PRODUCT MONOGRAPH

[Pr]PREZISTA®

darunavir tablets 75 mg, 150 mg, 400 mg, 600 mg, 800 mg
darunavir oral suspension 100 mg/mL

(as darunavir ethanolate)

Human Immunodeficiency Virus (HIV) Protease Inhibitor

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PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Pharmaceutical Form/Strength	Clinically Relevant Nonmedicinal Ingredients*
Oral	suspension, 100 mg/mL	None
Oral	tablet, 75 mg	None
Oral	tablet, 150 mg	None
Oral	tablet, 400 mg	None
Oral	tablet, 600 mg	None
Oral	tablet, 800 mg	None

For a complete listing see* **DOSAGE FORMS, COMPOSITION AND PACKAGING *section*

INDICATIONS AND CLINICAL USE

PREZISTA® (darunavir), co-administered with 100 mg ritonavir, and with other antiretroviral agents, is indicated for the treatment of HIV-1 infection.

For a description of the clinical data in support of this indication, refer to ***Product Monograph, Part II*: CLINICAL TRIALS**.

Pediatrics (from 3 to < 18 years of age)
PREZISTA® co-administered with low-dose ritonavir, and with other antiretroviral agents, is indicated for the treatment of HIV-1 infection in treatment-experienced pediatric patients 3 years of age and above (see **WARNINGS AND PRECAUTIONS** and **DOSAGE AND ADMINISTRATION**). This indication is based on 24-week and 48-week analyses of plasma HIV-1 RNA levels and CD4+ cell counts from two open-label Phase 2 trials in antiretroviral treatment-experienced pediatric patients 6 to less than 18 years of age and 3 to less than 6 years of age, respectively.

Geriatrics (> 65 years of age)
Clinical studies of PREZISTA® did not include sufficient numbers of patients aged 65 and over to determine whether they respond differently from younger patients. In general, caution should be exercised in the administration and monitoring of PREZISTA® in elderly patients, reflecting

the greater frequency of decreased hepatic, renal or cardiac function and of concomitant disease or other drug therapy (see **WARNINGS AND PRECAUTIONS**, **DOSAGE AND ADMINISTRATION,** and **ACTION AND CLINICAL PHARMACOLOGY**).

CONTRAINDICATIONS

PREZISTA® is contraindicated in patients who are hypersensitive to this drug or to any ingredient in the formulation or component of the container. For a complete listing, see the **DOSAGE FORMS, COMPOSITION AND PACKAGING** section of the Product Monograph.

PREZISTA® is contraindicated in patients with severe (Child-Pugh Class C) hepatic insufficiency.

Co-administration of PREZISTA®/rtv is contraindicated with drugs that are highly dependent on CYP3A4 for clearance and for which elevated plasma concentrations are associated with serious and/or life-threatening events (narrow therapeutic index). Co-administration of PREZISTA®/rtv is contraindicated with rifampin and St. John's Wort as it may reduce plasma concentrations of darunavir which may result in loss of therapeutic effect and development of resistance. These drugs are listed in Table 1 (also see **DRUG INTERACTIONS**, **Drug-Drug Interactions**, Table 8 and Table 9).

Table 1: Drugs that are Contraindicated with PREZISTA®/rtv

Drug Class	Drugs within Class that are Contraindicated with PREZISTA®/rtv
Alpha 1-Adrenoreceptor Antagonist	alfuzosin
Antiarrhythmics	amiodarone, bepridil[a], dronedarone, lidocaine (systemic), quinidine
Anti-coagulants	apixaban, rivaroxaban
Anti-gout	colchicine (in patients with renal and/or hepatic impairment)
Antihistamines	astemizole[a], terfenadine[a]
Antimycobacterial	rifampin
Ergot Derivatives	dihydroergotamine, ergonovine, ergotamine, methylergonovine
GI Motility Agents	cisapride[a]
Herbal Products	St. John's wort (*Hypericum perforatum*)
HMG-CoA Reductase Inhibitors	lovastatin, simvastatin
Neuroleptics	pimozide
PDE-5 Inhibitor	sildenafil (for treatment of pulmonary arterial hypertension)
Sedatives/Hypnotics	orally administered midazolam, triazolam

[a] Not marketed in Canada.

WARNINGS AND PRECAUTIONS

General

PREZISTA® (darunavir) must be administered with low-dose ritonavir to ensure its therapeutic effect (see **DETAILED PHARMACOLOGY, Pharmacokinetics, Drug-Drug Interactions**, Table 29**; DOSAGE AND ADMINISTRATION;** and **ACTION AND CLINICAL**

PHARMACOLOGY, **Pharmacokinetics**). Failure to correctly co-administer PREZISTA® with ritonavir will result in reduced plasma levels of PREZISTA® that may be insufficient to achieve the desired antiviral effect. Patients should be instructed accordingly. Please refer to the ritonavir Product Monograph for additional information on precautionary measures.

PREZISTA® is not a cure for HIV-1 infection or AIDS. Patients receiving darunavir or any other antiretroviral therapy may continue to develop opportunistic infections and other complications of HIV-1 infection.

PREZISTA® therapy has not been shown to reduce the risk of transmission of HIV-1 to others.

Due to inhibition of CYP3A by PREZISTA®, co-administration of PREZISTA® with quetiapine may results in increased quetiapine concentrations. Serious and life-threatening quetiapine-related adverse reactions have been reported with CYP3A inhibitors. PREZISTA® should not be used in combination with quetiapine (see **DRUG INTERACTIONS**). Monitoring and dose reductions may be required if necessary.

Carcinogenesis and Mutagenesis

Darunavir was evaluated for carcinogenic potential by oral gavage administration to mice and rats up to 104 weeks. A dose-related increase in the incidence of hepatocellular adenomas and carcinomas were observed in males and females of both species as well as an increase in thyroid follicular cell adenomas in male rats. These findings are considered to be of limited relevance to humans. Based on AUC measurements, exposure to darunavir at the dose levels studied was below or approximately equivalent to exposure in humans at the recommended therapeutic dose (see **TOXICOLOGY, Carcinogenesis and Mutagenesis**).

Darunavir was not mutagenic or genotoxic in a battery of *in vitro* and *in vivo* assays including bacterial reverse mutation (Ames), chromosomal aberration in human lymphocytes and *in vivo* micronucleus test in mice (see **TOXICOLOGY, Carcinogenesis and Mutagenesis**).

Endocrine and Metabolism

Diabetes Mellitus/Hyperglycemia
New onset diabetes mellitus, exacerbation of pre-existing diabetes mellitus, and hyperglycemia have been reported during postmarketing surveillance in HIV-infected patients receiving protease inhibitor (PI) therapy. Some patients required either initiation or dose adjustments of insulin or oral hypoglycemic agents for treatment of these events. In some cases, diabetic ketoacidosis has occurred. In those patients who discontinued PI therapy, hyperglycemia persisted in some cases. Because these events have been reported voluntarily during clinical practice, estimates of frequency cannot be made and causal relationships between PI therapy and these events have not been established.

Fat Distribution
Redistribution/accumulation of body fat, including central obesity, dorsocervical fat enlargement (buffalo hump), peripheral wasting, facial wasting, breast enlargement, and "cushingoid

appearance" have been observed in patients receiving antiretroviral therapy. The mechanism and long-term consequences of these events are currently unknown. A causal relationship has not been established.

Hematologic

There have been reports of increased bleeding, including spontaneous skin hematomas and hemarthrosis, in patients with hemophilia type A and B treated with protease inhibitors. In some patients, additional factor VIII was given. In more than half of the reported cases, treatment with protease inhibitors was continued or reintroduced. A causal relationship between protease inhibitor therapy and these events has not been established; however, the frequency of bleeding episodes should be closely monitored in patients on PREZISTA®/rtv.

Hepatic/Biliary/Pancreatic

Hepatic Impairment
PREZISTA® is contraindicated in patients with severe hepatic insufficiency (Child-Pugh Class C) (see **CONTRAINDICATIONS**). Patients with mild or moderate hepatic impairment (Child-Pugh Class A or B, respectively) should be closely monitored.

Patients with chronic hepatitis B or C and treated with combination antiretroviral therapy are at an increased risk for severe and potentially fatal hepatic adverse events. Limited data are currently available for the use of PREZISTA® co-administered with low dose ritonavir, in patients co-infected with hepatitis B or C.

Hepatotoxicity
Drug-induced hepatitis (e.g., acute hepatitis, cytolytic hepatitis) has been reported with PREZISTA®/rtv. During the clinical development program (n=3,063), hepatitis has been reported in 0.5% of patients receiving combination therapy with PREZISTA®/rtv.

Post-marketing cases of clinical hepatitis and hepatic decompensation, including some fatalities have been reported. These have generally occurred in patients with advanced HIV disease taking multiple concomitant medications, having co-morbidities including hepatitis B or C co-infection, and/or developing immune reconstitution inflammatory syndrome. A causal relationship with PREZISTA®/rtv therapy has not been established.

Patients with pre-existing liver dysfunction including chronic hepatitis B or C have an increased frequency of liver function abnormalities during combination antiretroviral therapy. Appropriate monitoring should be conducted prior to initiating therapy with PREZISTA®/rtv and increased monitoring should be considered in patients with elevated baseline transaminase levels, active hepatitis B or C and in patients with underlying liver disease, especially during the first several months of PREZISTA®/rtv treatment. Evidence of new or worsening liver dysfunction (including clinically significant elevation of liver enzymes and/or symptoms such as fatigue, anorexia, nausea, jaundice, dark urine, liver tenderness and hepatomegaly) in patients on PREZISTA®/rtv, should prompt consideration to interrupt or discontinue treatment.

For information on the multi-dose pharmacokinetics of darunavir in hepatically impaired patients, see **ACTION AND CLINICAL PHARMACOLOGY**.

Pancreatic
Pancreatitis has been observed in patients receiving PREZISTA®/rtv therapy, including those who developed marked triglyceride elevations. Although a causal relationship to PREZISTA® has not been established, marked triglyceride elevation is a risk factor for development of pancreatitis (see **WARNINGS AND PRECAUTIONS, Lipid Elevations**). Patients with advanced HIV disease may be at risk of elevated triglycerides and pancreatitis, and patients with a history of pancreatitis may be at increased risk for recurrence during PREZISTA®/rtv therapy.

Immune

Immune Reconstitution Inflammatory Syndrome
During the initial phase of treatment, patients responding to antiretroviral therapy may develop an inflammatory response to indolent or residual opportunistic infections (such as MAC, CMV, *Pneumocystis jirovecii* pneumonia, and TB), which may necessitate further evaluation and treatment.

Autoimmune disorders (such as Graves' disease, polymyositis and Guillain-Barré syndrome) have also been reported to occur in the setting of immune reconstitution; however, the time to onset is more variable, and can occur many months after initiation of treatment.

Lipid Elevations

Treatment with PREZISTA® has resulted in increases in the concentration of total cholesterol and triglycerides. Triglyceride and cholesterol testing should be performed prior to initiating PREZISTA® therapy and at periodic intervals during therapy. Lipid disorders should be managed as clinically appropriate. See Table 8 and Table 9 for additional information on potential drug interactions with PREZISTA® and HMG-CoA reductase inhibitors.

Renal

Renal Impairment
Population pharmacokinetic analysis showed that the pharmacokinetics of darunavir were not significantly affected in HIV-1-infected patients with moderate renal impairment (CrCL between 30–60 mL/min, n=20). There are no pharmacokinetic data available in HIV-1-infected patients with severe renal impairment or end stage renal disease. However, since the renal clearance of darunavir is limited, a decrease in total body clearance is not expected in patients with renal impairment. As darunavir and ritonavir are highly bound to plasma proteins, it is unlikely that they will be significantly removed by hemodialysis or peritoneal dialysis (see **DOSAGE AND ADMINISTRATION** and **ACTION AND CLINICAL PHARMACOLOGY**, **Special Populations and Conditions**, **Renal Insufficiency**).

Sensitivity

Darunavir contains a sulfonamide moiety. PREZISTA® (darunavir) should be used with caution in patients with a known sulfonamide allergy. The potential for cross-sensitivity between drugs in the sulfonamide class and darunavir is unknown. In clinical studies with darunavir/ritonavir, the incidence and severity of rash was similar in patients with or without a history of sulphonamide allergy.

Severe Skin Reactions

During the clinical development program (n=3,063), severe skin reactions, which may be accompanied by fever and/or elevations of transaminases, have been reported in 0.4% of patients. Stevens-Johnson Syndrome was rarely (< 0.1%) reported; and during post-marketing experience toxic epidermal necrolysis, DRESS (Drug Rash with Eosinophilia and Systemic Symptoms) and acute generalized exanthematous pustulosis have been reported very rarely (< 0.01%).

Discontinue PREZISTA® immediately if signs or symptoms of severe skin reactions develop. These can include but are not limited to severe rash or rash accompanied with fever, general malaise, fatigue, muscle or joint aches, blisters, oral lesions, conjunctivitis, hepatitis and/or eosinophilia.

Rash (all grades, regardless of causality) occurred in 10.3% of patients treated with PREZISTA® (see **ADVERSE REACTIONS**). Rash was mostly mild-to-moderate, often occurring within the first four weeks of treatment and resolving with continued dosing. The discontinuation rate due to rash in patients using PREZISTA®/rtv was 0.5%.

Darunavir contains a sulfonamide moiety. PREZISTA® should be used with caution in patients with a known sulfonamide allergy. In clinical studies with PREZISTA®/rtv, the incidence and severity of rash was similar in patients with or without a history of sulfonamide allergy.

Rash occurred more commonly in treatment-experienced patients receiving regimens containing PREZISTA®/rtv + raltegravir compared to subjects receiving PREZISTA®/rtv without raltegravir or raltegravir without PREZISTA®/rtv. However, rash that was considered drug related occurred at similar rates for all three groups. These rashes were mild to moderate in severity and did not limit therapy; there were no discontinuations due to rash.

Special Populations

Pregnant Women

PREZISTA® should not be used during pregnancy unless the potential benefit justifies the potential risk.

There are no adequate and well-controlled studies with darunavir in pregnant women.

Darunavir/ritonavir (600/100 mg twice daily. or 800/100 mg once daily) in combination with a background regimen was evaluated in a clinical trial of 34 pregnant women during the second and third trimesters, and postpartum. The pharmacokinetic data demonstrate that exposure to darunavir and ritonavir as part of an antiretroviral regimen was lower during pregnancy compared with postpartum (6-12 weeks). Virologic response was preserved throughout the study period in both arms. No mother to child transmission occurred in the infants born to the 29 subjects who stayed on the antiretroviral treatment through delivery. Darunavir/ritonavir was well tolerated during pregnancy and postpartum. There were no new clinically relevant safety findings compared with the known safety profile of darunavir/ritonavir in HIV-1 infected adults (see **ACTION AND CLINICAL PHARMACOLOGY**, **Pharmacokinetics**).

Studies in animals have not shown evidence of developmental toxicity or effect on reproductive function and fertility. However, due to limited bioavailability and/or dosing limitations, animal exposures (based on AUC) were only 50% (mice and rats) and 5% (rabbit) of those obtained in humans at the recommended clinical dose boosted with ritonavir (see **TOXICOLOGY**, **Reproductive and Developmental Toxicity**).

The Antiretroviral Pregnancy Registry has received prospective reports of exposure to darunavir-containing regimens during pregnancies resulting in live births. The majority had first exposure in the first trimester. No safety concern relating to darunavir has been identified to date.

Antiretroviral Pregnancy Registry: *To monitor maternal-fetal outcomes of pregnant women exposed to PREZISTA®, an Antiretroviral Pregnancy Registry has been established. Physicians are encouraged to register patients by calling 1-800-258-4263.*

Nursing Women

HIV-infected mothers should not breast-feed their infants to avoid risking postnatal transmission of HIV. It is not known whether darunavir is excreted in human milk. Studies in rats have demonstrated that darunavir is excreted into the milk of lactating rats and at high levels (1,000 mg/kg/day) resulted in toxicity. Because of both the potential for HIV transmission and the potential for serious adverse events in nursing infants, mothers should be instructed not to breast-feed if they are receiving PREZISTA® (see **TOXICOLOGY**, **Reproductive and Developmental Toxicity**).

Pediatrics (< 3 years of age)

The pharmacokinetics, safety, tolerability and efficacy of PREZISTA® in pediatric patients < 3 years of age have not been established. PREZISTA® should not be used in pediatric patients below 3 years of age in view of toxicity and mortality observed in juvenile rats dosed with darunavir (from 20 mg/kg to 1,000 mg/kg) up to days 23 to 26 of age (see **ACTION AND CLINICAL PHARMACOLOGY, Special Populations and Conditions, Pediatrics** and **TOXICOLOGY, Reproductive and Developmental Toxicity**).

Treatment-Naïve Pediatric Patients

The pharmacokinetics, safety, tolerability and efficacy of PREZISTA® in antiretroviral treatment-naïve pediatric patients have not been established.

Geriatrics (> 65 years of age)
Clinical studies of PREZISTA® did not include sufficient numbers of patients aged 65 and over to determine whether they respond differently from younger patients. In general, caution should be exercised in the administration and monitoring of PREZISTA® in elderly patients, reflecting the greater frequency of decreased hepatic, renal or cardiac function and of concomitant disease or other drug therapy.

ADVERSE REACTIONS

Adverse Drug Reaction Overview

During the clinical development program (n=3,063), 65.9% of patients experienced at least one ADR. The total mean exposure for patients was 57.5 weeks. The most common ADRs were diarrhea (23.7%), nausea (14.9%), headache (13.8%), and rash (10.3%). The majority of ADRs were mild or moderate in severity. The overall incidence of any Grade 3 or 4 ADR was 15.1%. The most common Grade 3 or 4 ADRs were diarrhea (1.3%) and those related to laboratory abnormalities, i.e., hepatic enzyme increased (3.4%), hypertriglyceridemia (2.9%), pancreatic enzyme increased (2.7%), and hypercholesterolemia (1.4%). Treatment discontinuation due to ADRs was infrequent (2.5%). The most common ADRs leading to treatment discontinuation were rash (0.5%), hepatic enzyme increased (0.6%), diarrhea (0.3%), and nausea (0.3%).

Clinical Trial Adverse Drug Reactions

Because clinical trials are conducted under very specific conditions the adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse drug reaction information from clinical trials is useful for identifying drug-related adverse events and for approximating rates.

Antiretroviral Treatment-Naïve Adult Patients
The safety assessment is based on all safety data from the randomized, controlled, open-label Phase 3 trial TMC114-C211 comparing PREZISTA®/rtv 800/100 mg q.d. versus lopinavir/ritonavir 800/200 mg per day in antiretroviral treatment-naïve HIV-1-infected adult patients. The total mean exposure in weeks for patients in the PREZISTA®/rtv 800/100 mg q.d. arm and the lopinavir/ritonavir 800/200 mg per day arm was 162.5 and 153.5, respectively.

Discontinuation due to adverse events/HIV-related events occurred in 4.7% of patients in the PREZISTA®/rtv group and in 12.7% of patients in the LPV/rtv group.

The majority of the ADRs reported during treatment with PREZISTA®/rtv 800/100 mg q.d. were mild in severity. The most common ADRs to PREZISTA®/rtv 800/100 mg q.d. (≥5%) of at least moderate intensity (≥ Grade 2) were diarrhea, headache and abdominal pain.

ADRs to PREZISTA®/rtv 800/100 mg q.d. of at least moderate intensity (≥ Grade 2) in antiretroviral treatment-naïve HIV-1-infected adult patients are presented in Table 2.

Table 2: Adverse Drug Reactions[a] of At Least Moderate Intensity (≥ Grade 2) Reported in ≥1% of Antiretroviral Treatment-Naïve HIV-1-Infected Adult Patients Who Received PREZISTA®/rtv 800/100 mg q.d.

	Randomized Study TMC114-C211 (through 192 weeks)	
System Organ Class, Preferred Term	**PREZISTA®/rtv 800/100 mg q.d. + TDF/FTC N=343**	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N=346**
Gastrointestinal Disorders		
Abdominal pain	5.8%	6.1%
Diarrhea	8.7%	15.9%
Nausea	4.1%	3.8%
Vomiting	2.0%	3.5%
Metabolism and Nutrition Disorders		
Anorexia	1.5%	0.9%
Nervous System Disorders		
Headache	6.7%	5.5%
Skin and Subcutaneous Tissue Disorders		
Pruritus	1.2%	0.9%
Rash	6.1%	6.6%
Urticaria[b]	1.2%	0.6%

N = total number of patients per treatment group TDF = tenofovir disoproxil fumarate FTC = emtricitabine

[a] Excluding laboratory abnormalities reported as ADRs [b] Adverse drug reaction also identified from post-marketing experience

Less Common Clinical Trial Adverse Drug Reactions (<1%)

Adverse drug reactions occurring in less than 1% of patients receiving PREZISTA®/rtv considered at least possibly related to treatment and of at least moderate intensity are listed below by body system:

Gastrointestinal Disorders: acute pancreatitis, dyspepsia, flatulence

General Disorders and Administration Site Conditions: asthenia, fatigue

Hepatobiliary Disorders: acute hepatitis

Immune System Disorders: (drug) hypersensitivity†, immune reconstitution inflammatory syndrome

Metabolism and Nutrition Disorders: diabetes mellitus

Musculoskeletal and Connective Tissue Disorders: myalgia, osteonecrosis

Psychiatric Disorders: abnormal dreams

Skin and Subcutaneous Tissue Disorders: angioedema†, lipodystrophy (lipohypertrophy, lipodystrophy and lipoatrophy), pruritus, Stevens-Johnson Syndrome, urticaria†

†Adverse drug reaction also identified from post-marketing experience

Abnormal Hematologic and Clinical Chemistry Findings

The percentages of antiretroviral treatment-naïve HIV-1-infected adult patients treated with PREZISTA®/rtv 800/100 mg q.d. with Grade 2 to 4 laboratory abnormalities, considered ADRs, are presented in Table 3.

Table 3: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Naïve HIV-1-Infected Adult Patients

		Randomized Study TMC114-C211 (through 192 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 800/100 mg q.d. + TDF/FTC N=343**	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N=346**
Biochemistry			
Alanine Aminotransferase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	8.8%	9.4%
Grade 3	> 5.0 to ≤ 10.0 X ULN	2.9%	3.5%
Grade 4	> 10.0 X ULN	0.9%	2.9%
Aspartate Aminotransferase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	7.3%	9.9%
Grade 3	> 5.0 to ≤ 10.0 X ULN	4.4%	2.3%
Grade 4	> 10.0 X ULN	1.2%	2.6%
Alkaline Phosphatase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	1.5%	1.5%
Grade 3	> 5.0 to ≤ 10.0 X ULN	0%	0.6%
Grade 4	> 10.0 X ULN	0%	0%
Hyperbilirubinemia			
Grade 2	>1.5 to ≤ 2.5 X ULN	0.9%	4.4%
Grade 3	> 2.5 to ≤ 5.0 X ULN	0.3%	0.6%
Grade 4	> 5.0 X ULN	0%	0%
Triglycerides			
Grade 2	5.65–8.48 mmol/L 500–750 mg/dL	2.6%	9.9%
Grade 3	8.49–13.56 mmol/L 751–1200 mg/dL	1.8%	5.0%
Grade 4	> 13.56 mmol/L > 1200 mg/dL	1.5%	1.2%
Total Cholesterol[a]			
Grade 2	6.20–7.77 mmol/L 240–300 mg/dL	22.9%	27.1%
Grade 3	> 7.77 mmol/L > 300 mg/dL	1.5%	5.5%
Low-Density Lipoprotein[a] Cholesterol			
Grade 2	4.13–4.90 mmol/L 160–190 mg/dL	14.1%	12.3%
Grade 3	≥ 4.91 mmol/L ≥ 191 mg/dL	8.8%	6.1%

Table 3: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Naïve HIV-1-Infected Adult Patients			
		Randomized Study TMC114-C211 (through 192 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 800/100 mg q.d. + TDF/FTC N=343**	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N=346**
Elevated Glucose Levels			
Grade 2	6.95–13.88 mmol/L 126–250 mg/dL	10.8%	9.6%
Grade 3	13.89–27.75 mmol/L 251–500 mg/dL	1.2%	0.3%
Grade 4	> 27.75 mmol/L > 500 mg/dL	0%	0%
Pancreatic Lipase			
Grade 2	> 1.5 to ≤ 3.0 X ULN	2.6%	1.7%
Grade 3	> 3.0 to ≤ 5.0 X ULN	0.6%	1.2%
Grade 4	> 5.0 X ULN	0%	0.9%
Pancreatic Amylase			
Grade 2	> 1.5 to ≤ 2.0 X ULN	4.7%	2.3%
Grade 3	> 2.0 to ≤ 5.0 X ULN	4.7%	4.1%
Grade 4	> 5.0 X ULN	0%	0.9%

N = total number of patients per treatment group
TDF = tenofovir disoproxil fumarate
FTC = emtricitabine
[a] Grade 4 data not applicable in Division of AIDS grading scale.

Antiretroviral Treatment-Experienced Adult Patients

The safety assessment is based on all safety data from the randomized, controlled, open-label Phase 3 trial TMC114-C214 comparing PREZISTA®/rtv 600/100 mg b.i.d. versus lopinavir/ritonavir 400/100 mg b.i.d. in antiretroviral treatment-experienced HIV-1-infected adult patients. The total mean exposure in weeks for patients in the PREZISTA®/rtv 600/100 mg b.i.d. arm and the lopinavir/ritonavir 400/100 mg b.i.d. arm was 80.7 and 76.4, respectively.

Discontinuation due to adverse events/HIV-related events occurred in 7.7% of patients in the PREZISTA®/rtv group and in 8.1% of patients in the LPV/rtv group.

The majority of the ADRs reported during treatment with PREZISTA®/rtv 600/100 mg b.i.d. were mild in severity. The most common ADRs to PREZISTA®/rtv 600/100 mg b.i.d. (≥5%) of at least moderate intensity (≥ Grade 2) were diarrhea, hypertriglyceridemia, hypercholesterolemia, nausea, abdominal pain, vomiting, lipodystrophy, hepatic enzymes increased and rash.

ADRs of at least moderate intensity (≥ Grade 2) and reported in ≥1% of patients treated with PREZISTA®/rtv 600/100 mg b.i.d. in antiretroviral treatment-experienced HIV-1-infected adult patients are presented in Table 4.

