicine, flavored liquid, or syrup.

- If you take too much VFEND, call your healthcare provider or go to the nearest hospital emergency room.

### **What should I avoid while taking VFEND?**

- You should not drive at night while taking VFEND. VFEND can cause changes in your vision such as blurring or sensitivity to light.
- Do not drive or operate machinery, or do other dangerous activities until you know how VFEND affects you.
- Avoid direct sunlight. VFEND can make your skin sensitive to the sun and the light from sunlamps and tanning beds. You could get a severe sunburn. Use sunscreen and wear a hat and clothes that cover your skin if you have to be in sunlight. Talk to your healthcare provider if you get sunburn.

### **What are possible side effects of VFEND?**

VFEND may cause serious side effects including:

- **liver problems.** Symptoms of liver problems may include:
  - itchy skin
  - yellowing of your eyes
  - feeling very tired
  - flu-like symptoms
  - nausea or vomiting
- **vision changes.** Symptoms of vision changes may include:
  - blurred vision
  - changes in the way you see colors
  - sensitivity to light (photophobia)
- **serious heart problems.** VFEND may cause changes in your heart rate or rhythm, including your heart stopping (cardiac arrest).

- **allergic reactions.** Symptoms of an allergic reaction may include:
  - fever
  - sweating
  - feels like your heart is beating fast (tachycardia)
  - chest tightness
  - trouble breathing
  - feel faint
  - nausea
  - itching
  - skin rash
- **kidney problems.** VFEND may cause new or worse problems with kidney function, including kidney failure. Your healthcare provider should check your kidney function while you are taking VFEND. Your healthcare provider will decide if you can keep taking VFEND
- **serious skin reactions.** Symptoms of serious skin reactions may include:
  - rash or hives
  - mouth sores
  - blistering or peeling of your skin
  - trouble swallowing or breathing

Call your healthcare provider or go to the nearest hospital emergency room right away if you have any of the symptoms listed above.

**The most common side effects of VFEND include:**

- vision changes
- rash
- vomiting
- nausea
- headache
- fast heart beat (tachycardia)
- hallucinations (seeing or hearing things that are not there)
- abnormal liver function tests

Tell your healthcare provider if you have any side effect that bothers you or that does not go away.

These are not all the possible side effects of VFEND. For more information, ask your healthcare provider 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 VFEND?**

- Store VFEND tablets and liquid at room temperature, 59° to 86° F (15° to 30°C). Do not refrigerate or freeze.
- VFEND suspension should be thrown away (discarded) after 14 days.
- Keep VFEND tablets and oral suspension in a tightly closed container.
- Safely throw away medicine that is out of date or no longer needed.
- **Keep VFEND, as well as all other medicines, out of the reach of children.**

**General information about the safe and effective use of VFEND**

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use VFEND for a condition for which it was not prescribed. Do not give VFEND to other people, even if they have the same symptoms that you have. It may harm them.

This Patient Information leaflet summarizes the most important information about VFEND. If you would like more information, talk to your healthcare provider. You can ask your healthcare provider or pharmacist for information about VFEND that is written for health professionals.

This product's label may have been updated. For current full prescribing information, please visit [www.pfizer.com](http://www.pfizer.com).

**What are the ingredients of VFEND?**

Active ingredient: voriconazole

Inactive ingredients:

**VFEND IV:** sulfobutyl ether beta-cyclodextrin sodium

**VFEND tablets:** lactose monohydrate, pregelatinized starch, croscarmellose sodium, povidone, magnesium stearate, and a coating containing hypromellose, titanium dioxide, lactose monohydrate, and triacetin

**VFEND oral suspension:** colloidal silicon dioxide, titanium dioxide, xanthan gum, sodium citrate dihydrate, sodium benzoate, anhydrous citric acid, natural orange flavor, and sucrose

This Patient Information has been approved by the U.S. Food and Drug Administration.

This product's label may have been updated. For current full prescribing information, please visit [www.pfizer.com](http://www.pfizer.com).



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