Table 4: Adverse Drug Reactions[a] of At Least Moderate Intensity (≥ Grade 2) Reported in ≥1% of Antiretroviral Treatment-Experienced HIV-1-Infected Adult Patients Who Received PREZISTA®/rtv 600/100 mg b.i.d.		
	Randomized Study TMC114-C214 (through 96 weeks)	
System Organ Class, Preferred Term	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR N=298**	**lopinavir/rtv 400/100 mg b.i.d. + OBR N=297**
Gastrointestinal Disorders		
Abdominal distension	2.0%	0.3%
Abdominal pain	5.7%	2.7%
Diarrhea	14.4%	19.9%
Dyspepsia	2.0%	1.0%
Nausea	7.0%	6.4%
Vomiting	5.4%	2.7%
General Disorders and Administration Site Conditions		
Asthenia	3.0%	1.0%
Fatigue	1.3%	1.3%
Metabolism and Nutrition Disorders		
Anorexia	1.7%	2.0%
Diabetes mellitus	1.7%	0.3%
Musculoskeletal and Connective Tissue Disorders		
Myalgia	1.0%	0.7%
Nervous System Disorders		
Headache	2.7%	3.0%
Skin and Subcutaneous Tissue Disorders		
Lipodystrophy (lipohypertrophy, lipodystrophy and lipoatrophy)	5.4%	4.4%
Pruritus	1.0%	1.0%
Rash	7.0%	3.0%

N = total number of patients per treatment group
OBR = optimized background regimen
[a] Excluding laboratory abnormalities reported as ADRs

Less Common Clinical Trial Adverse Drug Reactions (<1%)

Adverse drug reactions occurring in less than 1% of patients receiving PREZISTA®/rtv, considered at least possibly related to treatment and of at least moderate intensity are listed below by body system:

Gastrointestinal Disorders: acute pancreatitis, flatulence

Immune System Disorders: immune reconstitution inflammatory syndrome

Psychiatric Disorders: abnormal dreams

Reproductive and Breast Disorders: gynecomastia

Skin and Subcutaneous Tissue Disorders: urticaria†

†Adverse drug reaction also identified from post-marketing experience

Additional safety data was obtained from the randomized, controlled, open-label trial TMC114-C229 comparing PREZISTA®/rtv 800/100 mg q.d. to PREZISTA®/rtv 600/100 mg b.i.d. in treatment-experienced HIV-1-infected patients with screening genotype resistance test showing

no darunavir resistance associated mutations. ADRs of at least moderate intensity (≥ Grade 2) and reported in ≥1% of patients treated with PREZISTA® are presented in Table 5.

Table 5: Adverse Drug Reactions[a] of At Least Moderate Intensity (≥ Grade 2) Reported in ≥1% of Antiretroviral Treatment-Experienced HIV-1 Infected Adult Patients with No DRV-RAMs Who Received PREZISTA®/rtv 800/100 mg Once Daily and PREZISTA®/rtv 600/100 b.i.d.

	Randomized Study TMC114-C229 Week 48	
System Organ Class, Preferred Term	**PREZISTA®/rtv 800/100 mg once daily + OBR[b] N=294**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR[b] N=296**
Gastrointestinal Disorders		
Abdominal pain	3.1%	2.4%
Diarrhea	5.8%	5.4%
Dyspepsia	0.3%	1.4%
Nausea	4.8%	5.1%
Vomiting	3.4%	5.4%
Metabolism and Nutrition Disorders		
Anorexia	0.3%	1.4%
Diabetes mellitus	0.3%	1.0%
Musculoskeletal and Connective Tissue Disorders		
Myalgia	0.7%	1.4%
Nervous System Disorders		
Headache	3.4%	4.4%
Skin and Subcutaneous Tissue Disorders		
Rash	2.0%	0.3%

N = total number of patients per treatment group
[a] Excluding laboratory abnormalities reported as ADRs
[b] OBR = optimized background regimen

Less Common Clinical Trial Adverse Drug Reactions (<1%)

Adverse drug reactions occurring in less than 1% of patients receiving PREZISTA®/rtv considered at least possibly related to treatment and of at least moderate intensity are listed below by body system:

Gastrointestinal Disorders: abdominal distention, flatulence

General Disorders and Administration Site Conditions: asthenia, fatigue

Hepatobiliary Disorders: hepatitis acute

Skin and Subcutaneous Tissue Disorders: angioedema, lipodystrophy, pruritus, urticaria†

†Adverse drug reaction also identified from post-marketing experience

Abnormal Hematologic and Clinical Chemistry Findings

The percentages of antiretroviral treatment-experienced HIV-1-infected adult patients treated with PREZISTA®/rtv 600/100 mg b.i.d. with Grade 2 to 4 laboratory abnormalities, considered ADRs, are presented in Table 6.

Table 6: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Experienced HIV-1-Infected Adult Patients			
		Randomized Study TMC114-C214 (through 96 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR[a] N = 298**	**lopinavir/rtv 400/100 mg b.i.d. + OBR N = 297**
Biochemistry			
Alanine Aminotransferase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	6.9%	4.8%
Grade 3	> 5.0 to ≤ 10.0 X ULN	2.4%	2.4%
Grade 4	> 10.0 X ULN	1.0%	1.7%
Aspartate Aminotransferase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	5.5%	6.2%
Grade 3	> 5.0 to ≤ 10.0 X ULN	2.4%	1.7%
Grade 4	> 10.0 X ULN	0.7%	1.7%
Alkaline Phosphatase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	0.3%	0%
Grade 3	> 5.0 to ≤ 10.0 X ULN	0.3%	0.3%
Grade 4	> 10.0 X ULN	0%	0%
Hyperbilirubinemia			
Grade 2	>1.5 to ≤ 2.5 X ULN	0.3%	1.7%
Grade 3	> 2.5 to ≤ 5.0 X ULN	0.3%	0.3%
Grade 4	> 5.0 X ULN	0.3%	0%
Triglycerides			
Grade 2	5.65–8.48 mmol/L 500–750 mg/dL	10.4%	11.4%
Grade 3	8.49–13.56 mmol/L 751–1200 mg/dL	6.9%	9.7%
Grade 4	> 13.56 mmol/L > 1200 mg/dL	3.1%	6.2%
Total Cholesterol[b]			
Grade 2	6.20–7.77 mmol/L 240–300 mg/dL	24.9%	23.2%
Grade 3	> 7.77 mmol/L > 300 mg/dL	9.7%	13.5%
Low-Density Lipoprotein[b] Cholesterol			
Grade 2	4.13–4.90 mmol/L 160–190 mg/dL	14.4%	13.5%
Grade 3	≥ 4.91 mmol/L ≥ 191 mg/dL	7.7%	9.3%
Elevated Glucose Levels			
Grade 2	6.95–13.88 mmol/L 126–250 mg/dL	10.0%	11.4%
Grade 3	13.89–27.75 mmol/L 251–500 mg/dL	1.4%	0.3%
Grade 4	> 27.75 mmol/L > 500 mg/dL	0.3%	0%
Pancreatic Lipase			

Table 6: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Experienced HIV-1-Infected Adult Patients			
		Randomized Study TMC114-C214 (through 96 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR[a] N = 298**	**lopinavir/rtv 400/100 mg b.i.d. + OBR N = 297**
Grade 2	> 1.5 to ≤ 3.0 X ULN	2.8%	3.5%
Grade 3	> 3.0 to ≤ 5.0 X ULN	2.1%	0.3%
Grade 4	> 5.0 X ULN	0.3%	0%
Pancreatic Amylase			
Grade 2	> 1.5 to ≤ 2.0 X ULN	6.2%	7.3%
Grade 3	> 2.0 to ≤ 5.0 X ULN	6.6%	2.8%
Grade 4	> 5.0 X ULN	0%	0%
N = total number of patients per treatment group [a] OBR = optimized background regimen [b] Grade 4 data not applicable in Division of AIDS grading scale.			

Except for a lower incidence of diarrhea and Grade 3 or 4 increases in triglycerides, and a higher incidence of rash-related AEs with DRV/rtv than with LPV/rtv, there were no clinically relevant differences in the overall safety profile for DRV/rtv 600/100 mg b.i.d. and LPV/rtv 400/100 mg b.i.d in the TMC114-C214 study.

The percentages of antiretroviral treatment-experienced HIV-1-infected adult patients with no darunavir resistance-associated mutations treated with PREZISTA®/rtv 800/100 mg once daily and PREZISTA®/rtv 600/100 b.i.d. with Grade 2 to 4 laboratory abnormalities, considered ADRs, are presented in Table 7.

Table 7: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Experienced HIV-1-Infected Adult Patients with No RAMS			
		Randomized Study TMC114-C229 (48 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 800/100 mg once daily + OBR[a] N = 294**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR[a] N = 296**
Biochemistry			
Alanine Transaminase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	1.7%	2.5%
Grade 3	> 5.0 to ≤ 10.0 X ULN	0%	0.7%
Grade 4	> 10.0 X ULN	0%	0.4%
Aspartate Aminotransferase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	1.4%	2.5%
Grade 3	> 5.0 to ≤ 10.0 X ULN	0.7%	0.7%
Grade 4	> 10.0 X ULN	0%	0.4%
Alkaline Phosphatase			
Grade 2	> 2.5 to ≤ 5.0 X ULN	0.7%	0.4%

Table 7: Grade 2 to 4 Laboratory Abnormalities Observed in Antiretroviral Treatment-Experienced HIV-1-Infected Adult Patients with No RAMS			
		Randomized Study TMC114-C229 (48 weeks)	
Laboratory Parameter Preferred Term	**Limit**	**PREZISTA®/rtv 800/100 mg once daily + OBR[a] N = 294**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR[a] N = 296**
Grade 3	> 5.0 to ≤ 10.0 X ULN	0%	0%
Grade 4	> 10.0 X ULN	0%	0%
Triglycerides			
Grade 2	5.65–8.48 mmol/L 500–750 mg/dL	3.5%	7.1%
Grade 3	8.49–13.56 mmol/L 751–1200 mg/dL	1.4%	2.8%
Grade 4	> 13.56 mmol/L > 1200 mg/dL	0.3%	1.1%
Total Cholesterol [b]			
Grade 2	6.20–7.77 mmol/L 240–300 mg/dL	7.7%	14.9%
Grade 3	> 7.77 mmol/L > 300 mg/dL	2.4%	5.7%
Low-Density Lipoprotein Cholesterol [b]			
Grade 2	4.13–4.90 mmol/L 160–190 mg/dL	7.0%	12.8%
Grade 3	≥ 4.91 mmol/L ≥ 191 mg/dL	2.8%	3.9%
Elevated Glucose Levels			
Grade 2	6.95–13.88 mmol/L 126–250 mg/dL	6.6%	5.3%
Grade 3	13.89–27.75 mmol/L 251–500 mg/dL	0.7%	0.7%
Grade 4	> 27.75 mmol/L > 500 mg/dL	0%	0.4%
Pancreatic Lipase			
Grade 2	> 1.5 to ≤ 3.0 X ULN	1.0%	1.8%
Grade 3	> 3.0 to ≤ 5.0 X ULN	0.3%	0%
Grade 4	> 5.0 X ULN	0%	0%
Pancreatic Amylase			
Grade 2	> 1.5 to ≤ 2.0 X ULN	3.1%	2.5%
Grade 3	> 2.0 to ≤ 5.0 X ULN	2.4%	1.1%
Grade 4	> 5.0 X ULN	0.3%	0.4%

N=total number of patients per treatment group
[a] OBR = optimized background regimen
[b] Grade 4 data not applicable in Division of AIDS grading scale.

Additional ADRs to PREZISTA®/rtv Identified in Adult Patients in Other Clinical Trials
The additional ADRs of interest identified from other clinical trials were osteonecrosis (0.4%).

Serious ADRs
The following serious ADRs of at least moderate intensity (≥ Grade 2) occurred in the Phase 2b studies and Phase 3 studies with PREZISTA®/rtv: abdominal pain, acute hepatitis, acute pancreatitis, anorexia, asthenia, diabetes mellitus, diarrhea, fatigue, headache, hepatic enzyme increased, hypercholesterolemia, hyperglycemia, hypertriglyceridemia, immune reconstitution inflammatory syndrome, low density lipoprotein increased, nausea, pancreatic enzyme increased, rash, Stevens-Johnson Syndrome and vomiting.

Patients Co-infected with Hepatitis B and/or Hepatitis C Virus

The incidence of adverse events or clinical chemistry abnormalities, except for increased hepatic enzymes, was comparable in patients co-infected with hepatitis B or C virus and patients who were not co-infected. Patients co-infected with hepatitis B or C virus receiving PREZISTA®/rtv were more likely to have baseline and treatment-emergent hepatic transaminase elevations than those without chronic viral hepatitis. Patients with chronic hepatitis B and/or C co-infection should be monitored appropriately.

Clinical Trial Experience in Pediatric Patients

The safety assessment in children and adolescents is based on the safety data from two Phase 2 trials: DELPHI (TMC114-C212) in which 80 antiretroviral treatment-experienced HIV-1-infected pediatric patients 6 to < 18 years of age and weighing at least 44 lbs (20 kg) received PREZISTA® tablets in combination with low dose ritonavir and other antiretroviral agents (24-week data), and ARIEL (TMC114-C228) in which 21 antiretroviral treatment-experienced HIV-1 infected pediatric patients aged from 3 to < 6 years and weighing ≥ 10 kg to < 20 kg received PREZISTA® oral suspension in combination with low dose ritonavir and other antiretroviral agents (48-week data) (see **ACTION AND CLINICAL PHARMACOLOGY, Special Populations and Conditions**, **Pediatrics** and **CLINICAL TRIALS**).

In DELPHI (TMC114-C212), the mean duration of patient exposure to DRV/rtv at the time of the data cut-off for the Week 24 analysis was 39.5 (±11.02) weeks. Total patient years of exposure was 60.8 years. Frequency, type and severity of adverse drug reactions in pediatric patients were comparable to those observed in adults. The overall incidence of ADRs was 40%. The most common ADRs (all Grades, ≥ 3%) were vomiting (12.5%), diarrhea (11.3%), abdominal pain (10%), headache (9%), rash (5%), nausea (4%), and fatigue (3%). Grade 3 or 4 laboratory abnormalities were ALT increased (Grade 3: 3%; Grade 4: 1%), AST increased (Grade 3: 1%), pancreatic amylase increased (Grade 3: 4%; Grade 4: 1%), pancreatic lipase increased (Grade 3: 1%), total cholesterol increased (Grade 3: 1%), and LDL increased (Grade 3: 3%). The majority of ADRs were mild or moderate in severity. There were no ADRs leading to treatment discontinuation in the Week 24 analysis of the TMC114-C212 trial.

In ARIEL (TMC114-C228), the mean duration of patient exposure to DRV/rtv at the time of the data cut-off for the Week 48 analysis was 47.93 (±2.350) weeks. Total patient-years of exposure was 19.4. Frequency, type and severity of adverse drug reactions in pediatric patients were comparable to those observed in adults, and in treatment-experienced children between 6 and 18 years of age. The overall incidence of ADRs was 47.9%. The most common ADRs (all Grades, ≥ 3%) were diarrhea (24%), vomiting (19%), rash (14%), abdominal pain (5%), anorexia (5%), and hyperglycemia (5%). All laboratory abnormalities, except one (Grade 3 decreased neutrophils), were Grade 1 or 2 in severity.

Post-Market Adverse Drug Reactions

In addition to adverse events identified in clinical trials, the following post-marketing events have been included due to their seriousness, frequency of reporting, potential causal association with PREZISTA®/rtv, or a combination of these factors. Because they are reported spontaneously from a population of unknown size, estimates of incidence cannot be made.

Blood and Lymphatic System Disorders: anemia, pancytopenia, thrombocytopenia and neutropenia

Cardiac Disorders: bradycardia, myocarditis

Eye Disorders: eye swelling, uveitis, maculopathy, blurred vision

Gastrointestinal Disorders: pancreatitis, pancreatitis relapsing, rectal hemorrhage, gastritis

Hepatobiliary Disorders: bile duct obstruction, hepatic cirrhosis, hepatic failure, hepatitis, hepatotoxicity, jaundice

Infections and Infestations: clostridial infection, cryptosporidiosis infection, cytomegalovirus encephalitis, hepatitis B, esophageal candidiasis, progressive multifocal leukoencephalopathy, sepsis

Investigations: blood alkaline phosphatase increased, blood bilirubin increased, abnormal liver function test

Immune System Disorder: drug hypersensitivity, immune reconstitution inflammatory syndrome, autoimmune disorders such as Graves' disease

Injury, Poisoning and Procedural Complications: drug toxicity

Metabolism and Nutrition Disorders: dehydration, hyperkalemia, metabolic acidosis

Musculoskeletal and Connective Tissue Disorders: myositis, osteonecrosis, rhabdomyolysis, sensation of heaviness, arthritis, bone pain, pain in extremities, arthropathy

Neoplasms Benign, Malignant and Unspecified: diffuse large B-cell neoplasm, malignant hepatic neoplasm, lymphoma

Nervous System Disorders: altered state of consciousness, cerebrovascular accident, dizziness, facial palsy, grand mal convulsion, ischemic cerebral infarction, nervous system disorder, neuromyopathy, petit mal epilepsy

Psychiatric Disorders: completed suicide, anxiety, depression

Renal and Urinary Disorders: acute renal failure, hematuria, renal tubular necrosis, creatinine renal decreased, GFR decreased, renal failure, proteinuria

Respiratory, Thoracic and Mediastinal Disorders: acute respiratory distress syndrome, pharyngeal lesion, pneumothorax, respiratory failure, pulmonary edema, epistaxis

Skin and Subcutaneous Tissue Disorders: angioedema, rash, swelling face, Stevens-Johnson syndrome, toxic epidermal necrolysis, urticaria, acute generalized exanthematous pustulosis, DRESS (Drug Rash with Eosinophilia and Systemic Symptoms)

DRUG INTERACTIONS

Serious Drug Interactions

- Darunavir and ritonavir are both inhibitors of the cytochrome P450 3A4 (CYP3A4) isoform. PREZISTA®/rtv should not be co-administered with medicinal products that are highly dependent on CYP3A4 for clearance, and for which increased plasma concentrations are associated with serious and/or life-threatening events (narrow therapeutic index). These medicinal products include alfuzosin, amiodarone, apixaban, astemizole, bepridil, cisapride, colchicine (in patients with renal and/or hepatic impairment), dronedarone, lidocaine (systemic), lovastatin, oral midazolam, pimozide, quinidine, rivaroxaban, sildenafil (when used for the treatment of pulmonary arterial hypertension), simvastatin, terfenadine, the ergot alkaloids (e.g., ergotamine, dihydroergotamine, ergonovine, and methylergonovine), and triazolam (see **CONTRAINDICATIONS**).
- Rifampin and St John's Wort (*Hypericum perforatum*) are potent inducers of CYP450 metabolism. PREZISTA®/rtv should not be used in combination with these products as this may cause significant decreases in darunavir plasma concentrations. This may result in a loss of therapeutic effect of PREZISTA® and development of resistance (see **CONTRAINDICATIONS**).

Overview

Darunavir and ritonavir are both inhibitors of the cytochrome P450 isoform CYP3A4 and CYP2D6 and the transporter P-gp. Co-administration of darunavir and ritonavir with drugs primarily metabolized by CYP3A4, CYP2D6, or transported by P-gp may result in increased plasma concentrations of such drugs, which could increase or prolong their therapeutic effect and adverse events (see **CONTRAINDICATIONS** and **Drug-Drug Interactions**, Tables 8 and 9).

Drug-Drug Interactions

Drugs that are contraindicated and not recommended for co-administration with PREZISTA®/ rtv are included in Table 8. These recommendations are based on either drug interaction studies or predicted interactions due to the expected magnitude of interaction and potential for serious events or loss of efficacy.

Table 8: Drugs that are CONTRAINDICATED with PREZISTA®/rtv	
Drug Class: Drug Name	**Clinical Comment**
Alpha 1–Adrenoreceptor Antagonists: alfuzosin	Due to potential for serious and/or life-threatening reactions such as hypotension.
Antiarrhythmics: bepridil[a] dronedarone lidocaine (systemic) quinidine amiodarone	Concentrations of bepridil, dronedarone, lidocaine, quinidine and amiodarone may be increased when co-administered with PREZISTA®/rtv.
Anticoagulants: apixaban rivaroxaban	Concentrations of apixaban or rivaroxaban may be increased when co-administered with PREZISTA®/rtv (inhibition of CYP3A and/or P glycoprotein).
Anti-gout colchicine	Patients with renal or hepatic impairment should not be given colchicine with PREZISTA®/rtv. Concomitant use of PREZISTA®/rtv with colchicine may increase concentrations of colchicine (inhibition of CYP3A). Refer to Table 9 for dosing recommendations.
Antihistamines: astemizole[a] terfenadine[a]	Due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
Antimycobacterials: rifampin	Rifampin is a potent inducer of CYP450 metabolism. PREZISTA®/rtv should not be used in combination with rifampin, as this may cause significant decreases in darunavir plasma concentrations. This may result in loss of therapeutic effect of PREZISTA® and development of resistance.
Ergot Derivatives: dihydroergotamine ergonovine ergotamine methylergonovine	Due to potential for serious and/or life-threatening reactions such as acute ergot toxicity characterized by peripheral vasospasm and ischemia of the extremities and other tissues.
Gastrointestinal Motility Agents: cisapride[a]	Due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
Herbal Products: St. John's wort (*Hypericum perforatum*)	PREZISTA®/rtv should not be used concomitantly with products containing St. John's wort (*Hypericum perforatum)* because co-administration may cause significant decreases in darunavir plasma concentrations. This may result in loss of therapeutic effect of PREZISTA® and development of resistance.
HMG-CoA Reductase Inhibitors: lovastatin simvastatin	HMG-CoA reductase inhibitors, such as lovastatin and simvastatin, which are highly dependent on CYP3A4 metabolism, are expected to have markedly increased plasma concentrations when co-administered with darunavir/ritonavir. Increased concentrations of HMG-CoA reductase inhibitors may cause myopathy, including rhabdomyolysis. For information regarding atorvastatin and pravastatin see Table 9.
Neuroleptics: pimozide	Due to the potential for serious and/or life-threatening reactions such as cardiac arrhythmias.

Table 8: Drugs that are CONTRAINDICATED with PREZISTA®/rtv	
Drug Class: Drug Name	**Clinical Comment**
PDE-5 Inhibitors: sildenafil (for treatment of pulmonary arterial hypertension)	A safe and effective dose of the PDE-5 inhibitors for the treatment of pulmonary arterial hypertension has not been established when co-administered with PREZISTA®/rtv. There is an increased potential for sildenafil-associated adverse events (which include visual disturbances, hypotension, prolonged erection, and syncope).
Sedatives/Hypnotics: orally administered midazolam[b] orally administered triazolam	Due to the potential for serious and/or life-threatening reactions such as prolonged or increased sedation or respiratory depression.

[a] Not marketed in Canada.

Established and other potentially significant drug interactions with PREZISTA®/rtv are included in Table 9. These recommendations are based on either drug interaction studies or predicted interactions due to the expected magnitude of interaction and potential for serious events or loss of efficacy.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
HIV-Antiviral Agents: Non-Nucleoside Reverse Transcriptase Inhibitors (NNRTIs)		
delavirdine	↑ darunavir ↑ delavirdine	Co administration of PREZISTA®/rtv and delavirdine may increase darunavir and delavirdine concentrations (inhibition of CYP3A). The appropriate doses of PREZISTA®/rtv and delavirdine have not been established. The combination of PREZISTA®/rtv and delavirdine is not recommended.
efavirenz	↓ darunavir ↑ efavirenz	An interaction trial between darunavir (300 mg twice daily [b.i.d.]), low-dose ritonavir (100 mg b.i.d.), and efavirenz (600 mg once daily [q.d.]) has been performed. In the presence of efavirenz, a decrease of 13% for darunavir exposure was observed. Exposure to efavirenz increased by 21% when administered in combination with darunavir and ritonavir. Since this difference is not considered to be clinically relevant, the combination of PREZISTA®/rtv and efavirenz can be used without dose adjustments.
etravirine	↔ darunavir ↓ etravirine	In an interaction trial between PREZISTA®/rtv (600/100 mg b.i.d.) and etravirine (100 mg b.i.d.), there was a 37% decrease in etravirine exposure in the presence of PREZISTA®/rtv and no relevant change in exposure to darunavir. Therefore, PREZISTA®/rtv can be co-administered with etravirine at the recommended therapeutic dose of 200 mg b.i.d. without dose adjustments.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
nevirapine	↔ darunavir ↑ nevirapine	The results of an interaction trial with darunavir (400 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and nevirapine (200 mg b.i.d.) demonstrated that darunavir exposure was not affected when administered concomitantly with nevirapine. Exposure to nevirapine increased by 27% (compared to historical controls) when administered in combination with darunavir and ritonavir. No dose adjustment is currently recommended for the combination of PREZISTA®/rtv and nevirapine. However, the literature indicates that changes in plasma exposure of nevirapine can lead to significant safety concerns, specifically hepatotoxicity. For further information, please refer to the nevirapine Product Monograph.
rilpivirine	↔ darunavir ↑ rilpivirine	Concomitant use of rilpivirine with darunavir/ritonavir may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). An interaction trial between rilpivirine (150 mg q.d.) and darunavir/ritonavir (800 mg/100 mg q.d.) demonstrated that darunavir/ritonavir increased the mean exposure of rilpivirine by 2.3-fold and from 2.7-fold to 3.8-fold in a subset (31%) of subjects. Caution should be exercised when these drugs are co-administered.
HIV-Antiviral Agents: Nucleoside Reverse Transcriptase Inhibitors (NRTIs)		
didanosine	↔ darunavir ↔ didanosine	PREZISTA®/rtv (600/100 mg b.i.d.) did not significantly affect didanosine exposure. The combination of PREZISTA® co-administered with 100 mg ritonavir and didanosine can be used without dose adjustments. Dosing of enteric-coated didanosine and darunavir, co-administered with low-dose ritonavir, should be separated by at least 2 hours to avoid formulation incompatibility (see **Drug-Food Interactions** and **DOSAGE AND ADMINISTRATION**, **Dosing Considerations**).
tenofovir disoproxil fumarate	↔ darunavir ↑ tenofovir	The results of an interaction trial between darunavir (300 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and tenofovir disoproxil fumarate (300 mg q.d.) demonstrated that darunavir exposure was not significantly affected when administered concomitantly with tenofovir disoproxil fumarate. Exposure to tenofovir disoproxil fumarate increased by 22% when administered in combination with darunavir and ritonavir. This finding is not considered to be clinically relevant. There was no change in the urinary excretion of tenofovir disoproxil fumarate or darunavir during co-administration. The combination of PREZISTA®/rtv and tenofovir disoproxil fumarate can be used without dose adjustments.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
HIV-Antiviral Agents: CCR5 Antagonist		
maraviroc	↔ darunavir ↑ maraviroc	An interaction trial between darunavir (600 mg b.i.d.), low dose ritonavir (100 mg b.i.d.), and maraviroc (150 mg b.i.d.) demonstrated that in the presence of PREZISTA®/rtv the exposure of maraviroc was increased by 305%. There was no apparent effect of maraviroc on darunavir/ritonavir exposure. When used in combination with PREZISTA®/rtv, the dose of maraviroc should be 150 mg twice daily.
HIV-Antiviral Agents: Integrase strand transfer Inhibitors		
dolutegravir	↔ darunavir ↔ dolutegravir	PREZISTA®/rtv (600/100 mg b.i.d.) decreased the dolutegravir AUC_{24h} and the C_{24h} by 22% and 38%, respectively, with no effect on the maximum plasma concentration (C_{max}). No impact of dolutegravir on darunavir pharmacokinetics is expected which is confirmed with historical pharmacokinetic data for darunavir. The decrease in dolutegravir pharmacokinetics is not considered to be clinically relevant and no dose adjustment is recommended.
elvitegravir	↔ darunavir	When PREZISTA®/rtv (600/100 mg b.i.d.) is used in combination with elvitegravir, the dose of elvitegravir should be 150 mg once daily. The pharmacokinetics and dosing recommendations for other doses of darunavir or with elvitegravir/cobicistat have not been established. Therefore, co administration of PREZISTA®/rtv in doses other than 600/100 mg b.i.d. and elvitegravir is not recommended. Co administration of PREZISTA®/rtv and elvitegravir in the presence of cobicistat is not recommended.
raltegravir	↓ darunavir	Some clinical studies suggest raltegravir may cause a modest decrease in darunavir plasma concentrations. At present the effect of raltegravir on darunavir plasma concentrations does not appear to be clinically relevant. PREZISTA® co-administered with low dose ritonavir and raltegravir can be used without dose adjustments.
HIV-Antiviral Agents: HIV-Protease Inhibitors		
ritonavir	↑ darunavir	The overall pharmacokinetic enhancement effect by ritonavir was an approximate 14-fold increase in the systemic exposure of darunavir when a single dose of 600 mg darunavir was given orally in combination with ritonavir at 100 mg b.i.d. Therefore, PREZISTA® should only be used in combination with 100 mg of ritonavir as a pharmacokinetic enhancer (see **WARNINGS AND PRECAUTIONS**, **General** and **ACTION AND CLINICAL PHARMACOLOGY**, **Pharmacokinetics**, **Absorption and Bioavailability**).

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
atazanavir	↔ darunavir ↔ atazanavir	An interaction trial between darunavir (400 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and atazanavir (300 mg q.d.) demonstrated that exposure to darunavir and atazanavir was not significantly affected when co-administered. Atazanavir can be co-administered with PREZISTA®/rtv.
indinavir	↑ darunavir ↑ indinavir	An interaction trial between darunavir (400 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and indinavir (800 mg b.i.d.) demonstrated that darunavir exposure was increased by 24% when co-administered with indinavir and ritonavir; indinavir exposure was increased by 23% when administered concomitantly with darunavir/ritonavir. When used in combination with PREZISTA®/rtv, dose adjustment of indinavir may be warranted in case of intolerance.
lopinavir/ritonavir	↓ darunavir	Results of interaction trials with PREZISTA® with or without ritonavir and lopinavir/ritonavir (1200 mg darunavir b.i.d. with or without 100 mg ritonavir b.i.d. and lopinavir/ritonavir 400/100 mg b.i.d. or 533/133.3 mg b.i.d.) demonstrated a decrease in the exposure (AUC) of darunavir by 40%. The appropriate doses of the combination have not been established. Hence, it is not recommended to co-administer PREZISTA®/rtv with lopinavir/ritonavir.
saquinavir	↓ darunavir	An interaction trial between darunavir (400 mg b.i.d.), saquinavir (1000 mg b.i.d.), and low-dose ritonavir (100 mg b.i.d.) demonstrated that darunavir exposure was decreased by 26% when co-administered with saquinavir and ritonavir; saquinavir exposure was not affected when administered concomitantly with darunavir/ritonavir. It is not recommended to co-administer saquinavir and PREZISTA®, with or without low-dose ritonavir.
Other Agents		
Antacids aluminium/magnesium hydroxide calcium carbonate	↔ darunavir	No interaction is expected between antacids and PREZISTA/rtv. PREZISTA/rtv and antacids can be used concomitantly without dose adjustments.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Antiarrythmics digoxin	↑ digoxin	An interaction trial with PREZISTA®/rtv (600/100 mg b.i.d.) and a single dose of digoxin (0.4 mg) showed an increase of digoxin AUC_{last} of 77% (ratio of Least Square Means was 1.77 with a 90% CI of 0.90 to 3.50). It is recommended that the lowest dose of digoxin should initially be prescribed and digoxin dose should be titrated to obtain the desired clinical effect when co-administered with PREZISTA®/rtv. Serum digoxin concentrations should be monitored to assist in the titration.
disopyramide flecainide mexiletine propafenone	↑ antiarrhythmics	Co-administration of disopyramide or propafenone with PREZISTA®/rtv is expected to increase plasma concentration of disopyramide or propafenone. Disopyramide or propafenone plasma concentration should be monitored when co-administered with PREZISTA®/rtv. Exposure to flecainide or mexiletine may be increased when co-administered with PREZISTA®/rtv. Caution is warranted and therapeutic concentration monitoring of antiarrythmics is recommended when available.
Anticancer Agents: dasatinib nilotinib vinblastine vincristine	↑ anticancer agent	The plasma concentrations of these anticancer agents are expected to increase with co administration of PREZISTA®/rtv (inhibition of CYP3A), resulting in the potential for adverse events usually associated with these agents. Caution should be exercised and drug concentration monitoring should be conducted, if available, when combining one of these anticancer agents with PREZISTA®/rtv.
everolimus		Concomitant use of everolimus and PREZISTA®/rtv is not recommended.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Anticoagulants: dabigatran etexilate	↑ anticoagulant	The combination of PREZISTA®/rtv and dabigatran etexilate should be used with caution and is not recommended in subjects with severe renal impairment.
warfarin	↓ warfarin ↔ darunavir	Warfarin concentrations may be affected when co-administered with PREZISTA®/rtv. It is recommended that the international normalized ratio (INR) be monitored when warfarin is combined with PREZISTA®/rtv.
Anticonvulsant: carbamazepine	↔ darunavir ↑ carbamazepine	An interaction trial between darunavir/rtv (600/100 mg b.i.d.) and carbamazepine (200 mg b.i.d.) showed that the exposure to darunavir, co-administered with ritonavir, was unaffected by carbamazepine. Ritonavir exposure (AUC_{12h}) was decreased by 49% . For carbamazepine, AUC_{12h} was increased by 45%. No dose adjustment for PREZISTA®/rtv is recommended. If there is a need to combine PREZISTA®/rtv and carbamazepine, patients should be monitored for potential carbamazepine-related adverse events. Carbamazepine concentrations should be monitored and its dose should be titrated for adequate response. Based upon the findings, the dose of carbamazepine may need to be reduced by 25% to 50% in the presence of PREZISTA®/rtv.
Anticonvulsants: phenobarbital phenytoin	↓ darunavir	Phenobarbital and phenytoin are inducers of CYP450 enzymes. PREZISTA®/rtv should not be used in combination with phenobarbital, or phenytoin as co-administration may cause significant decreases in darunavir plasma concentrations. This may result in loss of therapeutic effect of PREZISTA®.
Anti-bacterials: clarithromycin	↔ darunavir ↑ clarithromycin	An interaction trial between darunavir (400 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and clarithromycin (500 mg b.i.d.) demonstrated an increase in exposure to clarithromycin by 57%, while exposure to darunavir was not affected. For patients with renal impairment, the following dose adjustments should be considered: • For patients with CLcr of 30-60 mL/min, the dose of clarithromycin should be reduced by 50%. • For patients with CLcr of < 30 mL/min, the dose of clarithromycin should be reduced by 75%.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Antifungals: ketoconazole itraconazole (not studied) posaconazole voriconazole (not studied)	↑ ketoconazole ↑ darunavir itraconazole (not studied) voriconazole (not studied)	Ketoconazole, itraconazole and posaconazole are potent inhibitors as well as substrates of CYP3A4. Concomitant systemic use of these antifungals, and darunavir and ritonavir may increase plasma concentrations of both darunavir and some of these antifungals. In an interaction trial, concomitant administration of ketoconazole (200 mg b.i.d.) with darunavir (400 mg b.i.d.) and ritonavir (100 mg b.i.d.) increased exposure of ketoconazole and darunavir by 212% and 42%, respectively. When co-administration is required, the daily dose of ketoconazole or itraconazole should not exceed 200 mg. Clinical monitoring is recommended when coadministering PREZISTA®/rtv with posaconazole. Co-administration of voriconazole with PREZISTA®/rtv has not been studied. Administration of voriconazole with ritonavir (100 mg twice daily) decreased the AUC of voriconazole by an average of 39%. Voriconazole should not be administered to patients receiving darunavir/ritonavir unless an assessment of the benefit/risk ratio justifies the use of voriconazole.
Anti-gout: colchicine	↑ colchicine	Exposure of colchicines, a CYP34A substrate, may be increased when co-administered with Darunavir/rtv. <u>Treatment of gout-flares – co-administration of colchicine in patients on PREZISTA®/rtv:</u> 0.6 mg (1 tablet) x 1 dose, followed by 0.3 mg (half tablet) 1 hour later. Treatment course to be repeated no earlier than 3 days. <u>Prophylaxis of gout-flares – co-administration of colchicine in patients on PREZISTA®/rtv:</u> If the original regimen was 0.6 mg twice a day, the regimen should be adjusted to 0.3 mg (half tablet) once a day. If the original regimen was 0.6 mg once a day, the regimen should be adjusted to 0.3 mg (half tablet) once every other day. <u>Treatment of familial Mediterranean fever – co-administration of colchicine in patients on PREZISTA®/ritonavir:</u> maximum daily dose of 0.6 mg (may be given as 0.3 mg twice a day). **The use of colchicine in patients with renal or hepatic impairment is contraindicated with PREZISTA®/ritonavir.**

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Antimalarials: artemether/lumefantrine	↑ lumefantrine ↔ darunavir ↓ artemether and its active metabolite, dihydroartemisinin	Artemether/lumefantrine are not approved for use in Canada. An interaction trial between PREZISTA®/rtv (600/100 mg b.i.d.) and artemether/lumefantrine (80/480 mg, 6 doses at 0, 8, 24, 36, 48, and 60 hours) showed an increase in exposure to lumefantrine by 2.75-fold, while exposure to darunavir was not affected. The exposure to artemether and its active metabolite, dihydroartemisinin, decreased by 16% and 18%, respectively.
Antimycobacterials: rifabutin rifapentine[a]	↑ darunavir ↑ rifabutin ↑ 25-*O*-desacetylrifabutin	Rifabutin is a substrate of CYP450 enzymes. In an interaction trial, an increase of systemic exposure to darunavir by 57% was observed when PREZISTA®/rtv (600/100 mg b.i.d.) was administered with rifabutin (150 mg once every other day (q.o.d.). Based on the safety profile of PREZISTA®/rtv, the increase in darunavir exposure in the presence of rifabutin does not warrant a dose adjustment for PREZISTA®/rtv. The interaction trial showed a comparable systemic exposure for rifabutin between treatment at 300 mg q.d. alone and at 150 mg q.o.d. in combination with PREZISTA®/rtv (600/100 mg b.i.d.) with an increase in exposure to the active metabolite 25-*O*-desacetyl-rifabutin. A dosage reduction of rifabutin by 75% of the usual dose of 300 mg/day (i.e., rifabutin 150 mg every other day) is warranted if rifabutin is co-administered with PREZISTA®/rtv. Increased monitoring for rifabutin-related adverse events is warranted in patients receiving the combination. Co-administration of PREZISTA®/rtv with rifapentine may decrease darunavir concentrations (induction of CYP3A), which may result in loss of therapeutic effect of PREZISTA®. Co-administration of PREZISTA®/rtv with rifapentine is not recommended
β-Blockers: carvedilol metoprolol timolol	↑ beta-blockers	Co administration of PREZISTA®/rtv and beta blockers may increase concentrations of the beta blocker (inhibition of CYP2D6). Clinical monitoring is recommended when co administering PREZISTA®/rtv with beta blockers and a lower dose of the beta blocker should be considered.
Calcium Channel Blockers: amlodipine diltiazem felodipine nifedipine nicardipine verapamil	↑ calcium channel blockers	Plasma concentrations of calcium channel blockers (e.g., amlodipine, diltiazem, felodipine, nifedipine, nicardipine, verapamil) may increase when PREZISTA®/rtv are co-administered. Caution is warranted and clinical monitoring of patients is recommended.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Corticosteroids: Systemic dexamethasone prednisone Inhaled/Nasal budesonide fluticasone	↓ darunavir ↑ corticosteroid	Use with caution. Systemic dexamethasone induces CYP3A4 and can thereby decrease darunavir plasma concentrations. This may result in loss of therapeutic effect to PREZISTA®. Concomitant use of systemic or inhaled/nasal corticosteroids and PREZISTA®/rtv may increase plasma concentrations of these corticosteroids. Concomitant use may increase the risk for development of systemic corticosteroid effects, including Cushing's syndrome and adrenal suppression. Clinical monitoring is recommended when coadministering PREZISTA®/rtv with prednisone. Alternatives should be considered, particularly for long-term use.
Endothelin Receptor Antagonists: bosentan	↑ bosentan	Co-administration of bosentan in patients on PREZISTA®/rtv: In patients who have been receiving PREZISTA®/rtv for at least 10 days, start bosentan at 62.5 mg once daily or every other day based upon individual tolerability. Co-administration of PREZISTA®/rtv in patients on bosentan: Discontinue use of bosentan at least 36 hours prior to initiation of PREZISTA®/rtv. After at least 10 days following the initiation of PREZISTA®/rtv, resume bosentan at 62.5 mg once daily or every other day based upon individual tolerability.
Estrogen-Based Contraceptives: ethinyl estradiol norethindrone	↓ ethinyl estradiol ↓ norethindrone	The results of an interaction trial between PREZISTA®/rtv (600/100 mg b.i.d.) and ethinyl estradiol and norethindrone demonstrated that at steady-state, systemic exposures to ethinyl estradiol and norethindrone are decreased by 44% and 14%, respectively. Therefore, alternative methods of non-hormonal contraception are recommended.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
HMG-CoA Reductase Inhibitors: atorvastatin	↑atorvastatin	An interaction trial between darunavir (300 mg b.i.d.), low-dose ritonavir (100 mg b.i.d.), and atorvastatin (10 mg q.d.) demonstrated that exposure to atorvastatin was only 15% lower when co-administered with darunavir and ritonavir than when atorvastatin (40 mg q.d.) was administered alone. When administration of atorvastatin and PREZISTA®/rtv is desired, it is recommended to start with an atorvastatin dose of 10 mg q.d. A gradual dose increase of atorvastatin may be tailored to the clinical response.
rosuvastatin	↑rosuvastatin	An interaction study evaluating PREZISTA®/rtv (600/100 mg b.i.d.) in combination with rosuvastatin (10 mg q.d.) resulted in a significant increase in rosuvastatin plasma exposures. It is recommended to start with the lowest possible dose of rosuvastatin and titrate up to the desired clinical effect while monitoring for safety.
pravastatin	↑pravastatin	An interaction trial between darunavir (600 mg b.i.d.), low dose ritonavir (100 mg b.i.d.) and pravastatin (40 mg single dose) demonstrated that DRV/rtv did not increase exposure of pravastatin in most patients but increased pravastatin exposure up to 5-fold in a limited subset of patients. When administration of pravastatin and PREZISTA®/rtv is required, it is recommended to start with the lowest possible dose of pravastatin and titrate up to the desired clinical effects while monitoring safety.
H2-Receptor Antagonists and Proton Pump Inhibitors: cimetidine famotidine nizatidine ranitidine esomeprazole lansoprazole omeprazole pantoprazole rabeprazole	↔ darunavir	Co-administration of omeprazole (20 mg q.d.) or ranitidine (150 mg b.i.d.) and darunavir (400 mg b.i.d.) in the presence of low-dose ritonavir (100 mg b.i.d.) did not affect the exposure to darunavir. Based on these results, PREZISTA®/rtv can be co-administered with H2-receptor antagonists and proton pump inhibitors without dose adjustments. The effects of PREZISTA®/rtv on omeprazole or ranitidine exposures were not evaluated.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Immunosuppressants: cyclosporine tacrolimus sirolimus everolimus	↑ immunosuppressants	Plasma concentrations of these immunosuppressants may be increased when co-administered with PREZISTA®/rtv. Therapeutic concentration monitoring of the immunosuppressive agent is recommended for immunosuppressant agents when co-administered with PREZISTA®/rtv. Concomitant use of everolimus and PREZISTA®/rtv is not recommended.
Inhaled Beta Agonist: salmeterol		Concurrent administration of salmeterol and PREZISTA®/rtv is not recommended. The combination may result in increased risk of cardiovascular adverse events associated with salmeterol, including QT prolongation, palpitations and sinus tachycardia.
Narcotic Analgesics/Treatment of Opioid Dependence: methadone	↓ methadone	An interaction trial investigating the effect of PREZISTA®/rtv (600/100 mg b.i.d.) on a stable methadone maintenance therapy showed an AUC decrease of 16% for R-methadone. Based on pharmacokinetic and clinical findings, no adjustment of methadone dosage is required when initiating co-administration of PREZISTA®/rtv. However, clinical monitoring is recommended, as maintenance therapy may need to be adjusted in some patients.
meperidine	↓ meperidine	PREZISTA®/rtv is expected to decrease meperidine concentrations and increase normeperidine metabolite concentrations. Dosage increase and long-term use of meperidine and PREZISTA®/rtv are not recommended due to the increased concentrations of the metabolite normeperidine, which has both analgesic and CNS stimulant activity (e.g., seizures).
buprenorphine/naloxone	↑ norbuprenorphine	The results of an interaction trial with PREZISTA®/rtv and buprenorphine/naloxone demonstrated that buprenorphine exposure was not affected when administered with PREZISTA®/rtv. Exposure of the active metabolite, norbuprenorphine, increased by 46%. No dose adjustment for buprenorphine was required. Careful clinical monitoring is recommended if PREZISTA®/rtv and buprenorphine are co-administered.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Neuroleptics: risperidone thioridazine quetiapine	↑ neuroleptics	Concomitant use of risperidone or thioridazine and PREZISTA®/rtv may increase the exposure to these antipsychotics (inhibition CYP2D6 and/or P-gp). Decrease of risperidone or thioridazine dose may be needed when co administered with PREZISTA®/rtv. PREZISTA®/rtv should not be used in combination with quetiapine. Due to CYP3A inhibition by PREZISTA®/rtv, concentrations of quetiapine are expected to increase, which can result in serious and/or life-threatening adverse reactions. Refer to Norvir® Product Monograph.
NS3-4A protease inhibitors [Hepatitis C Virus (HCV) direct-acting antivirals]**:** boceprevir telaprevir[a]	↓ darunavir ↓ boceprevir ↓ telaprevir	In an interaction trial between PREZISTA®/rtv (600/100 mg b.i.d.) and boceprevir (800 mg three times daily), darunavir exposure was reduced by 44% and boceprevir exposure was reduced by 32%. It is not recommended to co-administer PREZISTA®/rtv with boceprevir. In a drug interaction study in healthy volunteers where telaprevir was co-administered with darunavir/ritonavir, the steady-state exposure to telaprevir and darunavir was reduced. Coadministration of darunavir/ritonavir and telaprevir is not recommended.
simeprevir	↑ simeprevir	Co administration of PREZISTA®/rtv (800/100 mg q.d.) and simeprevir increased darunavir and simeprevir concentrations (inhibition of CYP3A). In an interaction trial between PREZISTA®/rtv (800/100 mg q.d.) and simeprevir (50 mg q.d.), simeprevir exposure increased 2.59-fold and darunavir exposure increased by 1.18 fold. The combination of PREZISTA®/rtv and simeprevir is not recommended.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
PDE-5 Inhibitors: sildenafil, tadalafil vardenafil	↑ PDE-5 inhibitors	In an interaction trial, a comparable systemic exposure to sildenafil was observed for a single dose of 100 mg sildenafil alone and a single dose of 25 mg sildenafil co-administered with darunavir (400 mg b.i.d.) and low-dose ritonavir (100 mg b.i.d.). Co-administration with PREZISTA®/rtv may result in an increase in PDE-5 inhibitor-associated adverse events, including hypotension, syncope, visual disturbances and priapism. Use of PDE-5 inhibitors for erectile dysfunction: Concomitant use of PDE-5 inhibitors, when used for the treatment of erectile dysfunction, should be done with caution. Co-administration of darunavir and low-dose ritonavir with sildenafil or tadalafil is expected to substantially increase the PDE-5 concentration and may result in an increase in PDE-5 inhibitor-associated adverse events including hypotension, visual changes, syncope and priapism. If concomitant use of PREZISTA®/rtv with sildenafil or tadalafil is required, sildenafil at a single dose not exceeding 25 mg in 48 hours or tadalafil at a single dose not exceeding 10 mg in 72 hours is recommended. Vardenafil should not be used with PREZISTA®/rtv. Use of PDE-5 inhibitors for pulmonary arterial hypertension (PAH): Use of sildenafil is contraindicated (see Table 8). Based on theoretical considerations, co-administration of PREZISTA® with tadalafil may increase concentrations of tadalafil (CYP3A inhibition). Co-administration of PREZISTA® with tadalafil is not recommended.
Sedatives/Hypnotics: buspirone clorazepate diazepam estazolam flurazepam zoldipem[a] parenterally administered midazolam	↑ sedatives/hypnotics	Co administration of PREZISTA®/rtv with these sedatives/hypnotics may increase concentrations of the sedative/hypnotic (inhibition of CYP3A). Clinical monitoring is recommended when co administering PREZISTA®/rtv with these sedatives/hypnotics and a lower dose of the sedatives/hypnotics should be considered. Co-administration of parenteral midazolam should be done in a setting that ensures close clinical monitoring and appropriate medical management in case of respiratory depression and/or prolonged sedation. Dose reduction for parenteral midazolam should be considered, especially if more than a single dose of midazolam is administered.

Table 9: Established and Other Potentially Significant Drug Interactions: Alterations in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction (See Tables 29 and 30)		
Concomitant Drug Class: Drug Name	**Effect on Concentration of Darunavir or Concomitant Drug**	**Clinical Comment**
Antidepressants: sertraline paroxetine	↔ darunavir ↓ sertraline ↓ paroxetine	An interaction trial between paroxetine (20 mg q.d.) or sertraline (50 mg q.d.) and darunavir (400 mg b.i.d.) and low-dose ritonavir (100 mg b.i.d.) demonstrated that exposure to darunavir was not affected by the co-administration of sertraline or paroxetine. Exposure to sertraline or paroxetine decreased by 49% and 39%, respectively, when co-administered with darunavir and ritonavir. If sertraline or paroxetine is co-administered with PREZISTA®/rtv, the recommended approach is a careful dose titration of the SSRI based on a clinical assessment of antidepressant response. In addition, patients on a stable dose of sertraline or paroxetine who start treatment with PREZISTA®/rtv should be monitored for antidepressant response.
amitriptyline desipramine imipramine nortriptyline trazodone	↑ antidepressants	Concomitant use of PREZISTA®/rtv and these antidepressants is expected to increase concentrations of the antidepressant (inhibition of CYP2D6 and/or CYP3A). Clinical monitoring and dose adjustment are recommended when co administering PREZISTA®/rtv with these antidepressants.

[a] Not marketed in Canada

Other NRTIs

Based on the different elimination pathways of the other NRTIs (zidovudine, zalcitabine, emtricitabine, stavudine, lamivudine and abacavir) that are primarily renally excreted, no drug interactions are expected for these drugs and PREZISTA®/rtv.

Other Protease Inhibitors

The co-administration of PREZISTA®/rtv and PIs other than lopinavir/ritonavir, saquinavir, atazanavir, and indinavir has not been studied. Therefore, such co-administration is not recommended.

CCR5 Antagonist

When used in combination with PREZISTA®/rtv, the dose of maraviroc should be 150 mg twice daily.

An interaction trial between PREZISTA®/rtv (600/100 mg b.i.d.) and maraviroc (150 mg b.i.d.) demonstrated that in the presence of PREZISTA®/rtv the exposure of maraviroc was increased by 305%. There was no apparent effect of maraviroc on darunavir/ritonavir exposure.

Drug-Food Interactions
Darunavir, when given as a tablet and co-administered with low-dose ritonavir as a pharmacokinetic enhancer, should be taken with food. The type of food does not affect the exposure to darunavir.

Drug-Herb Interactions
Concomitant use of PREZISTA®/rtv and St. John's wort (*Hypericum perforatum*) or products containing St. John's wort is contraindicated. Co-administration of protease inhibitors (PIs), including PREZISTA®/rtv, with St. John's wort is expected to substantially decrease PI concentrations and may result in suboptimal concentrations of darunavir and lead to loss of virologic response and possible resistance to PREZISTA®/rtv or to the class of PIs (see **Drug-Drug Interactions**, Table 8).

Interactions with other herbal products have not been established.

Drug-Laboratory Interactions
Interactions with laboratory tests have not been established.

DOSAGE AND ADMINISTRATION

Dosing Considerations

PREZISTA® (darunavir) must always be given with ritonavir as a pharmacokinetic enhancer, and in combination with other antiretroviral medicinal products. The prescribing information of ritonavir must, therefore, be consulted prior to initiation of therapy with PREZISTA®/rtv. The treatment history and, when available, genotypic or phenotypic testing should guide the use of PREZISTA®/rtv (see **DOSAGE AND ADMINISTRATION, Recommended Dose and Dosage Adjustment,** Adults).

Recommended Dose and Dosage Adjustment

Patients who have difficulty swallowing PREZISTA® tablets can use PREZISTA® oral suspension.

Adults
The recommended oral dosing regimens for adult patients are PREZISTA® 800 mg in combination with ritonavir (100 mg) once daily and with food, or 600 mg in combination with ritonavir (100 mg) twice daily and with food. The type of food does not affect exposure to darunavir (see **DRUG INTERACTIONS**, **Drug-Food Interactions**). The dosing schedule for PREZISTA®/rtv is presented in Table 10.

Table 10: Recommended Dose for Adult Patients for PREZISTA® Tablets with ritonavir (100 mg)		
Treatment-Naïve Adult Patients	**Treatment-Experienced Adult Patients**	
	With no darunavir resistance-associated mutations (DRV-RAMs)[a]	**With at least one darunavir resistance-associated mutation (DRV-RAM)[a]**
800 mg (two 400 mg tablets or one 800 mg tablet or 8 mL oral suspension[b]) PREZISTA® in combination with ritonavir 100 mg (one 100 mg Tablet/Capsule or 1.25 ml of the 80 mg/ml ritonavir solution) once daily and with food	800 mg (two 400 mg tablets or one 800 mg tablet or 8 mL oral suspension[b]) PREZISTA® in combination with ritonavir 100 mg (one 100 mg Tablet/Capsule or 1.25 ml of the 80 mg/ml ritonavir solution) once daily and with food	600 mg (one 600 mg tablet or 6 mL oral suspension) PREZISTA® in combination with 100 mg ritonavir (one 100 mg Tablet/Capsule or 1.25 ml of the 80 mg/ml ritonavir solution) twice daily and with food

[a] DRV-RAMs: V11I, V321, L33F, I47V, I50V, I54M, I54L, T74P, L76V, I84V and L89V
[b] An 8 mL dose should be taken as two 4 mL administrations with the included oral dosing syringe.

For antiretroviral treatment-experienced patients HIV genotypic testing is recommended. However, when HIV genotypic testing is not feasible, the PREZISTA®/rtv once-daily dosing regimen is recommended in protease inhibitor-naive patients and the twice daily dosing regimen is recommended in protease inhibitor-experienced patients.

The type of food does not affect exposure to darunavir. Ritonavir is used as a pharmacokinetic enhancer of PREZISTA® (see **DRUG INTERACTIONS**, **Drug-Drug Interactions**, and **ACTION AND CLINICAL PHARMACOLOGY, Pharmacokinetics,** Effects of Food on Oral Absorption). A further increase in the dose of darunavir or ritonavir is not likely to result in any clinically relevant increase in antiviral activity.

Geriatric Patients
In general, caution should be exercised in the administration and monitoring of PREZISTA® in elderly patients, reflecting the greater frequency of decreased hepatic, renal or cardiac function, and of concomitant disease or other drug therapy (see **INDICATIONS AND CLINICAL USE**, **WARNINGS AND PRECAUTIONS** and **ACTION AND CLINICAL PHARMACOLOGY**).

Pediatric Patients
PREZISTA®/rtv should not be used in pediatric patients below 3 years of age (see **WARNINGS AND PRECAUTIONS** and **TOXICOLOGY**).

Antiretroviral Treatment-Experienced Pediatric Patients 3 to < 18 years of age
The recommended dose of PREZISTA® for pediatric patients (3 to < 18 years of age and weighing at least 22 lbs (10 kg)) is based on body weight (see Tables 11 and 12) and should not exceed the recommended adult dose (PREZISTA®/rtv 600/100 mg b.i.d.) (see **CLINICAL TRIALS)**. PREZISTA® should be taken with ritonavir twice daily and with food. The type of food does not affect exposure to darunavir (see **ACTION AND CLINICAL PHARMACOLOGY, Pharmacokinetics,** Effects of Food on Oral Absorption).

Before prescribing PREZISTA®, children weighing greater than or equal to 15 kg should be assessed for the ability to swallow tablets. If a child is unable to reliably swallow a tablet, the use of PREZISTA® oral suspension should be considered.

The weight-based dose in pediatric patients weighing at least 10 kg and < 15 kg is PREZISTA® 20 mg/kg twice daily with ritonavir 3 mg/kg twice daily. The following table provides recommended weight-based dosing.

Table 11: Recommended Dose for Pediatric Patients weighing 10 kg to <15 kg who are treatment experienced with at least one darunavir resistance associated mutation (DRV-RAM)[a] using PREZISTA® Oral Suspension with ritonavir[b]

Body weight		Dose (twice Daily with food)
(kg)	(lbs)	
≥ 10 kg – < 11 kg	≥ 22 lbs – < 24.2 lbs	200 mg (2 mL) PREZISTA® with 32 mg (0.4 mL) ritonavir twice daily
≥ 11 kg – < 12 kg	≥ 24.2 lbs – < 26.4 lbs	220 mg (2.2 mL) PREZISTA® with 32 mg (0.4 mL) ritonavir twice daily
≥ 12 kg – < 13 kg	≥ 26.4 lbs – < 28.6 lbs	240 mg (2.4 mL) PREZISTA® with 40 mg (0.5 mL) ritonavir twice daily
≥ 13 kg – < 14 kg	≥ 28.6 lbs – < 30.8 lbs	260 mg (2.6 mL) PREZISTA® with 40 mg (0.5 mL) ritonavir twice daily
≥ 14 kg – < 15 kg	≥ 30.8 lbs – < 33 lbs	280 mg (2.8 mL) PREZISTA® with 48 mg (0.6 mL) ritonavir twice daily

[a] DRV-RAMs: V11I, V321, L33F, I47V, I50V, I54M, I54L, T74P, L76V, I84V and L89V
[b] Ritonavir oral solution: 80 mg/mL

The following table provides recommended weight-based dosing for pediatric patients weighing ≥ 15 kg:

Table 12: Recommended Dose for Pediatric Patients weighing ≥ 15 kg who are treatment experienced with at least one darunavir resistance associated mutation (DRV-RAM)[a] using PREZISTA® Tablets or Oral Suspension with ritonavir[b]

Body Weight		Dose (twice daily with food)
(kg)	(lbs)	
≥ 15 kg – < 30 kg	≥ 33 lbs – < 66 lbs	375 mg[c] (3.8 mL) PREZISTA® with 50 mg (0.6 mL) ritonavir twice daily
≥ 30 kg – < 40 kg	≥ 66 lbs – < 88 lbs	450 mg[d] (4.6mL) PREZISTA® with 60 mg (0.8 mL) ritonavir twice daily
≥ 40 kg	≥ 88 lbs	600 mg (6 mL) PREZISTA® with 100 mg (1.25 mL) ritonavir twice daily

[a] DRV-RAMs: V11I, V321, L33F, I47V, I50V, I54M, I54L, T74P, L76V, I84V and L89V
[b] Given as a tablet or capsule, or with ritonavir oral solution: 80 mg/mL

[c] The 375 mg dose refers to the dose using darunavir tablets for this weight group, which is rounded off to 3.8 mL for suspension dosing.
[d] The 450 mg dose refers to the dose using darunavir tablets for this weight group, which is rounded off to 4.6 mL for suspension dosing.

Antiretroviral Treatment-Naïve Pediatric Patients
The safety and efficacy of PREZISTA®/rtv in antiretroviral treatment-naïve pediatric patients have not been established.

Hepatic Impairment
The safety and efficacy of PREZISTA® have not been established in patients with severe hepatic insufficiency (see **CONTRAINDICATIONS**). No dose adjustment is required in patients with mild or moderate hepatic impairment (see **ACTION AND CLINICAL PHARMACOLOGY**, **Pharmacokinetics**).

Renal Impairment
No dose adjustment is required in patients with renal impairment (see **WARNINGS AND PRECAUTIONS**, **Renal** and **ACTION AND CLINICAL PHARMACOLOGY**, **Pharmacokinetics**).

Dosing with Didanosine
Dosing of enteric-coated didanosine and darunavir, co-administered with low-dose ritonavir, should be separated by at least 2 hours to avoid formulation incompatibility.

Missed Dose
Patients Taking 600 mg of PREZISTA® Twice Daily
The missed dose should be taken as soon as possible, if the dose was missed by less than 6 hours. The next dose of PREZISTA® and ritonavir should be taken at the regularly scheduled time. If the dose of PREZISTA® or ritonavir was missed by more than 6 hours, the next dose of PREZISTA® and ritonavir should be taken at the regularly scheduled time. Doses should not be doubled.

Patients Taking 800 mg of PREZISTA® Once Daily
The missed dose (two 400 mg tablets or one 800 mg tablet) should be taken as soon as possible, if the dose was missed by less than 12 hours. The next dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet) and ritonavir should be taken at the regularly scheduled time. If the dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet) or ritonavir was missed by more than 12 hours, the next dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet) and ritonavir should be taken at the regularly scheduled time. Doses should not be doubled.

OVERDOSAGE

For management of a suspected drug overdose, contact your regional Poison Control Centre.

Human experience of acute overdose with PREZISTA®/rtv is limited. Single doses up to 3,200 mg of the oral solution of darunavir alone and up to 1,600 mg of the tablet formulation of darunavir co-administered with ritonavir have been administered to healthy volunteers without untoward symptomatic effects.

There is no specific antidote for overdose with PREZISTA®. Treatment of overdose with PREZISTA® consists of general supportive measures including monitoring of vital signs and observation of the clinical status of the patient. Administration of activated charcoal may be used to aid in removal of unabsorbed active substance. Since PREZISTA® is highly protein bound, dialysis is unlikely to be beneficial in significant removal of the active substance.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

Darunavir is an inhibitor of the dimerization and of the catalytic activity of the HIV-1 protease. It selectively inhibits the cleavage of HIV-encoded Gag-Pol polyproteins in virus-infected cells, thereby preventing the formation of mature infectious virus particles.

Darunavir tightly binds to the HIV-1 protease with a K_D of 4.5 x 10^{-12} M.

Darunavir is not an inhibitor of any of 13 tested human cellular proteases.

Pharmacodynamics

Electrocardiogram (Effect on QT Intervals)

In an open-label, randomized, placebo- and active-controlled, four-way crossover trial, 40 healthy subjects were administered supratherapeutic doses of darunavir/ritonavir 1,600/100 mg once daily and 800/100 mg twice daily for seven days.

At the mean maximum darunavir concentration of 6,599 ng/mL observed in this study, the mean increase in QTcF was 2.2 ms with a 90% two-sided confidence interval (CI) of –2.0 to 6.3 ms. When evaluating the two-sided 90% CI on the time-matched mean changes in QTcF versus placebo control, the upper bounds of both darunavir/ritonavir groups never exceeded the 10 ms boundary. In the setting of this trial, darunavir/ritonavir did not appear to prolong the QTc interval.

Pharmacokinetics

General

Darunavir is primarily metabolized by CYP3A4. The overall pharmacokinetic enhancement effect by ritonavir was an approximate 14-fold increase in the systemic exposure of darunavir when a single dose of 600 mg darunavir was given orally co-administered with ritonavir at 100 mg b.i.d. Therefore, PREZISTA® should only be co-administered with 100 mg of ritonavir as a pharmacokinetic enhancer.

Table 13: Pharmacokinetic Parameters of Darunavir 600 mg and Darunavir/rtv at 600/100 mg b.i.d.

Pharmacokinetics of darunavir	Darunavir 600 mg Oral N=7 *(mean ±SD, t_{max}: median [range])*	Darunavir 600 mg Oral + rtv 100 mg b.i.d. N=7 *(mean ±SD, t_{max}: median [range])*
t_{max}, h	2.00 (1.00 – 4.00)	4.00 (2.00 – 5.00)
C_{max}, ng/mL	2,204 ± 1071	5,627 ± 923.5
AUC_{last}, ng.h/mL	7,748 ± 4867	91,390 ± 20050
AUC_{∞}, ng.h/mL	10,990 ± 4061	92,340 ± 20020
Bioavailability (F) (%)	36.93	81.93

Absorption and Bioavailability
Darunavir was rapidly absorbed following oral administration. Maximum plasma concentration of darunavir in the presence of low-dose ritonavir is generally achieved within 2.5-4.0 hours.

The absolute oral bioavailability of a single 600 mg dose of darunavir alone was approximately 37% and increased to approximately 82% in the presence of 100 mg b.i.d. ritonavir. Increasing the dose of ritonavir to above 100 mg b.i.d. did not significantly affect darunavir concentrations. It is not recommended to alter the dose of ritonavir. *In vivo* data suggest that darunavir/ritonavir is an inhibitor of the p-glycoprotein (p-gp) transporter.

Effects of Food on Oral Absorption
When administered without food, the relative bioavailability of darunavir in the presence of low-dose ritonavir is 30% lower as compared to intake with food. Therefore, PREZISTA® tablets should be taken with ritonavir and with food. The type of food does not affect exposure to darunavir.

Food has no effect on the oral bioavailability of darunavir when administered as a suspension formulation following a single 600 mg dose of darunavir taken with low-dose ritonavir.

Distribution
Darunavir is approximately 95% bound to plasma proteins. Darunavir binds primarily to plasma alpha-1-acid glycoprotein (AAG).

Metabolism
In vitro experiments with human liver microsomes (HLMs) indicate that darunavir primarily undergoes oxidative metabolism. Darunavir is extensively metabolized by the hepatic CYP system, and almost exclusively by isozyme CYP3A4. A ^{14}C-darunavir trial in healthy volunteers showed that a majority of the radioactivity in plasma after a single 400/100 mg darunavir/ritonavir dose was due to the parent drug. At least three oxidative metabolites of darunavir have been identified in humans; all showed activity that was at least 10-fold less than the activity of darunavir against wild-type HIV.

Excretion
After a 400/100 mg ^{14}C-darunavir/ritonavir dose, approximately 79.5% and 13.9% of the administered dose of ^{14}C-darunavir could be retrieved in feces and urine, respectively.

Unchanged darunavir accounted for approximately 41.2% and 7.7% of the administered dose in feces and urine, respectively. The terminal elimination half-life of darunavir was approximately 15 hours when combined with ritonavir. The intravenous clearance of darunavir alone (150 mg) and in the presence of low-dose ritonavir was 32.8 L/h and 5.9 L/h, respectively.

Special Populations and Conditions

Pediatrics

The pharmacokinetics of darunavir in combination with ritonavir in 74 antiretroviral treatment-experienced HIV-1-infected pediatric patients 6 to <18 years of age and weighing at least 44 lbs (20 kg) showed that the administered weight-based dosages resulted in darunavir exposure comparable to that in adults receiving PREZISTA®/rtv 600/100 mg b.i.d. (see **DOSAGE AND ADMINISTRATION**).

The pharmacokinetics of darunavir in combination with ritonavir in 19 antiretroviral treatment-experienced pediatric patients, aged 3 to < 6 years and weighing at least 10 kg to 20 kg showed that weight-based dosages resulted in darunavir exposure that was comparable to or slightly higher to that achieved in adults receiving PREZISTA®/rtv 600/100 mg b.i.d. (see **DOSAGE AND ADMINISTRATION**).

Table 14: Population Pharmacokinetic Estimates of Darunavir Exposure (Study TMC114-C230, Study TMC114-C212 and Study TMC114-C228) Following Administration of Doses in Tables 11 and 12[a]

Parameter	Study TMC114-C212 PREZISTA®/ritonavir twice daily N = 74	Study TMC114-C228 PREZISTA®/ ritonavir twice daily[b]	
		10 to less than 15 kg[d] N = 10	15 to less than 20 kg[e] N = 13
AUC_{24h} (ng·h/mL)[c]			
Mean ± Standard Deviation	126377 ± 34356	137896 ± 51420	157760 ± 54080
Median (Range)	127340 (67054-230720)	124044 (89688-261090)	132698 (112310 – 294840)
C_{0h} (ng/mL)			
Mean ± Standard Deviation	3948 ± 1363	4510 ± 2031	4848 ± 2143
Median (Range)	3888 (1836-7821)	4126 (2456-9361)	3927 (3046 – 10292)

N = number of subjects with data. [a] Summary statistics for population pharmacokinetic parameter estimates for DRV after administration of DRV/rtv at 800/100 mg q.d. in treatment-naïve HIV-1 infected subjects from 12 to < 18 years of age –Week-48 Analyses [b] Subjects may have contributed pharmacokinetic data to both the 10 kg to less than 15 kg weight group and the 15 kg to less than 20 kg weight group. [c] AUC24h is calculated as AUC12h*2 [d] Calculated from individual pharmacokinetic parameters estimated for Week 2 and Week 4, based on the Week 48 analysis that evaluated a darunavir dose of 20 mg/kg twice daily with ritonavir 3 mg/kg twice daily. [e] The 15 kg to less than 20 kg weight group received 380 mg (3.8 mL) PREZISTA® oral suspension twice daily with 48 mg (0.6 mL) ritonavir oral solution twice daily in TMC114-C228. Calculated from individual pharmacokinetic parameters estimated for Week 2 post-dose adjustment visit; Week 24 and Week 48 based on the -Week 48 analysis that evaluated a darunavir dose of 380 mg twice daily.

Geriatrics

Population pharmacokinetic analysis in HIV-1-infected patients showed that darunavir pharmacokinetics are not considerably different in the age range (18 to 75 years) evaluated in HIV-1-infected patients (n=12, age ≥ 65) (see **WARNINGS AND PRECAUTIONS**, **Special Populations**, **Geriatrics**).

Gender
Population pharmacokinetic analysis showed a slightly higher darunavir exposure (16.8%) in HIV-1-infected females (n=68) compared to males. This difference is not considered clinically relevant.

Race
Population pharmacokinetic analysis of darunavir in HIV-1-infected patients indicated that race had no apparent effect on the exposure to darunavir.

Hepatic Insufficiency
Hepatic Impairment
In a multiple dose study with PREZISTA® co-administered with ritonavir (600/100 mg) twice daily, it was demonstrated that the steady-state pharmacokinetic parameters of darunavir in patients with mild (Child-Pugh Class A, n=8) and moderate (Child-Pugh Class B, n=8) hepatic impairment were comparable with those in healthy patients. The effect of severe hepatic impairment of on the pharmacokinetics of darunavir has not been studied (see **CONTRAINDICATIONS, WARNINGS AND PRECAUTIONS**, and **DOSAGE AND ADMINISTRATION**).

Hepatitis B or Hepatitis C Virus Co-infection
The primary 48-week analysis of the data from Study TMC114-C211 and TMC114-C214 in HIV-1-infected patients indicated that hepatitis B and/or hepatitis C virus co-infection status had no apparent effect on the exposure to darunavir.

Renal Insufficiency
Results from a mass balance study with ^{14}C-darunavir/ritonavir showed that approximately 7.7% of the administered dose of darunavir is excreted in the urine as unchanged drug. As darunavir and ritonavir are highly bound to plasma proteins, it is unlikely that they will be significantly removed by hemodialysis or peritoneal dialysis. Although darunavir has not been studied in patients with renal impairment, population pharmacokinetic analysis showed that the pharmacokinetics of darunavir were not significantly affected in HIV-1-infected patients with moderate renal impairment (CrCL between 30–60 mL/min, n=20) (see **WARNINGS AND PRECAUTIONS**, **Renal** and **DOSAGE AND ADMINISTRATION**, **Recommended Dose and Dosage Adjustment**, **Renal Impairment**).

Pregnancy
Pregnancy and Postpartum
The exposure to total darunavir and ritonavir after intake of darunavir/ritonavir 600/100 mg b.i.d and darunavir/ritonavir 800/100 mg q.d. as part of an antiretroviral regimen was generally lower during pregnancy compared with postpartum (see Table 15 and Table 16). However, for unbound (i.e., active) darunavir, the pharmacokinetic parameters were less reduced during pregnancy compared to postpartum, due to an increase in the unbound fraction of darunavir during pregnancy compared to postpartum.

Table 14: Pharmacokinetic Results of Total darunavir After Administration of darunavir/ritonavir at 600/100 mg bid as Part of an Antiretroviral Regimen, During the 2nd Trimester of Pregnancy, the 3rd Trimester of Pregnancy and Postpartum

Pharmacokinetics of total darunavir (mean ± SD)	**2nd Trimester of Pregnancy (n=11)[a]**	**3rd Trimester of Pregnancy (n=11)**	**Postpartum (6-12 Weeks) (n=11)**
C_{max}, ng/mL	4601 ± 1125	5111 ± 1517	6499 ± 2411
AUC_{12h}, ng.h/mL	38950 ± 10010	43700 ± 16400	55300 ± 27020
C_{min}, ng/mL[b]	1980 ± 839.9	2498 ± 1193	2711 ± 2268

[a] n=10 for AUC_{12h}
[b] excluding C_{min} value below LLOQ, n=10 for reference

Table 15: Pharmacokinetic Results of Total darunavir After Administration of darunavir/ritonavir at 800/100 mg qd as Part of an Antiretroviral Regimen, During the 2nd Trimester of Pregnancy, the 3rd Trimester of Pregnancy and Postpartum

Pharmacokinetics of total darunavir (mean ± SD)	**2nd Trimester of Pregnancy (n=16)**	**3rd Trimester of Pregnancy (n=14)**	**Postpartum (6-12 Weeks) (n=15)**
C_{max}, ng/mL	4988 ± 1551	5138 ± 1243	7445 ± 1674
AUC_{24h}, ng.h/mL	61303 ± 16232	60439 ± 14052	94529 ± 28572
C_{min}, ng/mL[a]	1193 ± 509	1098 ± 609	1572 ± 1108

[a] N=12 for postpartum, N=15 for 2nd trimester and N=14 for 3rd trimester

In women receiving darunavir/ritonavir 600/100 mg b.i.d during the 2nd trimester of pregnancy, mean intra-individual values for total darunavir C_{max}, AUC_{12h} and C_{min} were 28%, 24% and 17% lower, respectively, as compared with postpartum; during the 3rd trimester of pregnancy, total darunavir C_{max}, AUC_{12h} and C_{min} values were 19%, 17% lower and 2% higher, respectively, as compared with postpartum.

In women receiving darunavir/ritonavir 800/100 mg q.d during the 2nd trimester of pregnancy, mean intra-individual values for total darunavir C_{max}, AUC_{24h} and C_{min} were 34%, 34% and 32% lower, respectively, as compared with postpartum; during the 3rd trimester of pregnancy, total darunavir C_{max}, AUC_{24h} and C_{min} values were 31%, 35% and 50% lower, respectively, as compared with postpartum.

STORAGE AND STABILITY

Store PREZISTA® tablets between 15 – 30°C.

Store PREZISTA® oral suspension at room temperature between 15 – 30°C. Do not refrigerate or freeze. Avoid exposure to excessive heat. Store in the original container.

DOSAGE FORMS, COMPOSITION AND PACKAGING

PREZISTA® 75 mg Tablets

PREZISTA® (darunavir) 75 mg tablets are available as white, caplet-shaped, film-coated tablets containing 75 mg of darunavir (corresponding to 81.31 mg of darunavir ethanolate). Each tablet is debossed with "75" on one side and "TMC" on the other side.

PREZISTA® 150 mg Tablets
PREZISTA® (darunavir) 150 mg tablets are supplied as white, oval-shaped, film-coated tablets containing 150 mg of darunavir (corresponding to 162.62 mg of darunavir ethanolate). Each tablet is debossed with "150" on one side and "TMC" on the other side.

PREZISTA® 400 mg Tablets
PREZISTA® (darunavir) 400 mg tablets are supplied as light orange, oval-shaped, film-coated tablets containing darunavir ethanolate equivalent to 400 mg of darunavir per tablet. Each tablet is debossed with "400MG" on one side and "TMC" on the other side.
PREZISTA® 600 mg Tablets
PREZISTA® (darunavir) 600 mg tablets are supplied as orange, oval-shaped, film-coated tablets containing darunavir ethanolate equivalent to 600 mg of darunavir per tablet. Each tablet is debossed with "600MG" on one side and "TMC" on the other side.

PREZISTA® 800 mg Tablets
PREZISTA® (darunavir) 800 mg tablets are supplied as dark red, oval-shaped, film-coated tablets containing darunavir ethanolate equivalent to 800 mg of darunavir per tablet. Each tablet is debossed with "800" on one side and "T" on the other side.

Each tablet also contains the inactive ingredients microcrystalline cellulose, colloidal silicon dioxide, crospovidone, and magnesium stearate. The 800 mg tablet core also contains hypromellose. The tablet film coatings contain either OPADRY® II White (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, talc, titanium dioxide) for the 75 and 150 mg tablets or OPADRY® II Orange (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, titanium dioxide, talc, sunset yellow FCF aluminum lake) for the 400 and 600 mg tablets or OPADRY® II Dark Red (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, titanium dioxide, talc, iron oxide red) for the 800 mg tablets.

PREZISTA® tablets are packaged in bottles in the following configuration:

75 mg tablets—bottles of 480 tablets
150 mg tablets—bottles of 240 tablets
400 mg tablets—bottles of 60
600 mg tablets—bottles of 60
800 mg tablets—bottles of 30

PREZISTA® 100 mg/mL Oral Suspension
PREZISTA® (darunavir) oral suspension is supplied as a white to off-white opaque suspension of 200 mL in an amber glass bottle with polypropylene child-resistant closure and a 6 mL low density polyethylene (LDPE) oral dosing pipette with 0.2 mL gradations. Each mL of the oral suspension contains 108.4 mg darunavir ethanolate equivalent to 100 mg darunavir. The supplied oral dosing pipette should not be used for any other medications.

The oral suspension contains the inactive ingredients hydroxypropyl cellulose, microcrystalline cellulose and carmellose sodium, sucralose, sodium methyl parahydroxybenzoate, masking flavor, citric acid monohydrate, and strawberry cream flavor.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

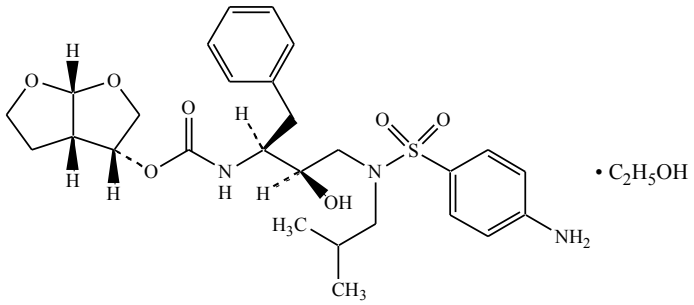
Common name:
Darunavir ethanolate

Chemical name:
[(1*S*,2*R*)-3-[[(4-aminophenyl)sulfonyl](2-methylpropyl)amino]-2-hydroxy-1-(phenylmethyl)propyl]-carbamic acid (3*R*,3a*S*,6a*R*)-hexahydrofuro[2,3-*b*]furan-3-yl ester ethanolate

Molecular formula and molecular mass:
Molecular formula: $C_{27}H_{37}N_3O_7S.C_2H_5OH$

Molecular mass: 547.66 (593.73 as the ethanolate)

Structural formula:



Physicochemical properties:

Physical Description: Darunavir ethanolate is a white to off-white powder.

Solubility: The solubility of darunavir (or darunavir ethanolate) is approximately 0.015 mg/mL in water at 20°C.

CLINICAL TRIALS

General

The evidence of efficacy of PREZISTA®/rtv is based on the analyses of 192-week data from a randomized, controlled open-label Phase 3 trial in treatment-naïve (TMC114-C211 [ARTEMIS]) HIV-1-infected adult subjects and 96-week data from a randomized, controlled, open-label Phase 3 trial in antiretroviral treatment-experienced (TMC114-C214 [TITAN]) HIV-1-infected adult patients, and the analysis of 48-week data from one randomized, open-label Phase 3 trial in early treatment experienced HIV-1-infected adult patients (TMC114-C229 [ODIN]).

In addition, 96-week data is included from 2 randomized, controlled Phase 2b trials, TMC114-C213 (POWER 1) and TMC114-C202 (POWER 2), in antiretroviral treatment-experienced HIV-1-infected adult patients, and 96-week data is included from the open label trial TMC114-C215 (POWER 3) in patients who initiated PREZISTA®/rtv at the recommended dose.

Antiretroviral Treatment-Naïve Adult Patients
TMC114-C211 (ARTEMIS)

Demographics and Trial Design
The evidence of efficacy of PREZISTA®/rtv 800/100 mg q.d. is based on the analyses of 192-week data from the randomized, controlled, open-label, Phase 3 trial TMC114-C211 in antiretroviral treatment-naïve HIV-1-infected patients comparing PREZISTA®/rtv 800/100 mg q.d. with lopinavir/rtv 800/200 mg per day (given as a twice-daily or as a once-daily regimen). Both arms used a fixed background regimen consisting of tenofovir disoproxil fumarate 300 mg q.d. (TDF) and emtricitabine 200 mg q.d. (FTC).

HIV-1 infected patients who were eligible for this trial had plasma HIV-1 RNA > 5000 copies/mL. Randomization was stratified by screening plasma viral load and screening CD4+ cell count. Virologic response was defined as a confirmed plasma HIV-1 RNA viral load < 50 copies/mL. Analyses included 689 patients in Study TMC114-C211 who had completed 48 and 192 weeks of treatment or discontinued earlier.

Demographics and baseline characteristics were balanced between the PREZISTA®/rtv arm and the lopinavir/rtv arm (see Table 17). Table 15 compares the demographic and baseline characteristics between patients in the PREZISTA®/rtv 800/100 mg q.d. arm and patients in the lopinavir/ritonavir 800/200 mg per day arm in the ARTEMIS trial. The 343 patients on PREZISTA®/rtv 800/100 mg q.d. had a median age of 34 years (range 18-70), 70% were male, 40% white, 23% black, 23% hispanic, and 13% asian. The mean baseline plasma HIV-1 RNA was 4.86 $\log_{10}$ copies/mL and the median baseline CD4+ cell count was 228 x 10^6 cells/L (range 4 – 750 x 10^6 cells/L).

Table 17: Demographic and Baseline Characteristics of Patients in TMC114-C211 Trial		
	Randomized TMC114-C211 Trial	
	PREZISTA®/rtv 800/100 mg q.d. + TDF/FTC N = 343	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N = 346**
Demographic Characteristics		
Median Age (years) (range, years)	34 (18-70)	33 (19-68)
Sex		
Male	70%	70%
Female	30%	30%
Race		
White	40%	45%
Black	23%	21%
Hispanic	23%	22%
Asian	13%	11%
Baseline Characteristics		
Mean Baseline Plasma HIV-1 RNA (log_{10} copies/mL)	4.86	4.84
Median Baseline CD4+ Cell Count (cells/mm^3) (range, cells/mm^3)	228 (4-750)	218 (2-714)
Percentage of Patients with Baseline Viral Load ≥ 100,000 copies/mL	34%	35%
Percentage of Patients with Baseline CD4+ Cell Count < 200 cells/mm^3	41%	43%

Study Results

According to the statistical methods for analysis at Week 48 and Week 192 in the TMC114-C211 protocol, if at each time point, the lower limit of the 95% two-sided CI of the difference between DRV/rtv and LPV/rtv exceeded -12% (for the On-Protocol (OP) population), noninferiority of DRV/rtv versus LPV/rtv could be concluded for that time point. Noninferiority in virologic response (HIV-1 RNA < 50 copies/mL) with PREZISTA®/rtv 800/100 mg q.d. compared to treatment with lopinavir/ritonavir 800/200 mg per day was demonstrated through 48 weeks of treatment (83.7% in the PREZISTA®/rtv 800/100 mg q.d arm versus 78.3% in the lopinavir/ritonavir 800/200 mg per day arm) ($p< 0.001$). The difference (95% CI) in response at Week 48 between DRV and LPV is 5.5% (-0.4 - 11.4) for <50 copies/mL and 2.7% (-2.4 - 7.8) for 400 copies/mL.

Analyses of the data at 192 weeks of treatment in the ARTEMIS trial demonstrated sustained antiretroviral efficacy and immunological benefit of the PREZISTA®/rtv arm. In the 192-week analysis, virologic response (HIV-1 RNA<50 copies/mL) in the ITT population was 68.8% and 57.2% for the PREZISTA®/rtv and lopinavir/rtv arm, respectively. Non-inferiority in virologic response was demonstrated ($p<0.001$) for both ITT and OP population. Furthermore, statistical superiority of the PREZISTA®/rtv arm over the lopinavir/rtv arm was demonstrated ($p=0.002$) for both ITT and OP population.

Week 48 and 192 week outcomes for patients in the On-Protocol population on PREZISTA®/rtv 800/100 mg q.d. from the ARTEMIS trial are shown in Table 18.

Table 18: Outcomes of Randomized Treatment Through Week 48 and 192 of the TMC114-C211 Trial for the On-Protocol Population

	Randomized Study TMC114-C211			
	Week 48		Week 192	
	PREZISTA®/ rtv 800/100 mg q.d. + TDF/FTC N = 340	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N = 346**	**PREZISTA®/rtv 800/100 mg q.d. + TDF/FTC N = 340**	**lopinavir/ritonavir 800/200 mg per day + TDF/FTC N = 346**
Virologic Responders HIV-1 RNA < 50 copies/mL	83.8%	78.3%	69.1%	57.1%
(HIV-1 RNA < 400 copies/mL)	(87.9%)	(85.3%)	(75.2%)	(65.0%)
Virologic failures	5.8%	10.1%	8.2%	12.5%
Rebounder[a]	1.5%	2.6%	7.0%	11.0%
Never suppressed[b]	4.4%	7.5%	1.2%	1.4%
Discontinuation due to adverse events	2.9%	5.5%	4.1%	11.3%
Death	0.3%	0.6%	0.3%	1.4%
Discontinuation due to other reasons	7.1%	5.5%	18.2%	17.7%

N = total number of patients in the On-Protocol population with data
[a] Patients with a confirmed viral load < 50 copies/mL before Week 48 or 192, but without a confirmed viral load < 50 copies/mL at Week 48 and Week 192
[b] Patients who never reached a confirmed viral load < 50 copies/mL before Week 48 or Week 192

The mean changes in plasma HIV-1 RNA from baseline in the ITT population were –2.77 $\log_{10}$ copies/mL at 48 weeks (-2.35 $\log_{10}$ copies/mL at 192 weeks) in the arm receiving PREZISTA®/rtv 800/100 mg q.d. and –2.65 $\log_{10}$ copies/mL at 48 weeks (-2.03 log10 copies/mL at 192 weeks) for the arm receiving lopinavir/ritonavir 800/200 mg per day. The median increase from baseline in CD4+ cell counts was comparable for both treatment groups (148 cells/mm^3 and 148 cells/mm^3 at 48 weeks (266 cells/mm^3 and 269 cells/mm^3 at 192 weeks) in the PREZISTA®/rtv 800/100 mg q.d. arm and the lopinavir/ritonavir 800/200 mg per day arm, respectively).

The virological response (< 50 copies/mL) by baseline viral load for all ITT patients is presented in Table 19. For patients with baseline VL < 100,000 copies/mL, responses were similar for PREZISTA®/rtv and lopinavir/ritonavir; patients with baseline VL ≥100,000 copies/mL receiving PREZISTA®/ritonavir had a statistically superior virological response (< 50 copies/mL) than lopinavir/rtv (67.5 % vs. 51.7%; p= 0.012).

Table 19: Virological Response (< 50 copies/mL) at 192 Weeks by BaselineViral Load and Baseline CD4+ Cell Count					
	PREZISTA®/rtv 800/100 mg q.d. n=343		lopinavir/ritonavir 800/200 mg per day n=346		Treatment difference
	N	number of responders at week 192 n (%)	N	number of responders at week 192 n (%)	Difference in % response (95% CI of difference in % response)
Baseline plasma viral load (copies/mL)					
< 100,000	226	157 (69.5)	226	136 (60.2)	9.3 (0.5; 18.1)
≥ 100,000	117	79 (67.5)	120	62 (51.7)	15.9 (3.5; 28.3)
Baseline CD4+ cell count (x 10^6/L)					
< 200	141	92 (65.2)	148	80 (54.1)	11.2 (-0.1; 22.5)
≥ 200	202	144 (71.3)	198	118 (59.6)	11.7 (2.4; 21.0)

Antiretroviral Treatment-Experienced Adult Patients

The evidence of comparable efficacy of PREZISTA®/rtv 800/100 mg q.d. and PREZISTA®/rtv 600/100 mg b.i.d. in treatment-experienced patients with no darunavir resistance associated mutations is based on the 48 week analysis of the Phase 3 trial TMC114-C229 (ODIN).

The evidence of efficacy of PREZISTA®/rtv 600/100 mg b.i.d. in treatment experienced patients is based on the 96 week analysis of the Phase 3 trial TMC114-C214 (TITAN) in treatment experienced, lopinavir/rtv naïve patients and on the analyses of 96 week data from the Phase 2b trials POWER 1, 2 and 3, in patients with high level of PI resistance.

TMC114-C229 (ODIN)

Demographics and Trial Design

Study TMC114-C229 is a randomized, open-label trial comparing PREZISTA®/rtv 800/100 mg q.d. to PREZISTA®/rtv 600/100 mg b.i.d. in treatment-experienced HIV-1-infected patients with screening genotype resistance test showing no darunavir resistance associated mutations (i.e. V11I, V32I, L33F, I47V, I50V, I54M, I54L, T47P, L76V, I84V, L89V) and a screening viral load of >1,000 HIV-1 RNA copies/mL. Both arms used an optimized background regimen consisting of ≥2 NRTIs selected by the investigator.

HIV-1-infected patients who were eligible for this trial were on a stable highly active antiretroviral therapy regimen (HAART) for at least 12 weeks. Virologic response was defined as a confirmed plasma HIV-1 RNA viral load <50 copies/mL. Analyses included 590 patients who had completed 48 weeks of treatment or discontinued earlier (ITT population). Table 20 compares the demographic and baseline characteristics between patients in the PREZISTA®/rtv 800/100 mg q.d. and patients in the PREZISTA®/rtv 600/100 mg b.i.d. arm in study TMC114-C229. No imbalances between the two arms were noted. The majority of the patient population (>87%) was not co-infected with hepatitis B and/or hepatitis C virus.

Table 20: Demographic and Baseline Characteristics of Patients in Study TMC114-C229 (ITT Population[a])		
	Randomized Study TMC114-C229	
	PREZISTA®/ritonavir 800/100 mg once daily + OBR N = 294	PREZISTA®/ritonavir 600/100 mg twice daily + OBR N = 296
Demographic Characteristics		
Median Age (years) (range, years)	40 (18-70)	40 (18-77)
Sex		
Male	61%	67%
Female	39%	33%
Race		
White	35%	37%
Black	28%	24%
Hispanic	16%	20%
Asian	16%	14%
Baseline Characteristics		
Mean Baseline Plasma HIV-1 RNA (log_{10} copies/mL)	4.19	4.13
Median Baseline CD4+ Cell Count (cells/mm^3) (range, cells/mm^3)	219 (24-1306)	236 (44-864)
Percentage of Patients with Baseline Viral Load ≥ 100,000 copies/mL	13%	11%
Percentage of Patients with Baseline CD4+ Cell Count < 200 cells/mm^3	43%	39%
Median Darunavir Fold Change (range)[b]	0.50 (0.1-1.8)	0.50 (0.1-1.9)
Median Number of Resistance-Associated[c]:		
PI mutations	3	4
NNRTI mutations	2	1
NRTI mutations	1	1
Percentage of Patients with Number of Baseline Primary Protease Inhibitor Mutations		
0	84%	84%
1	8%	9%
2	5%	4%
≥ 3	3%	2%
Median Number of ARVs Previously Used[d]:		
NRTIs	3	3
NNRTIs	1	1
PIs (excluding low-dose ritonavir)	1	1

[a] Intent-to-treat (ITT) population was defined as the set of patients who were randomized and who had taken ≥1 dose of trial medication, regardless of their compliance with the protocol.

[b] Based on phenotype [FC] (Antivirogram®) FC = $EC_{50}$of the patient virus/EC_{50} of reference wild type virus

[c] Johnson VA, Brun-Vézinet F, Clotet B, et al. Update of the drug resistance mutations in HIV-1: December 2008. Top HIV Med 2008; 16(5): 138-145

[d] Only counting ARVs, excluding low-dose ritonavir

Study Results

In the 48-week primary analysis, the virologic response defined as a confirmed plasma HIV-1 RNA viral load <50 copies/mL (ITT, TLOVR), was 72.1% for the PREZISTA®/rtv q.d. arm and 70.9% for the PREZISTA®/rtv b.i.d. arm (difference = 1.2%, 95% CI = [-6.1; 8.5]). Statistical comparisons between the treatment arms at Week 48 based on a normal approximation of the difference in virologic response, confirmed non-inferiority of PREZISTA®/rtv q.d. versus PREZISTA®/rtv b.i.d. for both the ITT and OP populations (p-value<0.001). A summary of all Week 48 outcomes for patients on PREZISTA®/rtv 800/100 mg q.d. from study TMC114-C229 are shown in Table 21.

Table 21: Outcomes of Randomized Treatment Through Week 48 of Study TMC114-C229

	Randomized Study TMC114-C229			
	PREZISTA®/rtv 800/100 mg q.d. + OBR N =294	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR N =296**	**Difference in virologic response [95% CI]**	**P-value for non-inferiority**
Virologic Responders (HIV-1 RNA < 50 copies/mL)	71.4% [a]	70.3% [a]	1.2 [-6.2; 8.5]	< 0.001 [b]
Virologic failures: [c] Lack of initial response [d]	10.5% [a]	8.1% [a]		
Rebounder [e]	2.4%	2.0%		
Never suppressed [f]	4.8%	5.1%		
Discontinuation due to adverse events	2.4%	2.0%		
Death	0.6%	1.7%		
Discontinuation due to other reasons [g]	7.8%	10.8%		

N = total number of patients in the ITT population with data

[a] Two patients in each arm with a confirmed virologic response at Week 48 demonstrated an initial lack of response at week 24, and were therefore not considered virologic responders in this outcome summary.

[b] Based on a normal approximation of the difference in response.

[c] Patients who discontinued prior to Week 48 for lack or loss of efficacy; patients who are ≥50 copies in Week 48 and patients who had a switch in their background regimen that was not permitted by the protocol.

[d] Patients are considered to show an initial lack of response if i) they have viral load data beyond the considered time point, and ii) confirmed virologic response (VL<400 copies/mL – TLOVR) at Week 24 equals 0.

[e] Patients who have a confirmed response (<50 copies/mL – TLOVR) before the considered time point (any response = 1) but who have a confirmed rebound at the considered time point (i.e. TLOVR response = No at the considered time point)

[f] Patients who never reached a confirmed viral load < 50 copies/mL before Week 48

[g] Includes: withdrew consent, loss to follow-up, moved etc.

At Week 48, the mean change in $\log_{10}$ viral load from baseline in the ITT population was -1.84 and -1.80 $\log_{10}$ copies/mL for the DRV/rtv q.d. and DRV/rtv b.i.d. groups, respectively. The difference in mean change and 95% confidence interval in $\log_{10}$ viral load from baseline between

treatment groups at Week 48 was -0.04 (-0.24, 0.16). Statistical comparison (ITT-ANCOVA) showed an estimated difference (difference between LS Means and the 95% CI) between the DRV/rtv q.d. and the b.i.d. treatment groups at 48 weeks of -0.003 (-0.188, 0.182) (p=0.977).

The mean increase from baseline in CD4+ cell counts was comparable for both treatment arms (108 cells/mm^3 and 112 cells/mm^3 in the PREZISTA®/rtv 800/100 mg q.d. arm and the PREZISTA®/rtv 600/100 mg b.i.d. arm, respectively).

Study TMC114-C214 (TITAN)

Demographics and Trial Design
Study TMC114-C214 was an ongoing randomized, controlled, open label Phase 3 trial comparing PREZISTA®/rtv 600/100 mg b.i.d. versus lopinavir/rtv 400/100 mg b.i.d. in antiretroviral treatment-experienced, lopinavir/rtv naïve HIV-1 infected adult patients. Both arms used an optimized background regimen (OBR) consisting of at least 2 antiretrovirals (NRTIs with or without NNRTIs).

HIV-1 infected patients who were eligible for this trial had plasma HIV-1 RNA > 1000 copies/mL and were on a highly active antiretroviral therapy regimen (HAART) for at least 12 weeks. Virologic response was defined as a confirmed plasma HIV-1 RNA viral load < 400 copies/mL. Analyses included 595 patients in the TITAN trial who had completed 96 weeks of treatment or discontinued earlier.

Demographics and baseline characteristics were balanced between the PREZISTA®/rtv arm and the lopinavir/ritonavir arm (see Table 22). Table 22 compares the demographic and baseline characteristics between patients in the PREZISTA®/rtv 600/100 mg b.i.d. arm and patients in the lopinavir/ritonavir 400/100 mg b.i.d. arm in study TMC114-C214.

Table 22: Demographic and Baseline Characteristics of Patients in Study TMC114-C214		
	Randomized Study TMC114-C214	
	PREZISTA®/rtv 600/100 mg b.i.d. + OBR N = 298	**lopinavir/ritonavir 400/100 mg b.i.d. + OBR N = 297**
Demographic Characteristics		
Median Age (years) (range, years)	40 (18-68)	41 (22-76)
Sex		
Male	77%	81%
Female	23%	19%
Race		
White	54%	57%
Black	18%	17%
Hispanic	15%	15%
Asian	9%	9%
Baseline Characteristics		
Mean Baseline Plasma HIV-1 RNA (log_{10} copies/mL)	4.33	4.28
Median Baseline CD4+ Cell Count (cells/mm^3) (range, cells/mm^3)	235 (3-831)	230 (2-1096)
Percentage of Patients with Baseline Viral Load ≥ 100,000 copies/mL	19%	17%
Percentage of Patients with Baseline CD4+ Cell Count < 200 cells/mm^3	40%	40%
Median Darunavir FC	0.60 (0.1 - 37.4)	0.60 (0.1 - 43.8)
Median Lopinavir FC	0.70 (0.4 – 74.4)	0.80 (0.3 – 74.5)
Median Number of Resistance-Associated[a]: PI mutations NNRTI mutations NRTI mutations	 4 1 2	 4 1 2
Percentage of Patients with Number of Baseline Primary Protease Inhibitor Mutations[a]: ≤ 1 2 ≥ 3	 78% 8% 13%	 80% 9% 11%
Median Number of ARVs Previously Used[b]: NRTIs NNRTIs PIs (excluding low-dose ritonavir)	 4 1 1	 4 1 1
Percentage of Patients Resistant[c] to All Available[d] PIs at Baseline, excluding Darunavir	2%	3%

[a] Johnson VA, Brun-Vezinet F, Clotet B, et al. Update of the drug resistance mutations in HIV-1: Fall 2006. Top HIV Med 2006; 14(3): 125-130

[b] Only counting ARVs, excluding low-dose ritonavir

[c] Based on phenotype (Antivirogram™)

[d] Commercially available PIs at the time of study enrollment

Study Results

According to the statistical methods in the TMC114-C214 protocol, if the lower limit of the 95% 2-sided CI of the difference between DRV/rtv and LPV/rtv exceeded –12% (for the On-protocol (OP) population), noninferiority of DRV/rtv versus LPV/rtv was concluded. Week 48 and 96 outcomes for patients on PREZISTA®/rtv 600/100 mg b.i.d. from study TMC114-C214 are shown in Table 23.

Table 23: Outcomes of Randomized Treatment Through Week 48 and 96 of Study TMC114-C214

	Randomized Study TMC114-C214			
	48 Weeks		**96 Weeks**	
	PREZISTA®/rtv 600/100 mg b.i.d. + OBR N = 274 N	**lopinavir/ritonavir 400/100 mg b.i.d. + OBR N = 280 N**	**PREZISTA®/rtv 600/100 mg b.i.d. + OBR N = 280 N**	**lopinavir/ritonavir 400/100 mg b.i.d. + OBR N = 294 N**
Virologic Responders HIV-1 RNA < 400 copies/mL (HIV-1 RNA < 50 copies/mL)	211 (77.0%) [196 (71.5%)]	189 (67.5%) [169 (60.4%)]	189 (67.5%) [172 (61.4%)]	175 (59.5%) [164 (55.8%)]
Virologic failures	28 (10.2%)	51 (18.2%)	31 (11.1%)	61 (20.7%)
Lack of initial response[a]	19 (6.9%)	29 (10.4%)	19 (6.8%)	35 (11.9%)
Rebounder[b]	9 (3.3%)	21 (7.5%)	12 (4.3%)	25 (8.5%)
Discontinued due to virologic failure: never suppressed[c]	0 (0%)	1 (0.4%)	0 (0%)	1 (0.3%)
Discontinuation due to adverse events	14 (5.1%)	13 (4.6%)	22 (7.9%)	23 (7.8%)
Death	2 (0.7%)	2 (0.7%)	20 (7.1%)	17 (5.8%)
Discontinuation due to other reasons	19 (6.9%)	25 (8.9%)	2 (0.7%)	3 (1.0%)

N = total number of patients with data

[a] Patients with viral load ≥ 400 copies/mL at Week 16 and without a confirmed viral load <400 copies/mL at Week 48 and Week 96

[b] Patients with a confirmed viral load < 400 copies/mL before Week 48 and Week 96, but without a confirmed viral load < 400 copies/mL at Week 48 and Week 96

[c] Patients who never reached a confirmed viral load < 400 copies/mL before Week 48 and Week 96

Through 96 weeks of treatment, there was a significantly greater proportion of patients with HIV-1 RNA < 400 copies/mL and with HIV-1 RNA < 50 copies/mL in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. (67.5% and 61.4%, respectively) compared to the arm receiving lopinavir/ritonavir 400/100 mg b.i.d. (59.5% and 55.8%, respectively). The difference (95% CI) in response at Week 96 between DRV and LPV is 8.0% (0.1 - 15.8) for <400 copies/mL and 5.6% (2.4 – 13.7) for <50 copies/mL.

Noninferiority in virologic response (HIV-1 RNA < 400 copies/mL) with PREZISTA®/rtv 600/100 mg b.i.d. compared to treatment with lopinavir/ritonavir 400/100 mg b.i.d. was demonstrated ($p < 0.001$), furthermore superiority of PREZISTA®/rtv over the lopinavir/rtv arm was demonstrated ($p = 0.033$).

The proportion of patients with at least 1 $\log_{10}$ HIV-1 RNA below baseline was 77.7% in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. compared to 69.3% in the arm receiving lopinavir/ritonavir 400/100 mg b.i.d. At week 96, the mean changes in plasma HIV-1 RNA from baseline were -1.72 $\log_{10}$ copies/mL in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. and -1.54 $\log_{10}$ copies/mL for the arm receiving lopinavir/ritonavir 400/100 mg b.i.d. The median increase from baseline in CD4+ cell counts was comparable for both treatment groups (81 cells/mm^3 and 96 cells/mm^3 in the PREZISTA®/ rtv 600/100 mg b.i.d. arm and lopinavir/ritonavir 400/100 mg b.i.d. arm, respectively).

TMC114-C213 (POWER 1) and TMC114-C202 (POWER 2)

Demographics and Trial Design
Studies TMC 114-C213 (Power 1) and TMC114-C202 (Power 2) are randomized, controlled Phase 2b trials in patients with a high level of PI resistance, consisting of 2 parts: an initial partially blinded, dose finding part and a second long term part in which all patients randomized to PREZISTA®/rtv received the recommended dose of 600/100 mg b.i.d.

HIV-1-infected patients who were eligible for these trials had plasma HIV-1 RNA > 1000 copies/mL, had prior treatment with PI(s), NNRTI(s) and NRTI(s), had at least one primary PI mutation (D30N, M46I/L, G48V, I50L/V, V82A/F/S/T, I84A/C/V, L90M) at screening, and were on a stable PI-containing regimen at screening for at least 8 weeks. Randomization was stratified by the number of PI mutations, screening viral load, and the use of enfuvirtide.

The virologic response rate was evaluated in patients receiving PREZISTA®/rtv plus an optimized background regimen (OBR) versus a control group receiving an investigator-selected PI(s) regimen plus an OBR. The OBR consisted of at least 2 NRTIs with or without enfuvirtide. Based on resistance testing and prior medical history, selected PIs in the control arm included: lopinavir/ritonavir in 36%, (fos)amprenavir in 34%, saquinavir in 35% and atazanavir in 17%; 23% of the control patients used dual-boosted PIs. Approximately 47% of all patients used enfuvirtide (ENF), and 35% of the use was in patients who were ENF-naïve. Virologic response was defined as a decrease in plasma HIV-1 RNA viral load of at least 1.0 $\log_{10}$ versus baseline.

In the pooled analysis for TMC114-C213 and TMC114-C202, demographics and baseline characteristics were balanced between the PREZISTA®/rtv arm and the comparator PI arm. Table 24 compares the demographic and baseline characteristics between patients in the PREZISTA®/rtv 600/100 mg b.i.d. arm and patients in the comparator PI arm in the pooled analysis of studies TMC114-C213 and TMC114-C202.

Table 24: Demographic and Baseline Characteristics of Patients in Studies TMC114-C213 and TMC114-C202 (Pooled Analysis)		
	Randomized Studies TMC114-C213 and TMC114-202	
	PREZISTA®/rtv 600/100 mg b.i.d. + OBR N = 131	**Comparator PI(s) + OBR N = 124**
Demographic Characteristics		
Median Age (years) (range, years)	43 (27-73)	44 (25-65)
Sex		
Male	89%	88%
Female	11%	12%
Race		
White	81%	73%
Black	10%	15%
Hispanic	7%	8%
Baseline Characteristics		
Mean Baseline Plasma HIV-1 RNA (log_{10} copies/mL)	4.61	4.49
Median Baseline CD4+ Cell Count (cells/mm^3) (range, cells/mm^3)	153 (3-776)	163 (3-1274)
Percentage of Patients with Baseline Viral Load > 100,000 copies/mL	24%	29%
Percentage of Patients with Baseline CD4+ Cell Count < 200 cells/mm^3	67%	58%
Median Darunavir FC	4.3	3.3
Median Number of Resistance-Associated[a]:		
PI mutations	12	12
NNRTI mutations	1	1
NRTI mutations	6	5
Percentage of Patients with Number of Baseline Primary Protease Inhibitor Mutations[a]:		
≤ 1	8%	9%
2	22%	21%
≥ 3	70%	70%
Median Number of ARVs Previously Used[b]:		
NRTIs	6	6
NNRTIs	1	1
PIs (excluding low-dose ritonavir)	4	4
Percentage of Patients Resistant[b] to All Available[c] PIs at Baseline, excluding Tipranavir and Darunavir	66%	61%
Percentage of Patients with Prior Use of Enfuvirtide	20%	17%

[a] Johnson VA, Brun-Vezinet F, Clotet B, et al. Update of the drug resistance mutations in HIV-1: Fall 2006. Top HIV Med 2006; 14(3): 125-130

[b] Based on phenotype (Antivirogram™)

[c] Commercially available PIs at the time of study enrollment

Study Results

Week 48 outcomes for patients on the recommended dose PREZISTA®/rtv 600/100 mg b.i.d. from the pooled studies TMC114-C213 and TMC114-C202 are shown in Table 25.

Table 25: Outcomes of Randomized Treatment Through Week 48 of the Studies TMC114-C213 and TMC114-C202 (Pooled Analysis)

	Randomized Studies TMC114-C213 and TMC114-C202	
	PREZISTA®/rtv 600 mg b.i.d. + OBR N=131	**Comparator PI + OBR N=124**
Virologic Responders confirmed at least 1 $\log_{10}$ HIV-1 RNA below baseline through Week 48 (< 50 copies/mL at Week 48)	61.1%[d] (45.0%)[d]	16.1% (11.3%)
Virologic failures	29.0%	75.0%
Lack of initial response[a]	8.4%	53.2%
Rebounder[b]	16.0%	13.7%
Never Suppressed[c]	4.6%	8.1%
Discontinuation due to adverse events	4.6%	2.4%
Deaths	2.3%	0.8%
Discontinuation due to other reasons	3.1%	5.6%

[a] Patients who did not achieve at least a confirmed 0.5 $\log_{10}$ HIV-1 RNA drop from baseline at Week 12

[b] Patients with an initial response (confirmed 1 $\log_{10}$ drop in viral load), but without a confirmed 1 $\log_{10}$ drop in viral load at Week 48

[c] Patients who never reached a confirmed 1 $\log_{10}$ drop in viral load before Week 48

[d] p<0.001, based on logistic regression model; p-values [PREZISTA®/rtv 600/100 mg b.i.d vs. comparator PI + OB]

In the pooled TMC114-C213 and TMC114-C202 analysis through 48 weeks of treatment, the proportion of patients with HIV-1 RNA < 400 copies/mL in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. compared to the comparator PI arm was 55.0% and 14.5%, respectively (p<0.001). In addition, the mean changes in plasma HIV-1 RNA from baseline were –1.69 $\log_{10}$ copies/mL in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. and –0.37 $\log_{10}$ copies/mL for the comparator PI arm (p<0.001). The mean increase from baseline in CD4+ cell counts was higher in the arm receiving PREZISTA®/rtv 600/100 mg b.i.d. (103 cells/mm^3) than in the comparator PI arm (17 cells/mm^3) (p<0.001).

Analyses of the data through 96 weeks of treatment in the pooled TMC114-C213 and TMC114-C202 trials demonstrated sustained antiretroviral efficacy and immunological benefit. Treatment with PREZISTA®/rtv 600/100 mg b.i.d. resulted in 56.5% of patients with a decrease of at least 1 $\log_{10}$ HIV-1 RNA versus baseline and 38.9% of patients reaching less than 50 HIV-1 RNA copies/mL. At Week 96, 49.6% of patients reached less than 400 HIV-1 RNA copies/mL, and the mean change in plasma HIV-1 RNA from baseline was -1.58 $\log_{10}$ copies/mL. The mean increase in CD4+ cell count versus baseline was 133 cells/mm^3 at Week 96.

TMC114-C215 (POWER 3)

Demographics and Trial Design
Additional data on the efficacy of PREZISTA®/rtv 600/100 mg b.i.d. have been obtained in treatment-experienced patients participating in the non-randomized trial TMC114-C215. The 318 patients included in the 96-week efficacy analysis initiated therapy with PREZISTA®/rtv with the recommended dose of 600/100 mg b.i.d. The OBR consisted of at least two NRTIs with or without enfuvirtide. Entry criteria for Study TMC114-C215 were the same as those for TMC114-C213 (POWER 1) and TMC114-C202 (POWER 2) trials.

Baseline characteristics of the patients included in TMC114-C215 trial were comparable to those patients in TMC114-C213 and TMC114-C202 trials.

Study Results
The TMC114-C215 48-week efficacy analysis supported the viral load reduction and CD4+ cell count increases observed in the TMC114-C213 and TMC114-C202 trials. Of the 334 patients at Week 48, 58.7% had a virologic response defined as a decrease of at least 1.0 $\log_{10}$ HIV-1 RNA versus baseline and 46.4% of the patients reached less than 50 HIV-1 RNA copies/mL. At Week 48, 54.8% of the patients reached less than 400 HIV-1 RNA copies/mL, and the mean changes in plasma HIV-1 RNA from baseline were -1.62 $\log_{10}$ copies/mL. The mean increase in CD4+ cell count versus baseline was 105 cells/mm^3 at Week 48.

Analyses of the data through 96 weeks of treatment in the non-randomized trial TMC114-C215 demonstrated sustained antiretroviral efficacy and immunological benefit**.** Of the 318 patients at Week 96, 52.2% had a virologic response defined as a decrease of at least 1.0 $\log_{10}$ HIV-1 RNA versus baseline and 42.1% of the patients reached less than 50 HIV-1 RNA copies/mL. At Week 96, 50.0% of the patients reached less than 400 HIV-1 RNA copies/mL, and the mean change in plasma HIV-1 RNA from baseline was -1.43 $\log_{10}$ copies/mL. The mean increase in CD4+ cell count versus baseline was 103 cells/mm^3 at Week 96.

TMC114-C213, TMC114-C202 and TMC114-C215 Trials
Out of the 206 patients who responded with complete viral suppression (< 50 HIV-1 RNA copies/mL) at Week 48, 86% of patients remained responders at Week 96.

Pediatric Patients

Treatment-Experienced Pediatric Patients

The evidence of efficacy of PREZISTA®/rtv in antiretroviral treatment-experienced pediatric patients is based on two Phase 2 trials.

TMC114-C212 (DELPHI)

Demographics and Trial Design
Study TMC114-C212 (DELPHI) is an open-label, Phase 2 trial of 48 week duration evaluating the pharmacokinetics, safety, tolerability, and efficacy of PREZISTA®/rtv in 80 antiretroviral

treatment-experienced HIV-1-infected pediatric patients 6 to < 18 years of age and weighing at least 44 lbs (20 kg).

At Week 24, the virologic response rate was evaluated in pediatric patients receiving PREZISTA®/rtv in combination with other antiretroviral agents (see **DOSAGE AND ADMINISTRATION** for dosage recommendations per body weight). Virologic response was defined as a decrease in plasma HIV-1 RNA viral load of at least 1 $\log_{10}$ versus baseline. The mean baseline plasma HIV-1 RNA was 4.64 $\log_{10}$ copies/mL, and the median baseline CD4+ cell count was 330 cells/mm^3 (range: 6 to 1505 cells/mm^3).

In the study, pediatric patients who were at risk of discontinuing therapy due to intolerance of ritonavir oral solution (e.g., taste aversion) were allowed to switch to the capsule formulation. Of the 44 pediatric patients taking ritonavir oral solution, 23 patients switched to the 100 mg capsule formulation and exceeded the weight-based ritonavir dose without changes in observed safety.

Study Results
At Week 24, 73.8% of pediatric patients had at least 1 $\log_{10}$ HIV-1 RNA decrease from baseline. The proportion of pediatric patients reaching undetectable viral load (< 50 HIV-1 RNA copies/mL) was 50.0%, and the proportion of pediatric patients with < 400 HIV-1 RNA copies/mL was 63.8%. The mean change in plasma HIV-1 RNA from baseline was -1.98 $\log_{10}$ copies/mL. The mean CD4+ cell count increase from baseline was 117 cells/mm^3 and the median CD4+ cell count increase was 96 cells/mm^3 (range: -232 to 465 cells/mm^3).

TMC114-C228 (ARIEL)

Demographics and Trial Design
Study TMC114- C228 (ARIEL) is an open-label, Phase 2 trial that evaluated the pharmacokinetics, safety, tolerability and efficacy of PREZISTA® oral suspension with low dose ritonavir in 21 antiretroviral treatment-experienced HIV-1-infected pediatric patients aged 3 to < 6 years and weighing ≥ 10 kg to < 20 kg.

At week 24, the virologic response rate was evaluated in pediatric patients receiving PREZISTA®/rtv in combination with other antiretroviral agents (see **DOSAGE AND ADMINISTRATION** for dosage recommendations per body weight). Final analysis was performed at week 48. Virologic response was defined as a decrease in plasma viral load to < 50 HIV-1 RNA copies/mL. The mean baseline plasma HIV-1 RNA was 4.34 $\log_{10}$ copies/mL, the median baseline CD4+ cell count was 927 x 10^6 cells/L (range: 209 to 2,429 x 10^6 cells/L) and the median baseline CD4+ percentage was 27.7% (range: 15.6% to 51.1%).

Study Results
The proportions of patients reaching undetectable viral load (< 50 HIV-1 RNA copies/mL) at weeks 24 and 48 were 57.1% and 81.0%, respectively. At week 48, the proportion of patients with < 400 HIV-1 RNA copies/mL was 85.7%. A ≥ 1.0 $\log_{10}$ HIV-1 RNA decrease from baseline was achieved in 90.5% of patients. The mean change in plasma HIV-1 RNA from

baseline was -2.14 $\log_{10}$ copies/mL. The mean CD4+ cell count increase and mean change in CD4+ percentage from baseline was 187 x 10^6 cells/L and 4%, respectively.

Pivotal Comparative Bioavailability Studies

TMC114-TiDP3-C162

In a Phase 1, open-label, randomized, two-panel, two-way crossover bioavailability trial the rate and extent of absorption of darunavir following administration of two tablet strengths (in the presence of low-dose ritonavir) under fed and fasted conditions was assessed in 96 healthy subjects.

In Panel 1, 47 male and female subjects randomly received under fasted conditions a single oral 600 mg dose of darunavir formulated as the 300 mg tablet (2 x 300 mg; Treatment A) in one session, followed by a single oral 600 mg dose formulated as the 600 mg tablet (1 x 600 mg; Treatment B) in the second session. The results indicate that the bioavailability of a 1x 600 mg dose of darunavir is comparable to the bioavailability of a 2 x 300 mg dose of darunavir.

In Panel 2, 46 male and female subjects randomly received under fed conditions a single oral 600 mg dose of darunavir formulated as the 300 mg tablet (2 x 300 mg; Treatment C) in one session and a single oral 600 mg dose of darunavir formulated as the 600 mg tablet (1 x 600 mg; Treatment D) in the second session. The results indicate that the bioavailability of a 1x 600 mg dose of darunavir is comparable to the bioavailability of a 2 x 300 mg dose of darunavir.

The summary of results is presented in Table 26.

Table 26: Summary Table of the Comparative Bioavailability Data Under Fed and Fasting Conditions

Darunavir (TMC114)
1 x 600 mg tablet and 2 x 300 mg tablet
From measured data

Geometric Mean[a]
Arithmetic Mean (CV%)

	Fed Conditions			Fasted Conditions		
Parameter	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval)	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval)
AUC_{last} (ng.h/mL)	103900 111800 (44.3)	99650 103900 (32.4)	95.87 (89.67 - 102.5)	81250 85470 (31.5)	78140 82500 (34.0)	96.17 (89.75 - 103.0)
AUC_{∞} (ng.h/mL)	108600 117300 (45.7)	105000 110600 (36.5)	96.66 (90.35 - 103.4)	87870 93700 (36.5)	85000 92440 (43.6)	96.73 (89.35 - 104.7)

Table 26: Summary Table of the Comparative Bioavailability Data Under Fed and Fasting Conditions						
Darunavir (TMC114) 1 x 600 mg tablet and 2 x 300 mg tablet From measured data Geometric Mean[a] Arithmetic Mean (CV%)						
Fed Conditions				Fasted Conditions		
Parameter	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval)	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval)
C_{max} (ng/mL)	5843 6024 (26.6)	5706 5803 (18.6)	97.66 (92.84 - 102.7)	4056 4134 (19.7)	4076 4213 (28.0)	100.5 (94.31 - 107.1)
T_{max}[d] (h)	4.0 (1.0 - 5.0)	4.0 (1.0 - 6.0)		2.0 (1.0 - 5.0)	2.0 (1.0 - 5.0)	
$T_{½}$[e] (h)	15.81 (33.4)	15.94 (42.9)		19.09 (38.0)	18.97 (58.5)	

[a] Based on least square mean estimates
[b] Darunavir 300 mg tablet (F016)
[c] Darunavir 600 mg tablet (F032)
[d] Expressed arithmetic median (range) only
[e] Expressed as the arithmetic mean (CV%) only

TMC114-TiDP3-C176

In a Phase 1, open-label, randomized two-panel, two-way crossover bioavailability trial the rate and extent of absorption of darunavir following administration of two tablet strengths (in the presence of low-dose ritonavir) under fed and fasted conditions was assessed in 124 healthy subjects.

In Panel 1, 78 male and female subjects randomly received under fasted conditions a single oral 800 mg dose of darunavir formulated as the 400 mg tablet (2 x 400 mg; Treatment A) in one session, followed by a single oral 800 mg dose formulated as the 800 mg tablet (1 x 800 mg; Treatment B) in the second session. The results indicate that the bioavailability of a 1 x 800 mg dose of darunavir is comparable to the bioavailability of a 2 x 400 mg dose of darunavir.

In Panel 2, 40 male and female subjects randomly received under fed conditions a single oral 800 mg dose of darunavir formulated as the 400 mg tablet (2 x 400 mg; Treatment C) in one session, followed by a single oral 800 mg dose formulated as the 800 mg tablet (1 x 800 mg; Treatment D) in the second session. The results indicate that the bioavailability of a 1 x 800 mg dose of darunavir is comparable to the bioavailability of a 2 x 400 mg dose of darunavir.

The summary of results is presented in Table 27.

Table 27: Summary Table of the Comparative Bioavailability Data Under Fed and Fasting Conditions						
Darunavir (TMC114) 1 x 800 mg tablet and 2 x 400 mg tablet From measured data Geometric Mean[a] Arithmetic Mean (CV%)						
Fed Conditions				Fasted Conditions		
Parameter	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval	Reference[b]	Test[c]	% Ratio of Geometric Means[a] (90% Confidence Interval
AUC_{last} (ng.h/mL)	101800 105900 (30.72)	99350 105100 (36.04)	97.59 (93.82 – 101.51)	86000 96120 (57.04)	85300 91140 (44.16)	99.18 (94.35 - 104.27)
AUC_{∞} (ng.h/mL)	105100 109700 (32.32)	103100 109600 (37.91)	98.12 (94.11– 101.31)	92420 105000 (62.36)	92720 99540 (49.60)	100.33 (94.80 - 106.19)
C_{max} (ng/mL)	6890 7031 (23.74)	6580 6773 (24.63)	95.50 (92.15 – 98.97)	4658 4866 (29.62)	4750 4914 (27.10)	101.97 (98.03 - 106.07)
T_{max}[d] (h)	2.98 (1.00 – 5.97)	2.98 (0.97 – 5.00)		2.02 (0.97 - 23.93)	2.00 (1.00 - 4.98)	
$T_{½}$[e] (h)	13.45 (35.47)	14.03 (32.94)		16.09 (52.05)	16.96 (101.30)	

[a] Based on least square mean estimates
[b] Darunavir 400 mg tablet (F030)
[c] Darunavir 800 mg tablet (G002)
[d] Expressed as the median (range) only
[e] Expressed as the mean (CV%) only

TMC114-TiDP3-C169

In a Phase 1, open-label, randomized, three-way crossover bioavailability trial the rate and extent of absorption of darunavir following administration of two different formulations (tablets or suspension) (in the presence of low-dose ritonavir) under fed and fasted conditions was assessed in 20 healthy subjects.

Seventeen male and female subjects randomly received a single oral 600 mg dose of darunavir formulated as the 300 mg tablet (2 x 300 mg; Treatment A) under fed conditions, followed by a single oral 600 mg dose formulated as the 100 mg/mL suspension (6 x 100 mg/mL; Treatment B) under fasted conditions, followed by a single oral 600 mg dose of darunavir formulated as the 100 mg/mL suspension (6 x 100 mg/mL; Treatment C) under fed conditions. The results indicate that the bioavailability of a 2 x 300 mg dose of darunavir under fed conditions is comparable to the bioavailability of a 6 mL x 100 mg/mL dose of darunavir under fasted or fed conditions.

The summary of results for the comparison between the 300 mg tablet and the 100 mg/mL suspension under fed conditions is presented in Table 28.

Table 28: Summary Table of the Comparative Bioavailability Data Under Fed Conditions

Darunavir (TMC114)
6 mL x 100 mg/mL and 2 x 300 mg tablet
From measured data

Geometric Mean[a]
Arithmetic Mean (CV%)

Fed conditions				
Parameter	Reference[b]	Test[c]	% Ratio of Geometric Means[a]	90% Confidence Interval
AUC_{last} (ng.h/mL)	77480 85240 (44.61%)	83320 88410 (36.86%)	107.5	101.1 - 114.4
AUC_{∞} (ng.h/mL)	81270 87330 (46.82%)	86940 92270 (36.35%)	107.0	100.3 - 114.1
C_{max} (ng/mL)	5434 5654 (26.13%)	5676 5885 (29.29%)	104.4	99.38 - 109.8
T_{max}[d] (h)	3.0 (2.5 – 5.0)	4.0 (1.5 – 4.0)		
$T_{½}$[e] (h)	15.04 (52.42%)	15.36 (41.93%)		

[a] Based on least square mean estimates
[b] Darunavir 300 mg tablet (F016)
[c] Darunavir 100 mg/mL suspension (F051)
[d] Expressed as the median (range) only
[e] Expressed as the mean (CV%) only

DETAILED PHARMACOLOGY

Pharmacodynamics

Electrocardiogram
See ***Product Monograph Part I*: ACTION AND CLINICAL PHARMACOLOGY**, **Pharmacodynamics**.

Pharmacokinetics

See ***Product Monograph Part I*: ACTION AND CLINICAL PHARMACOLOGY**; **Pharmacokinetics Absorption and Bioavailability**

Absorption
Maximum plasma concentration of darunavir in the presence of low-dose ritonavir is generally achieved within 2.5-4.0 hours. The absolute oral bioavailability of a single 600 mg dose of

darunavir alone was approximately 37% and increased to approximately 82% in the presence of 100 mg b.i.d. ritonavir. Increasing the dose of ritonavir to above 100 mg b.i.d. did not significantly affect darunavir concentrations. It is not recommended to alter the dose of ritonavir.

The pharmacokinetic properties of darunavir, co-administered with ritonavir, have been evaluated in healthy adult volunteers and in HIV-1-infected patients. Exposure to darunavir was higher in HIV-1-infected patients than in healthy patients. Darunavir is primarily metabolized by CYP3A. Ritonavir inhibits CYP3A, thereby increasing the plasma concentrations of darunavir considerably.

Table 29 displays the mean plasma concentrations of darunavir at steady-state for the darunavir/ritonavir 800/100 mg q.d. dose.

Table 29: Mean Steady-State Plasma Concentration-Time Profiles of Darunavir and Ritonavir at 800/100 mg q.d. at Week 4 (Study TMC114-C211)

Scheduled Time	Darunavir		Ritonavir	
	Mean ± SD (ng/mL)	CV (%)	Mean ± SD (ng/mL)	CV (%)
0 h	1826 ±1003	54.92	141.2± 156.0	110.5
1 h	3964 ± 1805	45.52	154.4± 122.6	79.36
2 h	4692 ± 1135	24.19	264.5± 247.2	93.46
3 h	4949 ± 1344	27.15	386.8± 357.6	92.45
4 h	4426 ± 1300	29.38	465.9± 256.5	55.06
6 h	3532 ± 1065	30.16	463.1± 205.3	44.33
9 h	2664 ± 1002	37.59	282.7± 137.5	48.64
12 h	2353 ± 919.6	39.08	229.4± 159.8	69.67
24 h	1440 ±513.9	35.68	97.10±99.08	102.0

Table 30 displays the mean plasma concentrations of darunavir and ritonavir at steady-state for the darunavir/ritonavir 600/100 mg b.i.d. dose.

Table 30: Mean Steady-State Plasma Concentration-Time Profiles of Darunavir and Ritonavir at 600/100 mg b.i.d. at Week 4 (Integrated Data from POWER 1 and POWER 2, Primary 24-Week Analysis)

Scheduled Time	Darunavir		Ritonavir	
	Mean ± SD (ng/mL)	CV (%)	Mean ± SD (ng/mL)	CV (%)
0 h	4010 ± 1635	40.78	495.6 ± 258.6	52.18
1 h	5386 ± 1538	28.55	634.0 ± 462.2	72.90
2 h	6125 ± 1750	28.57	658.1 ± 439.7	66.81
3 h	6471 ± 2066	31.92	713.8 ± 371.5	52.04
4 h	5307 ± 1796	33.84	719.5 ± 274.7	38.18
6 h	4856 ± 2007	41.33	701.9 ± 318.3	45.36
9 h	3618 ± 1893	52.33	516.9 ± 224.4	43.42
12 h	2813 ± 1612	57.30	321.9 ± 180.1	55.97

Effect of food on oral absorption
When administered without food, the relative bioavailability of darunavir in the presence of low-dose ritonavir is 30% lower as compared to intake with food. Therefore, to achieve optimal exposure, PREZISTA® tablets should be taken with ritonavir and with food. The type of food does not affect exposure to darunavir. (See **DOSAGE AND ADMINISTRATION** and **DRUG INTERACTIONS, Drug-Food Interactions**).

Distribution
Darunavir is approximately 95% bound to plasma proteins. Darunavir binds primarily to plasma alpha-1-acid glycoprotein (AAG).

Metabolism
In vitro experiments with human liver microsomes indicate that darunavir primarily undergoes oxidative metabolism. Darunavir is extensively metabolized by the hepatic CYP system, and almost exclusively by isozyme CYP3A4. At least 3 oxidative metabolites of darunavir have been identified in humans; all showed activity that was at least 10-fold less than the activity of darunavir against wild-type HIV.

Elimination
The terminal elimination half-life of darunavir was approximately 15 hours when combined with ritonavir. After administration of ^{14}C-darunavir with low-dose ritonavir, approximately 79.5% and 13.9% of the administered dose of ^{14}C -darunavir could be retrieved in feces and urine, respectively.

Drug-Drug Interactions
See also **CONTRAINDICATIONS**, **WARNINGS AND PRECAUTIONS** and **DRUG INTERACTIONS**.

Darunavir co-administered with ritonavir is an inhibitor of CYP3A and CYP2D6. Co-administration of darunavir and ritonavir with drugs primarily metabolized by CYP3A and CYP2D6 may result in increased plasma concentrations of such drugs, which could increase or prolong the therapeutic effect and adverse events.

Darunavir and ritonavir are metabolized by CYP3A. Drugs that induce CYP3A activity would be expected to increase the clearance of darunavir and ritonavir, resulting in lowered plasma concentrations of darunavir and ritonavir. Co-administration of darunavir and ritonavir and other drugs that inhibit CYP3A may decrease the clearance of darunavir and ritonavir and may result in increased plasma concentrations of darunavir and ritonavir.

Drug interaction studies were performed with darunavir and other drugs likely to be co-administered and some drugs commonly used as probes for pharmacokinetic interactions. The effects of co-administration of darunavir on the AUC, C_{max} and C_{min} values are summarized in Table 31 (effect of other drugs on darunavir) and Table 32 (effect of darunavir on other drugs) (see **ACTION AND CLINICAL PHARMACOLOGY**). For information regarding clinical recommendations, see **DRUG INTERACTIONS**.

Table 31: Drug Interactions: Pharmacokinetic Parameters for Darunavir in the Presence of Co-administered Drugs

Co-Administered Drug	Dose/Schedule		N	PK	LS Mean Ratio (90% CI) of Darunavir Pharmacokinetic Parameters With/Without Co-administered Drug No Effect =1.00		
	Co-Administered Drug	Darunavir/ ritonavir			C_{max}	AUC	C_{min}
Co-Administration With Other Protease Inhibitors							
Atazanavir	300 mg q.d.[a]	400/100 mg b.i.d. [b]	13	↔	1.02 (0.96-1.09)	1.03 (0.94-1.12)	1.01 (0.88-1.16)
Indinavir	800 mg b.i.d.	400/100 mg b.i.d.	9	↑	1.11 (0.98-1.26)	1.24 (1.09-1.42)	1.44 (1.13-1.82)
Lopinavir/ Ritonavir	400/100 mg b.i.d.	1200/100 mg b.i.d.[c]	14	↓	0.79 (0.67-0.92)	0.62 (0.53-0.73)	0.49 (0.39-0.63)
	533/133.3 mg b.i.d.	1200 mg b.i.d.[c]	15	↓	0.79 (0.64-0.97)	0.59 (0.50-0.70)	0.45 (0.38-0.52)
Ritonavir	Titrated: 300 to 600 mg b.i.d. over 6 days	Darunavir 800 mg single dose	9	↑	1.97 (1.40-2.77)	9.23 (6.62-12.88)	-
Saquinavir hard gel capsule	1000 mg b.i.d.	400/100 mg b.i.d.	14	↓	0.83 (0.75-0.92)	0.74 (0.63-0.86)	0.58 (0.47-0.72)
Co-Administration With Other Antiretrovirals							
Didanosine	400 mg q.d.	600/100 mg b.i.d.	17	↔	0.93 (0.86-1.00)	1.01 (0.95-1.07)	1.07 (0.95-1.21)
Efavirenz	600 mg q.d.	300/100 mg b.i.d.	12	↓	0.85 (0.72-1.00)	0.87 (0.75-1.01)	0.69 (0.54-0.87)
Etravirine	200 mg b.i.d.	600/100 mg b.i.d.	15	↔	1.11 (1.01-1.22)	1.15 (1.05-1.26)	1.02 (0.90-1.17)
Nevirapine	200 mg b.i.d.	400/100 mg b.i.d.	8	↑	1.40 [d] (1.14-1.73)	1.24 [d] (0.97-1.57)	1.02 [d] (0.79-1.32)
Rilpivirine	150 mg q.d. [e]	800/100 mg q.d.	15	↔	0.90 (0.81- 1.00)	0.89 (0.81-0.99)	0.89 (0.68-1.16)
Tenofovir Disoproxil Fumarate	300 mg q.d.	300/100 mg b.i.d.	12	↑	1.16 (0.94-1.42)	1.21 (0.95-1.54)	1.24 (0.90-1.69)
Co-Administration With Other Drugs							
Carbamazepine	200 mg b.i.d.	600/100 mg b.i.d.	16	↔	1.04 (0.93-1.16)	0.99 (0.90-1.08)	0.85 (0.73-1.00)
Clarithromycin	500 mg b.i.d.	400/100 mg b.i.d.	17	↔	0.83 (0.72-0.96)	0.87 (0.75-1.01)	1.01 (0.81-1.26)
Ketoconazole	200 mg b.i.d.	400/100 mg b.i.d.	14	↑	1.21 (1.04-1.40)	1.42 (1.23-1.65)	1.73 (1.39-2.14)
Omeprazole	20 mg q.d.	400/100 mg b.i.d.	16	↔	1.02 (0.95-1.09)	1.04 (0.96-1.13)	1.08 (0.93-1.25)
Paroxetine	20 mg q.d.	400/100 mg b.i.d.	16	↔	0.97 (0.92-1.02)	1.02 (0.95-1.10)	1.07 (0.96-1.19)

Table 31: Drug Interactions: Pharmacokinetic Parameters for Darunavir in the Presence of Co-administered Drugs

Co-Administered Drug	Dose/Schedule		N	PK	LS Mean Ratio (90% CI) of Darunavir Pharmacokinetic Parameters With/Without Co-administered Drug No Effect =1.00		
	Co-Administered Drug	Darunavir/ ritonavir			C_{max}	AUC	C_{min}
Ranitidine	150 mg b.i.d.	400/100 mg b.i.d.	16	↔	0.96 (0.89-1.05)	0.95 (0.90-1.01)	0.94 (0.90-0.99)
Rifabutin	150 mg q.o.d. [f]	600/100 mg b.i.d.	11	↑	1.42 (1.21-1.67)	1.57 (1.28-1.93)	1.75 (1.28-2.37)
Sertraline	50 mg q.d.	400/100 mg b.i.d.	13	↔	1.01 (0.89-1.14)	0.98 (0.84-1.14)	0.94 (0.76-1.16)
Telaprevir	750 mg q8h for 10 days	600/100 mg b.i.d. for 20 days	11[g]	↓	0.60 (0.56- 0.64)	0.60 (0.57- 0.63)	0.58 (0.52-0.64)
	1125 mg q12h for 4 days	600/100 mg b.i.d. for 24 days	15	↓	0.53 (0.47- 0.59)	0.49 (0.43-0.55)	0.42 (0.35- 0.51)

N = number of patients with data; - = no information available.
[a] q.d. = once daily
[b] b.i.d. = twice daily
[c]The pharmacokinetic parameters of darunavir in this study were compared with the pharmacokinetic parameters following administration of darunavir/ritonavir 600/100 mg b.i.d.
[d] Ratio based on between-study comparison.
[e] This interaction study has been performed with a dose higher than the recommended dose for rilpivirine (25 mg once daily) assessing the maximal effect on the co-administered drug.
[f] q.o.d. = every other day
[g] N=14 for C_{max}

Table 32: Drug Interactions: Pharmacokinetic Parameters for Co-administered Drugs in the Presence of Darunavir/Ritonavir

Co-Administered Drug	Dose/Schedule		N	PK	LS Mean Ratio (90% CI) of Co-Administered Drug Pharmacokinetic Parameters With/Without Darunavir No effect =1.00		
	Co-Administered Drug	Darunavir/ ritonavir			C_{max}	AUC	C_{min}
Co-Administration With Other Protease Inhibitors							
Atazanavir	300 mg q.d.[a] /100 mg ritonavir q.d. when administered alone 300 mg q.d. when administered with darunavir/ ritonavir	400/100 mg b.i.d. [b]	13	↔	0.89 (0.78-1.01)	1.08 (0.94-1.24)	1.52 (0.99-2.34)
Indinavir	800 mg b.i.d. /100 mg ritonavir b.i.d. when administered alone 800 mg b.i.d. when administered with darunavir/ ritonavir	400/100 mg b.i.d.	9	↑	1.08 (0.95-1.22)	1.23 (1.06-1.42)	2.25 (1.63-3.10)
Lopinavir/ Ritonavir	400/100 mg b.i.d[c]	1200/100 mg b.i.d.	14	↔	0.98 (0.78-1.22)	1.09 (0.86-1.37)	1.23 (0.90-1.69)
	533/133.3 mg b.i.d.[c]	1200 mg b.i.d.	15	↔	1.11 (0.96-1.30)	1.09 (0.96-1.24)	1.13 (0.90-1.42)
Saquinavir hard gel capsule	1000 mg b.i.d. /100 mg ritonavir b.i.d. when administered alone 1000 mg b.i.d. when administered with darunavir/ ritonavir	400/100 mg b.i.d.	12	↔	0.94 (0.78-1.13)	0.94 (0.76-1.17)	0.82 (0.52-1.30)
Co-Administration With Other Antiretrovirals							
Didanosine	400 mg q.d.	600/100 mg b.i.d.	17	↔	0.84 (0.59-1.20)	0.91 (0.75-1.10)	-
Efavirenz	600 mg q.d.	300/100 mg b.i.d.	12	↑	1.15 (0.97-1.35)	1.21 (1.08-1.36)	1.17 (1.01-1.36)
Etravirine	100 mg b.i.d.	600/100 mg b.i.d.	14	↓	0.68 (0.57-0.82)	0.63 (0.54-0.73)	0.51 (0.44-0.61)

Table 32: Drug Interactions: Pharmacokinetic Parameters for Co-administered Drugs in the Presence of Darunavir/Ritonavir

Co-Administered Drug	Dose/Schedule		N	PK	LS Mean Ratio (90% CI) of Co-Administered Drug Pharmacokinetic Parameters With/Without Darunavir No effect =1.00		
	Co-Administered Drug	Darunavir/ ritonavir			C_{max}	AUC	C_{min}
Rilpivirine	150 mg q.d. [a]	800/100 mg q.d.	14	↑	1.79 (1.56-2.06)	2.30 (1.98-2.67)	2.78 (2.39-3.24)
Nevirapine	200 mg b.i.d.	400/100 mg b.i.d.	8	↑	1.18 (1.02-1.37)	1.27 (1.12-1.44)	1.47 (1.20-1.82)
Tenofovir Disoproxil Fumarate	300 mg q.d.	300/100 mg b.i.d.	12	↑	1.24 (1.08-1.42)	1.22 (1.10-1.35)	1.37 (1.19-1.57)
Maraviroc	150 mg b.i.d.	600/100 mg b.i.d.	12	↑	2.29 (1.46-3.59)	4.05 (2.94-5.59)	8.00 (6.35-10.1)
Co-Administration With Other Drugs							
Atorvastatin	40 mg q.d. when administered alone 10 mg q.d. when administered with darunavir/ Ritonavir	300/100 mg b.i.d.	15	↑	0.56 (0.48-0.67)	0.85 (0.76-0.97)	1.81 (1.37-2.40)
Buprenorphine/ Naloxone	8/2 mg to 16/4 mg q.d.	600/100 mg b.i.d.	17	↔	0.92 [d] (0.79-1.08)	0.89[d] (0.78-1.02)	0.98[d] (0.82-1.16)
Norbuprenorphine			17	↑	1.36 (1.06-1.74)	1.46 (1.15-1.85)	1.71 (1.29-2.27)
Carbamazepine	200 mg b.i.d.	600/100 mg b.i.d.	16	↑	1.43 (1.34-1.53)	1.45 (1.35-1.57)	1.54 (1.41-1.68)
Carbamazepine epoxide			16	↓	0.46 (0.43-0.49)	0.46 (0.44-0.49)	0.48 (0.45-0.51)
Clarithromycin	500 mg b.i.d.	400/100 mg b.i.d.	17	↑	1.26 (1.03-1.54)	1.57 (1.35-1.84)	2.74 (2.30-3.26)
Digoxin	0.4 mg	600/100 mg b.i.d.	8	↑	1.15 (0.89-1.48)	1.36 (0.81-2.27)	-
Ethinyl Estradiol (EE)	Ortho-Novum 1/35 (35 µg EE / 1 mg NE)	600/100 mg b.i.d.	11	↓	0.68 (0.61-0.74)	0.56 (0.50-0.63)	0.38 (0.27-0.54)
Norethindrone (NE)			11	↓	0.90 (0.83-0.97)	0.86 (0.75-0.98)	0.70 (0.51-0.97)
Ketoconazole	200 mg b.i.d.	400/100 mg b.i.d.	15	↑	2.11 (1.81-2.44)	3.12 (2.65-3.68)	9.68 (6.44-14.55)

Table 32: Drug Interactions: Pharmacokinetic Parameters for Co-administered Drugs in the Presence of Darunavir/Ritonavir

Co-Administered Drug	Dose/Schedule		N	PK	LS Mean Ratio (90% CI) of Co-Administered Drug Pharmacokinetic Parameters With/Without Darunavir No effect =1.00		
	Co-Administered Drug	Darunavir/ ritonavir			C_{max}	AUC	C_{min}
R-Methadone	55-150 mg q.d.	600/100 mg b.i.d.	16	↓	0.76 (0.71-0.81)	0.84 (0.78-0.91)	0.85 (0.77-0.94)
Paroxetine	20 mg q.d.	400/100 mg b.i.d.	16	↓	0.64 (0.59-0.71)	0.61 (0.56-0.66)	0.63 (0.55-0.73)
Pravastatin	40 mg single dose	600/100 mg b.i.d.	14	↑	1.63 (0.95-2.82)	1.81 (1.23-2.66)	-
Rifabutin	150 mg q.o.d.[e] when administered with PREZISTA®/ritonavir	600/100 mg b.i.d. [f]	11	↑	0.72 (0.55-0.93)	0.93 (0.80-1.09)	1.64 (1.48-1.81)
25-*O*-desacetyl-rifabutin	300 mg q.d. when administered alone		11	↑	4.77 (4.04-5.63)	9.81 (8.09-11.9)	27.1 (22.2-33.2)
Sertraline	50 mg q.d.	400/100 mg b.i.d.	13	↓	0.56 (0.49-0.63)	0.51 (0.46-0.58)	0.51 (0.45-0.57)
Sildenafil	100 mg (single dose) administered alone 25 mg (single dose) when administered with darunavir/ ritonavir	400/100 mg b.i.d.	16	↑	0.62 (0.55-0.70)	0.97 (0.86-1.09)	-
Telaprevir	750 mg q8h for 10 days	600/100 mg b.i.d. for 20 days	11[g]	↓	0.64 (0.61-0.67)	0.65 (0.61-0.69)	0.68 (0.63-0.74)

N = number of patients with data;- = no information available.
[a] q.d. = once daily
[b] b.i.d. = twice daily
[c] The pharmacokinetic parameters of lopinavir in this study were compared with the pharmacokinetic parameters following administration of lopinavir/ritonavir 400/100 mg b.i.d.
[d] ratio is for buprenorphine; mean C_{max} and AUC_{24} for naloxone were comparable when buprenorphine/naloxone was administered with or without PREZISTA®/rtv
[e] q.o.d. = every other day
[f] In comparison to rifabutin 300 mg q.d.
[g] N=14 for C_{max}

Population Pharmacokinetics

Population pharmacokinetic analysis in HIV-1-infected patients showed that darunavir pharmacokinetics is not considerably different in the age range (18 to 75 years) evaluated in HIV-1-infected patients. Population pharmacokinetic analysis showed a slightly higher darunavir exposure in HIV-1-infected females compared to males. This difference is not considered clinically relevant.

Population pharmacokinetic analysis of darunavir in HIV-1-infected patients indicated that race had no apparent effect on the exposure to darunavir. The steady-state pharmacokinetic

parameters of darunavir in patients with mild and moderate hepatic impairment were comparable with those in healthy patients, therefore, no dose adjustment is required in patients with mild or moderate hepatic impairment. PREZISTA® has not been studied in patients with severe hepatic impairment.

Population pharmacokinetic analysis showed that the pharmacokinetics of darunavir were not significantly affected in HIV-1-infected patients with moderate renal impairment. There are no pharmacokinetic data available in HIV-1-infected patients with severe renal impairment or end stage renal disease. However, since the renal clearance of darunavir is limited, a decrease in total body clearance is not expected in patients with renal impairment.

The population pharmacokinetics derived geometric mean (SD) C_{0h} and AUC_{12h} for darunavir in 119 HIV-1-infected patients (TMC114-C213 and TMC114-C202, Primary 24-Week Analysis) receiving [600/100 mg b.i.d. darunavir/ritonavir] is 3578 (±1151) ng/mL and 62349 (±16143) ng.h/mL, respectively.

The pharmacokinetics of darunavir, co-administered with low dose ritonavir (100 mg), has been evaluated in healthy adult volunteers and in HIV-1-infected patients. Table 33 displays the population pharmacokinetic estimates of darunavir after oral administration of PREZISTA®/rtv 600/100 mg twice daily [based on sparse sampling in 285 patients in study TMC114-C214 and 119 patients (integrated data) from Studies TMC114-C202 and TMC114-C213] and PREZISTA®/rtv 800/100 mg once daily [based on sparse sampling in 335 patients in Study TMC114-C211 and 280 patients in Study TMC114-C229] to HIV-1-infected patients.

Table 33: Population Pharmacokinetic Estimates of Darunavir at PREZISTA®/rtv 800/100 mg once daily (Study TMC114-C211, 48-Week Analysis and Study TMC114-C229, 48-Week Analysis) and PREZISTA®/rtv 600/100 mg twice daily (Study TMC114-C214, 48-Week Analysis and Integrated data from Studies TMC114-C213 and TMC114-C202, Primary 24-Week Analysis)

Parameter	Study TMC114-C211 PREZISTA®/ rtv 800/100 mg once daily N = 335	Study TMC114-C229 PREZISTA®/ rtv 800/100 mg once daily N = 280	Study TMC114-C214 PREZISTA®/ rtv 600/100 mg twice daily N = 285	Study TMC114-C229 PREZISTA®/ rtv 600/100 mg twice daily N = 278	Studies TMC114-C213 and TMC114-C202 (integrated data) PREZISTA®/ rtv 600/100 mg twice daily N =119
AUC_{24h} (ng·h/mL)[a]					
Mean ± Standard Deviation	93026 ± 27050	93334 ± 28626	116796 ± 33594	114302 ± 32681	124698 ± 32286
Median (Range)	87854 (45000-219240)	87788 (45456-236920)	111632 (64874-355360)	109401 (48934-323820)	123336 (67714-212980)
C_{0h} (ng/mL)					
Mean ± Standard Deviation	2282 ± 1168	2160 ± 1201	3490 ± 1401	3386 ± 1372	3578 ± 1151
Median (Range)	2041 (368-7242)	1896 (184-7881)	3307 (1517-13198)	3197 (250-11865)	3539 (1255-7368)

N = number of patients with data. [a] AUC_{24h} is calculated as AUC_{12h}*2

MICROBIOLOGY

Antiviral Activity *In Vitro*

Darunavir exhibits activity against laboratory strains and clinical isolates of HIV-1 and laboratory strains of HIV-2 in acutely infected T-cell lines, human peripheral blood mononuclear cells and human monocytes/macrophages with median EC_{50} values ranging from 1.2 to 8.5 nM (0.7 to 5.0 ng/mL). Darunavir demonstrates antiviral activity *in vitro* against a broad panel of HIV-1 group M (A, B, C, D, E, F, G) and group O primary isolates with EC_{50} values ranging from < 0.1 to 4.3 nM.

These EC_{50} values are well below the 50% cellular toxicity concentration range of 87 µM to > 100 µM. The EC_{50} value of darunavir increases by a median factor of 5.4 in the presence of human serum.

Darunavir showed synergistic antiviral activity when studied in combination with the PIs ritonavir, nelfinavir, or amprenavir, and additive antiviral activity when studied in combination with the PIs indinavir, saquinavir, lopinavir, atazanavir, or tipranavir, the nucleoside (nucleotide) reverse transcriptase inhibitors (N(t)RTIs) zidovudine, lamivudine, zalcitabine, didanosine, stavudine, abacavir, emtricitabine, or tenofovir, the non-nucleoside reverse transcriptase inhibitors (NNRTIs) nevirapine, delavirdine, rilpivirine or etravirine, or efavirenz, and the fusion inhibitor enfuvirtide. No antagonism was observed between darunavir and any of these antiretrovirals *in vitro*.

Resistance *In Vitro*

In vitro selection of darunavir-resistant virus from wild-type HIV-1 was lengthy (more than 2 years). The selected viruses were unable to grow in the presence of darunavir concentrations above 220 nM. Viruses selected in these conditions and showing decreased susceptibility to darunavir (range: 23- to 50-fold) harboured 2 to 4 amino acid substitutions in the protease gene. The decreased susceptibility to darunavir of the emerging viruses in the selection experiment could not be explained by the emergence of these protease mutations.

In vitro selection of darunavir-resistant HIV-1 (range: 53- to 641-fold change in EC_{50} values) from 9 HIV-1 strains harbouring multiple PI resistance-associated mutations (RAMs) resulted in the overall emergence of 22 mutations in the protease, of which L10F, V32I, L33F, S37N, M46I, I47V, I50V, L63P, A71V, and I84V were present in more than 50% of the 9 darunavir-resistant isolates. A minimum of 8 of these darunavir *in vitro* selected mutations, from which at least 2 were already present in the protease prior to selection, were required in the HIV-1 protease to render a virus resistant (fold change (FC) > 10) to darunavir.

In 1113 clinical isolates resistant to amprenavir, atazanavir, indinavir, lopinavir, nelfinavir, ritonavir, saquinavir and/or tipranavir, and in 886 baseline isolates from the patients enrolled in the TMC114-C213 (POWER 1) and TMC114-C202 (POWER 2) trials and in the TMC114-C215 (POWER 3) analysis, only the subgroups with > 10 PI resistance-associated mutations showed a median FC for darunavir > 10.

Cross-Resistance *In Vitro*

Cross-resistance has been observed among PIs. Darunavir has a < 10-fold decreased susceptibility against 90% of 3309 clinical isolates resistant to amprenavir, atazanavir, indinavir, lopinavir, nelfinavir, ritonavir, saquinavir and/or tipranavir showing that viruses resistant to most PIs remain susceptible to darunavir.

Seven of the 9 darunavir-resistant viruses selected from PI-resistant viruses had phenotypic data for tipranavir. Six of those showed a FC in EC_{50} value < 3 for tipranavir, indicative of limited cross-resistance between these 2 PIs.

Cross-resistance between darunavir and the nucleoside/nucleotide reverse transcriptase inhibitors, the non-nucleoside reverse transcriptase inhibitors the entry inhibitors, or the integrase inhibitor is unlikely because the viral targets for those inhibitors are different.

***In Vivo* Selection of Viral Resistance During Darunavir/ritonavir Therapy**

In the 192-week analysis of the TMC114-C211 (ARTEMIS) trial, the number of virologic failures was lower in the group of patients receiving PREZISTA®/rtv 800/100 mg q.d. than in patients receiving lopinavir/ritonavir 800/200 mg per day (16.0% vs. 20.5%, respectively). In the virologic failures of the PREZISTA®/rtv arm with paired baseline/endpoint genotype data, four patients with developing PI RAMs were identified. In the virologic failures of the lopinavir/ritonavir arm with paired baseline/endpoint genotype data, nine patients with developing PI RAMs at endpoint were identified. This was not associated with a loss in susceptibility to lopinavir. None of the developing mutations in the PREZISTA®/rtv group or in the lopinavir/rtv group were primary (i.e., major) PI mutations. In four virologic failures in the PREZISTA®/rtv arm and seven virologic failures in the lopinavir/ritonavir arm, a maximum of two developing NRTI RAMs were identified. The development of the NRTI RAM at position 184 (n= 9) was identified, which was associated with a decreased susceptibility to emtricitabine (FTC) included in the fixed background regimen.

In the 48-week analysis of the TMC114-C229 (ODIN) trial, the number of virologic failures was comparable in the PREZISTA®/rtv 800/100 mg q.d. group and the PREZISTA®/rtv 600/100 mg b.i.d. group (22.1% vs. 18.2%, respectively). Of the virologic failures, the PREZISTA®/rtv 800/100 mg q.d. group reported 7 (12%) patients with developing PI RAMs compared to 4 (10%) patients in the PREZISTA®/rtv 600/100 mg b.i.d group. Only 1 subject, in the DRV/rtv q.d. group, developed primary (major) PI mutations (V32I, M46I, L76V and I84V), which included 3 DRV RAMs (V32I, L76Vand I84V). The emergence of these DRV RAMs was associated with loss of DRV susceptibility.

All virologic failures from the PREZISTA®/rtv 600/100 mg b.i.d. group retained susceptibility to darunavir. Four (6.7%) and 3 (7.1%) virologic failures developed 1 or 2 NRTI RAMs in the PREZISTA®/rtv 800/100 mg q.d. and the PREZISTA®/rtv 600/100 mg b.i.d. groups, respectively. In 3 and 2 of these virologic failures in the PREZISTA®/rtv 800/100 mg q.d. and the PREZISTA®/rtv 600/100 mg b.i.d. groups, respectively, the development of these NRTI RAMs (V75I+M184V; M184V; T215Y in the q.d. group and M184V; M41L+T215Y in the

b.i.d. group) was associated with a decreased susceptibility to a NRTI included in the background regimen.

In the 96-week analysis of the TMC114-C214 (TITAN) trial, the number of virologic failures was lower in the group of patients receiving PREZISTA®/rtv 600/100 mg b.i.d. than in patients receiving lopinavir/ritonavir 400/100 mg b.i.d. (41/298, 13.8% versus 76/297, 25.6%, respectively). Examination of patients who failed on PREZISTA®/rtv 600/100 mg twice daily and had post-baseline genotypes and phenotypes showed that 6 patients (6/39; 15%) developed PI substitutions on darunavir/ritonavir treatment resulting in decreased susceptibility to darunavir. Five of the 6 had baseline PI resistance-associated substitutions and baseline darunavir phenotypes > 7. The most common emerging PI substitutions in the virologic failures were V32I, I47V, T74P, and L76V.

For the lopinavir/ritonavir arm, baseline and endpoint genotype was available for 72 of 76 virologic failures. Comparing patients with available baseline and endpoint genotypes, fewer virologic failures treated with PREZISTA®/rtv 600/100 mg b.i.d. than with lopinavir/ritonavir 400/100 mg b.i.d. developed primary (i.e. major) PI mutations (7 vs. 25, respectively) or NRTI RAMS (4 vs. 20, respectively) or lost susceptibility to the PI (3 vs. 17, respectively) or NRTI(s) (4 vs. 20, respectively) used in the treatment regimen.

In a pooled analysis of the POWER and DUET trials, the percentage of rebounders (patients who lost a viral load $\geq$1.0 $\log_{10}$ below baseline) was 17.6% (188 out of 1071 patients). Baseline and endpoint genotype was available for 185 out of 188 rebounders. The most common protease mutations that developed in $\geq$ 20% of the isolates from patients who experienced virological failure by rebound were V32I, I54L, and L89V. Amino acid substitutions that developed in 10 to 20% of the isolates were V11I, I13V, L33F, I50V, and F53L. The percentage of never suppressed patients (patients that never achieved a viral load $\geq$1.0 $\log_{10}$ below baseline at 2 consecutive visits) was 19.7% (211 out of 1071 patients). Baseline and endpoint genotype was available for 197 out of the 211 never suppressed patients. The most common protease mutations that developed in $\geq$20% of the analyzed never suppressed patients were V32I and I54L. Mutations that developed in 10 to 20% of these never suppressed patients were V11I, I15V, L33F, I47V and L89V.

***In Vivo* Cross-Resistance with Other Protease Inhibitors**

In the virologic failures of the ARTEMIS trial, no cross-resistance with other PIs was observed.

Of the viruses isolated from patients receiving PREZISTA®/rtv 800/100 mg q.d. experiencing virologic failure in the ODIN trial, 98% remained susceptible to darunavir after treatment. In the same group of patients, 96% to 100% that were susceptible at baseline to amprenavir, atazanavir, indinavir, lopinavir, saquinavir or tipranivir remained susceptible to these protease inhibitors after treatment. In the virologic failures receiving PREZISTA®/rtv 600/100 mg b.i.d. no cross-resistance with other PIs was observed.

In the TITAN trial, the number of virologic failures was lower in the DRV/rtv group than in the LPV/rtv group and fewer virologic failures treated with DRV/rtv than with LPV/rtv lost susceptibility to PIs. Of the viruses isolated from patients receiving PREZISTA®/rtv 600/100 mg b.i.d. experiencing virologic failures in the TITAN trial, 8% of those susceptible to darunavir at baseline developed decreased susceptibility to darunavir during treatment. In the same group of patients, 97% to 100% that were susceptible at baseline to amprenavir, atazanavir, indinavir, lopinavir, saquinavir or tipranavir remained susceptible after PREZISTA®/rtv treatment.

Of the viruses isolated from patients experiencing virologic failure by rebound from the PREZISTA®/rtv 600/100 mg b.i.d. group, of the POWER and DUET trials, 85% that were susceptible to darunavir at baseline developed decreased susceptibility to darunavir during treatment. In the same group of patients, 71% of viruses that were susceptible to tipranavir at baseline remained susceptible after treatment. In the POWER trials, patients with resistance to tipranavir (FC > 3) at baseline showed a mean change in viral load at Week 24 of -1.38 $\log_{10}$. Cross resistance with the other PIs could not be studied in the POWER and DUET trials, since most of the baseline viruses were already resistant to these PIs. Patients with no susceptible PI at baseline (excluding tipranavir) showed a mean change in viral load at Week 24 of -1.57 $\log_{10}$.

Baseline Genotype or Phenotype and Virologic Outcome

In a pooled analysis of the 600/100 mg b.i.d. groups of the POWER and DUET trials, the presence at baseline of three or more of the darunavir- specific mutations (V11I, V32I, L33F, I47V, I50V, I54L or M, T74P, L76V, I84V or L89V) was associated with a decreased virologic response to PREZISTA®/rtv. In early treatment-experienced patients (TITAN) three or more of these mutations were only found in 4% of the patients at baseline.

Table 34: Response (HIV-1 RNA < 50 copies/mL at week 24) to PREZISTA®/rtv 600/100 mg b.i.d. by baseline genotype[a] and by use of enfuvirtide: As-treated analysis of the POWER and DUET trials

Number of mutations at baseline[a]	All % n/N	No/non-naïve use of ENF % n/N	Naïve use of ENF % n/N
All Ranges	45% 455/1014	39% 290/741	60% 165/273
0 – 2	54% 359/660	50% 238/477	66% 121/183
3	39% 67/172	29% 35/120	62% 32/52
≥4	12% 20/171	7% 10/135	28% 10/36

[a] Number of mutations from the list of mutations associated with a diminished response to PREZISTA®/rtv (V11I, V32I, L33F, I47V, I50V, I54L or M, T74P, L76V, I84V or L89V)

Baseline darunavir phenotype (shift in susceptibility relative to reference) was shown to be the most predictive factor of virologic outcome. Response rates assessed by baseline darunavir phenotype are shown in Table 35. The data are provided to give clinicians information on the likelihood of virologic success based on pre-treatment susceptibility to darunavir.

Table 35: Response (HIV-1 RNA < 50 copies/mL at week 24) to PREZISTA®/rtv 600/100 mg b.i.d. by baseline darunavir phenotype and by use of enfuvirtide: As-treated analysis of the POWER and DUET trials

Baseline darunavir phenotype	All % n/N	No/non naïve use of ENF % n/N	Naïve use of ENF % n/N
All ranges	45% 455/1014	39% 290/741	60% 165/273
≤ 10	55% 364/659	51% 244/477	66% 120/182
10-40	29% 59/203	17% 25/147	61% 34/56
>40	8% 9/118	5% 5/94	17% 4/24

In deciding on a new regimen for patients who have failed an antiretroviral regimen, careful consideration should be given to the treatment history and to resistance testing results where available.

TOXICOLOGY

Animal toxicology studies have been conducted with darunavir alone, in mice, rats and dogs and in combination with ritonavir in rats and dogs.

In chronic toxicology studies in rats and dogs, there were only limited effects of treatment with darunavir. In the rat the key target organs identified were the hematopoietic system, the blood coagulation system, liver and thyroid, observed at 100 mg/kg/day and above and at exposures below clinical levels. A variable but limited decrease in red blood cell-related parameters was observed, together with increases in activated PTT. The observed liver and thyroid changes were considered to reflect an adaptive response to enzyme induction in the rat rather than an adverse effect. In combination toxicity studies with ritonavir, no additional target organs of toxicity were reported in rats. In the dog, no major toxicity findings or key target organs were identified at doses up to 120 mg/kg/day and exposures equivalent to clinical exposure at the recommended dose.

Carcinogenesis and Mutagenesis

Darunavir was evaluated for carcinogenic potential by oral gavage administration to mice and rats up to 104 weeks. Daily doses of 150, 450 and 1000 mg/kg were administered to mice and doses of 50, 150 and 500 mg/kg were administered to rats. A dose related increase in the

incidences of hepatocellular adenomas and carcinomas were observed in males and females of both species. Thyroid follicular cell adenomas were noted in male rats. Administration of darunavir did not cause a statistically significant increase in the incidence of any other benign or malignant neoplasm in mice or rats. The observed hepatocellular findings in rodents are considered to be of limited relevance to humans. Repeated administration of darunavir to rats caused hepatic microsomal enzyme induction and increased thyroid hormone elimination, which predispose rats, but not humans, to thyroid neoplasms. At the highest tested doses, the systemic exposures (based on AUC) to darunavir were between 0.4- and 0.7- fold (mice) and 0.7- and 1-fold (rats), relative to those observed in humans at the recommended therapeutic doses (600/100 mg twice daily or 800/100 mg once daily). Darunavir was not mutagenic or genotoxic in a battery of *in vitro* and *in vivo* assays including bacterial reverse mutation (Ames), chromosomal aberration in human lymphocytes and *in vivo* micronucleus test in mice.

Reproductive and Developmental Toxicity

Investigation of fertility and early embryonic development was performed in rats, teratogenicity studies were conducted in mice, rats and rabbits, and the pre- and post-natal development study was conducted in rats.

In the fertility and early embryonic development study, a significant decrease in body weight gain with subsequent related reduction in the number of ovulations resulting in a reduction in the number of live fetuses was observed in female rats treated with 1000 mg/kg. Otherwise, there were no effects on mating or fertility with darunavir treatment up to 1000 mg/kg/day and exposure levels below (AUC 0.5-fold) that in humans at the clinically recommended dose. Up to the same dose levels, there was no teratogenicity with darunavir in rats and rabbits when treated alone nor in mice when treated in combination with ritonavir. The exposure levels were lower than those observed with the recommended clinical dose in humans. In addition, rats treated in combination with ritonavir showed no teratogenicity when exposed to higher levels of darunavir than those achieved with the recommended clinical dose in humans. In a pre- and post-natal development assessment in rats, darunavir with and without ritonavir caused a transient reduction in body weight gain of the offspring during lactation. This was attributed to drug exposure via the milk. No post-weaning functions were affected with darunavir alone or in combination with ritonavir.

In juvenile rats directly dosed with darunavir (from 20 mg/kg to 1000 mg/kg) up to days 23 to 26 of age, mortality was observed and, in some of the animals, convulsions. Within this age range exposures in plasma, liver and brain were dose and age dependent and were considerably greater than those observed in adult rats. These findings were attributed to the ontogeny of the CYP450 liver enzymes involved in the metabolism of darunavir and the immaturity of the blood-brain barrier. No treatment-related mortalities were noted in juvenile rats dosed at 1000 mg/kg darunavir (single dose) on day 26 of age or at 500 mg/kg (repeated dose) from day 23 to 50 of age, and the exposures and toxicity profile were comparable to those observed in adult rats. In humans, the activity of drug-metabolizing enzymes approaches adult values by 3 years of age.

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PART III: CONSUMER INFORMATION

PrPREZISTA®
darunavir tablets
darunavir oral suspension

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

ABOUT THIS MEDICATION

What the medication is used for:
PREZISTA® is used for the treatment of HIV (Human Immunodeficiency Virus) infection in adults and in children 3 years of age and older who have taken anti-HIV medicines in the past, when co-administered with ritonavir and other antiretroviral medications. HIV is the virus that causes AIDS (Acquired Immune Deficiency Syndrome). PREZISTA® is a type of anti-HIV medicine called a protease (PRO-tee-ase) inhibitor. PREZISTA® is available as an oral tablet and an oral suspension.

PREZISTA® does not reduce the risk of passing HIV to others through sexual contact, sharing needles, or being exposed to your blood. For your health and the health of others, it is important to always practice safer sex by using a latex or polyurethane condom or other barrier method to lower the chance of sexual contact with any body fluids such as semen, vaginal secretions, or blood. Never use or share dirty needles.

Ask your doctor if you have any questions on how to prevent passing HIV to other people.

What it does:
PREZISTA® blocks HIV protease, an enzyme which is needed for HIV to multiply. When used with other anti-HIV medicines, PREZISTA® can help to reduce the amount of HIV in your blood (called "viral load") and increase your CD4+ (T) cell count. HIV infection destroys CD4+ (T) cells, which are important to the immune system. The immune system helps fight infection. Reducing the amount of HIV and increasing the CD4+ (T) cell count may improve your immune system.

PREZISTA® is always taken with and at the same time as 100 mg of ritonavir (NORVIR®), in combination with other anti-HIV medicines. PREZISTA® should also be taken with food.

PREZISTA® does not cure HIV infection or AIDS. At present, there is no cure for HIV infection. People taking PREZISTA® may still develop infections or other conditions associated with HIV infection. Because of this, it is very important for you to remain under the care of a doctor.

When it should not be used:
Together with your doctor, you need to decide whether taking PREZISTA® is right for you.

Do not take PREZISTA®:

- if you are allergic to darunavir or any of the other ingredients in PREZISTA® (see **What the nonmedicinal ingredients are**)
- if you are allergic to ritonavir (NORVIR®)
- if you have severe liver disease
- if you take any of the following types of medicines because you could experience serious side effects:

Type of Drug	Examples of Generic Names (Brand Names)
Alpha1-Adrenoreceptor Antagonists (to treat enlarged prostate)	alfuzosin
Anti-coagulant	apixaban (Eliquis®) rivaroxaban (Xarelto®)
Antiarrhythmics (to treat abnormal heart rhythms)	bepridil[1] dronedarone (Multaq®) lidocaine (when taken by injection) quinidine amiodarone (Cordarone®)
Anti-gout (to treat gout and familial Mediteraranean fever) if you have renal hepatic impairment	colchicine
Antihistamines (to treat allergy symptoms)	astemizole[1] terfenadine[1]
Antimycobacterials (to treat tuberculosis)	rifampin (Rifadin®, Rifater®)
Ergot Derivatives (to treat migraine and headaches)	dihydroergotamine (Migranal®) ergonovine

Type of Drug	Examples of Generic Names (Brand Names)
	ergotamine (Cafergot®) methylergonovine
Gastrointestinal Motility Agents (to treat some digestive conditions)	cisapride[1]
Herbal products (to improve mood)	St. John's Wort
HMG-CoA Reductase Inhibitors also known as statins (to lower cholesterol)	lovastatin (Mevacor®) simvastatin (Zocor®)
Neuroleptics (to treat psychiatric conditions)	pimozide (Orap®)
PDE-5 Inhibitor (to treat pulmonary arterial hypertension)	sildenafil (Revatio®)
Sedatives/Hypnotics (to treat trouble with sleeping and/or anxiety)	midazolam (when taken by mouth) triazolam (Halcion®)

[1]Bepridil, astemizole, terfenadine and cisapride are not marketed in Canada.

What the medicinal ingredient is:
The active substance is darunavir.

Each 75 mg tablet contains 81.31 mg of darunavir ethanolate corresponding to 75 mg of darunavir.

Each 150 mg tablet contains 162.62 mg of darunavir ethanolate corresponding to 150 mg of darunavir.

Each 400 mg tablet contains 433.64 mg of darunavir ethanolate, corresponding to 400 mg of darunavir.

Each 600 mg tablet contains 650.46 mg of darunavir ethanolate, corresponding to 600 mg of darunavir.

Each 800 mg tablet contains 867.28 mg of darunavir ethanolate, corresponding to 800 mg of darunavir.

Each mL of the 100 mg/mL oral suspension contains 100 mg of darunavir (corresponding to 108.4 mg of darunavir ethanolate).

What the nonmedicinal ingredients are:
The other ingredients in the PREZISTA® tablets are colloidal anhydrous silica, crospovidone, magnesium stearate, and microcrystalline cellulose. The 800 mg tablet core also contains hypromellose. The tablet film coatings contain either OPADRY® II White (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, talc, titanium dioxide) for the 75 and 150 mg tablets or OPADRY® II Orange (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, titanium dioxide, talc, sunset yellow FCF aluminum lake) for the 400 and 600 mg tablets or OPADRY® II Dark Red (polyethylene glycol, polyvinyl alcohol - partially hydrolyzed, titanium dioxide, talc, iron oxide red) for the 800 mg tablets.

The other ingredients in the PREZISTA® oral suspension are hydroxypropyl cellulose, microcrystalline cellulose and carmellose sodium, sucralose, sodium methyl parahydroxybenzoate, masking flavor, citric acid monohydrate, and strawberry cream flavor.

What dosage forms it comes in:
100 mg/mL oral suspension
75 mg tablets
150 mg tablets
400 mg tablets
600 mg tablets
800 mg tablets

WARNINGS AND PRECAUTIONS

BEFORE you use PREZISTA® talk to your doctor or pharmacist if you:

- have diabetes. In general, anti-HIV medicines, such as PREZISTA®, might increase sugar levels in the blood.
- have liver problems, including hepatitis B and/or C.
- have hemophilia. Anti-HIV medicines, such as PREZISTA®, might increase the risk of bleeding.
- are pregnant or planning to become pregnant. It is not known if PREZISTA® can harm your unborn baby. PREZISTA® should not be used in pregnancy unless your doctor believes the benefit is greater than the risk to the fetus. If you take PREZISTA® while you are pregnant, talk to your doctor about how you can be included in the Antiretroviral Pregnancy Registry.
- are breast-feeding or planning to breast-feed. Do not breast-feed if you are taking PREZISTA®. You should not breast-feed if you have HIV because of the chance of passing HIV to your baby. Talk with your doctor about the best way to feed your baby.
- are allergic to sulpha medications.

INTERACTIONS WITH THIS MEDICATION

Tell your doctor about all the medicines you take including prescription and nonprescription medicines, vitamins, and herbal supplements, including St. John's wort (*Hypericum perforatum*). PREZISTA® and many other medicines can interact. Sometimes serious side effects will happen if PREZISTA® is taken with certain other medicines (see "**When it should not be used**").

PREZISTA® should not be combined with phenobarbital, phenytoin, rifapentine, rifampin or St. John's wort because the combination may significantly lower the amount of PREZISTA® in your blood and reduce the effects of PREZISTA®.

PREZISTA® should not be combined with vardenafil, because you may be at increased risk of side effects of vardenafil such as low blood pressure, visual changes and penile erection lasting more than 4 hours.

Tell your doctor if you are taking estrogen-based contraceptives. PREZISTA® might reduce the effectiveness of estrogen-based contraceptives (birth control). Therefore, alternative methods of non-hormonal contraception, such as a condom, are recommended.

Tell your doctor if you take other anti-HIV medicines (e.g., rilpivirine). PREZISTA® can be combined with some other anti-HIV medicines while other combinations are not recommended.

Tell your doctor if you are taking any of the following medicines:

Type of Drug	Examples of Generic Names (Brand Names)
Antiarrhythmics (for the heart)	Digoxin disopyramide flecainide mexiletine propafenone
Anticancer Agents	dasatinib (Sprycel®) nilotinib (Tasigna®) vinblastine vincristine everolimus (Afinitor®)
Anticoagulants (to prevent the clotting of red blood cells)	apixaban (Eliquis®) dabigatran etexilate (Pradaxa®) rivaroxaban (Xarelto®) warfarin (Coumadin®)
Anticonvulsants (to treat epilepsy and prevent seizures)	carbamazepine (Tegretol®) phenobarbital phenytoin (Dilantin®)
Antigout (to treat gout and familial Mediterranean fever)	colchicine
Anti-bacterials (to treat bacterial infections)	clarithromycin (Biaxin®)
Antidepressants (to treat depression, anxiety, or panic disorder)	amitriptyline desipramine imipramine nortriptyline paroxetine (Paxil®) sertraline (Zoloft®) trazodone (Oleptro®)
Antifungals (to treat fungal infections)	ketoconazole (Nizoral®) itraconazole (Sporanox®) voriconazole (Vfend®) posaconazole (Posanol®)
Antimalarials (to treat malarial infections)	artemether/lumefantrine (Riamet® and Coartem®) Artemether/lumefantrine are not approved for use in Canada.
Antimycobacterials (to treat bacterial infections)	rifabutin (Mycobutin®) rifampin (Rifadin®, Rifater®) rifapentine
Beta-Blockers (to treat heart disease)	carvedilol metoprolol (Betaloc®, Lopresor®) timolol
Calcium Channel Blockers (to treat heart disease)	amlodipine (Caduet®. Twynsta®) diltiazem (Cardizem®, Tiazac®) felodipine nifedipine (Adalat®) nicardipine verapamil (Isoptin®, Verelan®)

Type of Drug	Examples of Generic Names (Brand Names)
Corticosteroids (to treat inflammation or asthma)	Budesonide (Pulmicort®, Rhinocort®, Symbicort®) dexamethasone fluticasone (Advair Diskus®, Cutivate®, Flonase®, Flovent Diskus®) prednisone (Winpred®)
Endothelin receptor Antagonists (to treat pulmonary arterial hypertension)	bosentan (Tracleer®)
HIV- CCR5 Antagonist (to treat HIV infection)	maraviroc (Celsentri®)
HIV- Integrase strand transfer Inhibitors (to treat HIV infection)	dolutegravir (Tivicay®) elvitegravir (Stribild®)
HIV- Non-Nucleoside Reverse Transcriptase Inhibitors (NNRTIs) (to treat HIV infection)	delavirdine (Rescriptor®)
HIV- Protesase Inhibitors (to treat HIV infection)	lopinavir/ritonavir (Kaletra®) saquinavir (Invirase®) indinavir (Crixivan®)
HMG-CoA Reductase Inhibitors (to lower cholesterol levels)	atorvastatin (Lipitor®) pravastatin (Pravachol®) rosuvastatin (Crestor®)
Immunosuppressants (to prevent organ transplant rejection)	cyclosporine (Sandimmune®, Neoral®) tacrolimus (Prograf®) sirolimus (Rapamune®) everolimus (Afinitor®)
Inhaled beta agonist	salmeterol (Advair®)
Narcotic Analgesics (to treat opioid dependence)	methadone meperidine buprenorphine/naloxone (Suboxone™)
Neuroleptics (to treat psychotic disorders)	risperidone (Risperdal®, Risperdal Consta®) thioridazine quetiapine (Seroquel®)
NS3-4A Protease Inhibitors: (to treat Hepatitis C Virus [HCV])	boceprevir (Victrelis™) telaprevir (Incivek™) simeprevir (Galexos™)
PDE-5 Inhibitors (to treat erectile dysfunction)	sildenafil (Viagra®) vardenafil (Levitra®) tadalafil (Cialis®)
Sedatives/Hypnotics (to treat trouble with sleeping and/or anxiety)	buspirone (Bustab®) clorazepate diazepam (Diazemuls®, Valium®) estazolam flurazepam (Dalmane®, Som-Pam®) zoldipem midazolam (taken by injection)

Tell your doctor if you are taking any medicines that you obtained without a prescription.

This is **not** a complete list of medicines that you should tell your doctor that you are taking. Know and keep track of all the medicines you take and have a list of them with you. Show this list to all of your doctors and pharmacists any time you get a new medicine. Both your doctor and your pharmacist can tell you if you can take these other medicines with PREZISTA®. Do not start any new medicines while you are taking PREZISTA® without first talking with your doctor or pharmacist. You can ask your doctor or pharmacist for a list of medicines that can interact with PREZISTA®.

PROPER USE OF THIS MEDICATION

Always use PREZISTA® exactly as your doctor has told you. You must check with your doctor if you are not sure.

Usual dose:

Take PREZISTA® (tablets or oral suspension) every day exactly as prescribed by your doctor. You must take ritonavir (NORVIR®) at the same time as PREZISTA®.

Children, adolescents and adults who have difficulty swallowing PREZISTA® tablets should be prescribed PREZISTA® oral suspension.

Adults:

- For adults who have never taken anti-HIV medicines the usual dose is 800 mg (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) of PREZISTA®, together with 100 mg (one 100 mg capsule) of ritonavir (NORVIR®), once daily every day.

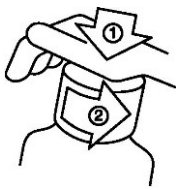
- For adults who have taken anti-HIV medicines in the past, the dose is either 800 mg (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) of PREZISTA® together with 100 mg (one 100 mg capsule) of ritonavir (NORVIR®), once daily *every day* **OR** 600 mg (one 600 mg tablet of 6 mL suspension) of PREZISTA® together with 100 mg (one 100 mg capsule) ritonavir (NORVIR®), twice daily *every day*. Please discuss with your doctor which dose is right for you.

Children:

- For children at least 3 years of age weighing at least 22 lbs (10 kg) who have taken anti-HIV medicines in the past, your child's doctor will decide the right dose based on your child's weight. Your child's doctor will inform you exactly on how much PREZISTA® (tablets or suspension) and how much ritonavir (NORVIR®) (capsules or solution) your child should take. In case your child does not tolerate ritonavir oral solution, ask your child's doctor for advice.

If your child is taking PREZISTA® oral suspension, shake the bottle well before each use. A 6 millilitre oral dosing pipette with 0.2 millilitre gradations is supplied with the pack so you can measure your dose accurately.PREZISTA® oral suspension should only be given with the supplied oral dosing pipette. The supplied oral dosing pipette should not be used for any other medication.

Removing the child-resistant cap



The amber glass bottle comes with a child-resistant cap and should be opened as follows:

- Push the plastic screw cap down while turning it counter clockwise.
- Remove the unscrewed cap.

If you have questions about when to take PREZISTA® and ritonavir (NORVIR®), your doctor can help you decide which schedule works for you.

You should always take PREZISTA® and ritonavir (NORVIR®) together with food. The type of food is not important.

Continue taking PREZISTA® and ritonavir (NORVIR®) unless your doctor tells you to stop. Take the exact amount of PREZISTA® and ritonavir (NORVIR®) that your doctor tells you to take, right from the very start. To help make sure you will benefit from PREZISTA® and ritonavir (NORVIR®), you must not skip doses or interrupt therapy. If you don't take PREZISTA® and ritonavir (NORVIR®) as prescribed, the beneficial effects of PREZISTA® and ritonavir (NORVIR®) may be reduced or even lost.

If you have also been prescribed enteric-coated didanosine as well as PREZISTA® and ritonavir, take didanosine 2 hours before or after the PREZISTA®/ritonavir combination.

Overdose:

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

Missed dose:

Patients taking 800 mg of PREZISTA® once daily:
If you miss a dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) or ritonavir (NORVIR®) by more than 12 hours, wait and then take the next dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) and ritonavir (NORVIR®) at the regularly scheduled time. If you miss a dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) or ritonavir (NORVIR®) by less than 12 hours, take your missed dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) and ritonavir (NORVIR®) immediately. Then take your next dose of PREZISTA® (two 400 mg tablets or one 800 mg tablet or 8 mL suspension) and ritonavir (NORVIR®) at the regularly scheduled time.

Patients taking 600 mg of PREZISTA® twice daily:

If you miss a dose of PREZISTA® or ritonavir (NORVIR®) by more than 6 hours, wait and then take the next dose of PREZISTA® and ritonavir (NORVIR®) at the regularly scheduled time. If you miss a dose of PREZISTA® or ritonavir (NORVIR®) by less than 6 hours, take your missed dose of PREZISTA®

and ritonavir (NORVIR®) immediately. Then take your next dose of PREZISTA® and ritonavir (NORVIR®) at the regularly scheduled time.

If a dose of PREZISTA® or ritonavir (NORVIR®) is skipped, do not double the next dose. Do not take more or less than your prescribed dose of PREZISTA® or ritonavir (NORVIR®) at any one time.

Do not stop using PREZISTA® without talking to your doctor first.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Like all prescription drugs, PREZISTA® can cause side effects. The following is **not** a complete list of side effects reported with PREZISTA® when taken either alone or with other anti-HIV medicines. Do not rely on this leaflet alone for information about side effects. Your doctor can discuss with you a more complete list of side effects.

Liver problems that may occasionally be severe have been reported. Your doctor should do blood tests prior to initiating PREZISTA®. If you have chronic hepatitis B or C infection, your doctor should check your blood tests more often because you have an increased chance of developing liver problems. Talk to your doctor about the signs and symptoms of liver problems. These may include yellowing of your skin or whites of your eyes, dark (tea coloured) urine, pale coloured stools (bowel movements), nausea, vomiting, loss of appetite, or pain, aching, or sensitivity on your right side below your ribs.

Rash has been reported in 10.3% of patients receiving PREZISTA®. In patients taking PREZISTA® and raltegravir, rashes (generally mild or moderate) may occur more frequently than in patients taking either drug separately. Contact your doctor immediately if you develop a rash. Your doctor will advise you whether your symptoms can be managed on therapy or whether PREZISTA® should be stopped.

In some patients, PREZISTA® has been reported to cause a severe or life-threatening rash. **If you develop a severe rash (e.g. blisters, peeling skin) which may be accompanied with symptoms such as fever, fatigue, muscle aches and pain, and liver problems, immediately discontinue PREZISTA® and contact your doctor.**

Other relevant severe side effects reported at an uncommon or rare frequency were inflammation of the liver or pancreas, increased blood fat levels, diabetes, and changes in body fat.

The most common side effects include diarrhea, nausea, headache, abdominal pain and vomiting.

Some side effects are typical for anti-HIV medicines in the same family as PREZISTA®. These are:

- high blood sugar (hyperglycemia) and diabetes. This can happen in patients taking PREZISTA® or other protease inhibitor medicines. Some patients have diabetes before starting treatment with PREZISTA®, which gets worse. Some patients get diabetes during treatment with PREZISTA®. Some patients will need changes in their diabetes medicine. Some patients may need new diabetes medicine.
- increased bleeding in patients with hemophilia. This may happen in patients taking PREZISTA® as it has been reported with other protease inhibitor medicines.
- changes in body fat. These changes can happen in patients taking anti-HIV medicines. The changes may include an increased amount of fat in the upper back and neck, breast, and around the back, chest, and stomach area. Loss of fat from the legs, arms, and face may also happen. The exact cause and long-term health effects of these conditions are not known.
- increases in triglycerides and cholesterol (forms of fat that are found in your blood). Your doctor may order blood testing for you.
- development of pancreatitis (inflammation of the pancreas) with symptoms such as abdominal pain, nausea and vomiting. If you suffer these symptoms while taking PREZISTA®, contact your doctor.
- immune reconstitution inflammatory syndrome. In some patients with advanced HIV infection (AIDS) and a history of opportunistic infection, signs and symptoms of inflammation from previous infections may occur soon after anti-HIV treatment is started. It is believed that these symptoms are due to an improvement in the body's immune response, enabling the body to fight infections that may have been present with no obvious symptoms. You could develop an autoimmune disease in which your immune system reacts against your own body (e.g., Graves' disease, which affects your thyroid gland, Guillain-Barré syndrome, which affects the nervous system, or polymyositis, which affects the muscles) and it may develop at any time, sometimes months after the start of HIV therapy.

Tell your doctor promptly about these or any other unusual symptoms. If the condition persists or worsens, seek medical attention.

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

Symptom / effect		Talk with your doctor or pharmacist		Stop taking drug and call your doctor or pharmacist
		Only if severe	In all cases	
Un-common	Severe and sometimes life-threatening rash (blisters, peeling skin) which may be accompanied by fever, fatigue, swelling of the face or lymph glands, muscle aches and pain, and liver problems.			✓
	Liver problems with symptoms such as yellowing of the skin or whites of the eyes, dark (tea coloured) urine, pale coloured stools (bowel movements), nausea, vomiting, loss of appetite, or pain, aching, or sensitivity on right side below ribs.		✓	
	Diabetes with symptoms such as excessive thirst, excessive urination, excessive eating, unexplained weight loss, poor wound healing, infections.		✓	
	Inflammation of the pancreas with symptoms such as abdominal pain, nausea and vomiting.		✓	

HOW TO STORE IT

Keep out of the reach and sight of children.

Store PREZISTA® oral suspension at room temperature between 15-30°C. Do not refrigerate or freeze. Avoid exposure to excessive heat. Store in the original container.

Store PREZISTA® tablets at room temperature between 15-30°C. Ask your doctor or pharmacist if you have any questions about storing your tablets.

This medication is prescribed for your particular condition. Do not use it for any other condition or give it to anybody else. Keep PREZISTA® and all of your medicines out of the reach of children. If you suspect that more than the prescribed dose of this medicine has been taken, contact your local poison control centre or emergency room immediately.

This leaflet provides a summary of information about PREZISTA®. If you have any questions or concerns about either PREZISTA® or HIV, talk to your doctor.

REPORTING SUSPECTED SIDE EFFECTS

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at www.healthcanada.gc.ca/medeffect
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
  - Fax toll-free to 1-866-678-6789, or
  - Mail to: Canada Vigilance Program
    Health Canada
    Postal Locator 0701E
    Ottawa, ON K1A 0K9

Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect® Canada Web site at www.healthcanada.gc.ca/medeffect.

NOTE: Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

This document plus the full product monograph, prepared for health professionals can be found at:
http://www.janssen.com/canada
or by contacting the sponsor, Janssen Inc., at:
1-800-567-3331 or 1-800-387-8781

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