
PREZISTA[®]

DATA SHEET

NAME OF THE DRUG

Darunavir

DESCRIPTION

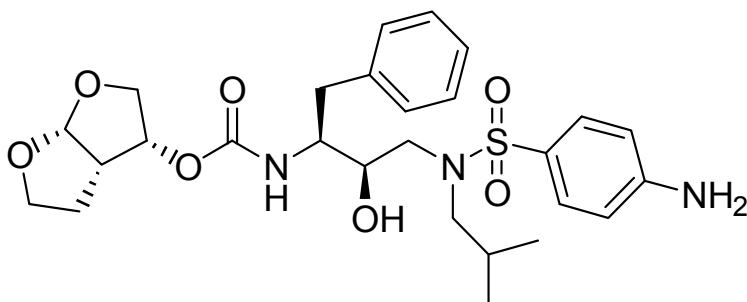
Antiviral for systemic use. Darunavir is an inhibitor of Human Immunodeficiency Virus type 1 (HIV-1) protease.

PREZISTA darunavir is available as 75 mg, 150 mg, 300 mg, 400 mg and 600 mg film-coated tablets (the 75 mg tablets are not currently marketed). Each film-coated tablet contains 75 mg, 150 mg, 300 mg, 400 mg or 600 mg darunavir, as 81.31 mg, 162.62 mg, 325.23 mg, 433.64 mg or 650.46 mg darunavir ethanolate, respectively. Inactive ingredients: microcrystalline cellulose, colloidal anhydrous silica, crospovidone, magnesium stearate, polyvinyl alcohol – partially hydrolysed, macrogol 3350, titanium dioxide (E171), talc, and (300 mg, 400 mg, 600 mg tablets only) sunset yellow FCF (E110).

Darunavir is isolated as darunavir ethanolate, a pseudo-polymorphic form of darunavir. Darunavir ethanolate is a white to off-white powder that is very slightly soluble in water, sparingly soluble in methanol, slightly soluble in ethanol, and freely soluble in acetone and dichloromethane.

The chemical name for darunavir is [(1*S*,2*R*)-3-[[[4-aminophenyl)sulfonyl]](2-methylpropyl)amino]-2-hydroxy-1-(phenylmethyl)propyl]-carbamic acid (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl ester.

Darunavir has the following chemical structure:



C₂₇H₃₇N₃O₇S

MW 547.66 daltons

CAS Registry No: 206361-99-1.

PHARMACOLOGY

Pharmacodynamics

Mechanism of action

Darunavir is an inhibitor of the dimerisation 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 binds to HIV-1 protease with a K_D of 4.5×10^{-12} M.

Darunavir was not a significant inhibitor of any of 13 tested human cellular proteases.

Antiviral activity *in vitro*

Darunavir exhibits activity against laboratory strains and clinical isolates of HIV-1 in acutely infected T-cell lines, human peripheral blood mononuclear cells and human monocytes/macrophages and laboratory strains of HIV-2 in acutely infected T-cell lines, 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 μ M to > 100 μ M.

The EC_{50} value of darunavir increased by a median factor of 5.4 in the presence of 50% human serum *in vitro*.

Darunavir showed synergistic antiviral activity when studied in combination with the protease inhibitors ritonavir, nelfinavir, or amprenavir and additive antiviral activity when studied in combination with the protease inhibitors 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, or efavirenz and the fusion inhibitor enfuvirtide. No antagonism was observed between darunavir and any of those antiretrovirals.

Resistance *in vitro*

In vitro selection of darunavir-resistant virus from wildtype HIV-1 was lengthy (> 3 years). The selected viruses were unable to grow in the presence of darunavir concentrations above 400 nM. Viruses selected in these conditions and showing decreased susceptibility to darunavir (range: 23 – 50-fold) harboured 2 to 4 amino acid substitutions in the protease gene.

In vitro selection of darunavir-resistant HIV-1 (range: 53 – 641-fold change in EC_{50} values [FC]) 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.

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 POWER 1 (TMC114-C213) and POWER 2 (TMC114-C202) trials and in the POWER 3 analysis (TMC114-C215 + TMC114-C208), only the subgroups with > 10 PI RAMs showed a median FC for darunavir > 10.

Cross-resistance *in vitro*

Cross-resistance has been observed among protease inhibitors. PREZISTA 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 PREZISTA.

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

Cross-resistance between PREZISTA and N(t)RTIs, NNRTIs or fusion inhibitors, is unlikely because the viral targets for those inhibitors are different.

Selection of viral resistance during PREZISTA/rtv therapy *in vivo*

In the 48 week analysis of the 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). In the virologic failures in the PREZISTA/rtv 800/100 mg q.d. group 7 subjects (12%) with developing PI RAMs were identified, compared to 4 subjects (10%) in the PREZISTA/rtv 600/100 mg b.i.d. group. One subject in the PREZISTA/rtv 800/100 mg q.d. group developed primary (i.e. major) PI mutations, which included 3 DRV RAMs, resulting in decreased susceptibility to darunavir. All the 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 was associated with a decreased susceptibility to a NRTI included in the treatment regimen.

In the 96 week analysis of the 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 (11.7% vs. 17.1%, respectively). In the virologic failures of the PREZISTA/rtv group, 3 patients with developing PI RAMs were identified. In the virologic failures of the lopinavir/rtv group, 5 patients with developing PI RAMs were identified. None of the developing mutations in the PREZISTA/rtv group or in the lopinavir/rtv group were primary (i.e. major) PI mutations. In 1 virologic failure of the PREZISTA/rtv group and 4 virologic failures of the lopinavir/rtv group, 1 developing NRTI RAM was identified. The development of the NRTI RAM at position 184 (n=4) was associated with a decreased susceptibility to FTC included in the background regimen.

In the 96 week analysis of the TITAN trial, the number of virologic failures was lower in the group of subjects receiving PREZISTA/rtv 600/100 mg b.i.d. than in subjects receiving lopinavir/ritonavir 400/100 mg b.i.d. (13.8% vs. 25.6%, respectively). Fewer virologic failures treated with PREZISTA/rtv 600/100 mg b.i.d. than with lopinavir/rtv 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 identified amino acid substitutions that developed on PREZISTA/rtv 600/100 mg b.i.d. 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.

Cross-resistance with other protease inhibitors *in vivo*

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

Of the viruses isolated from subjects 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 subjects, 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 subjects 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 subjects, 96% to 100% that were susceptible at baseline to amprenavir, atazanavir, indinavir, lopinavir, saquinavir or tipranavir 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.

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 or 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 PREZISTA/rtv 600/100mg b.i.d. groups of the POWER and DUET trials, the presence at baseline of three or more of the 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 1: Response (HIV-1 RNA < 50 copies/mL at Week 24) to PREZISTA/rtv 600/100 mg b.i.d. by baseline genotype* and by use of enfuvirtide: As-treated analysis of the POWER and DUET trials

Number of mutations at baseline*	All % n/N	No/non-naïve use of enfuvirtide % n/N	Naïve use of enfuvirtide % 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

*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 a predictive factor of virologic outcome. Response rates assessed by baseline darunavir phenotype are shown in the table below. The data are provided to give clinicians information on the likelihood of virologic success based on pre-treatment susceptibility to darunavir.

Table 2: 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 (fold change ranges)	All % n/N	No/non-naïve use of enfuvirtide % n/N	Naive use of enfuvirtide % 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

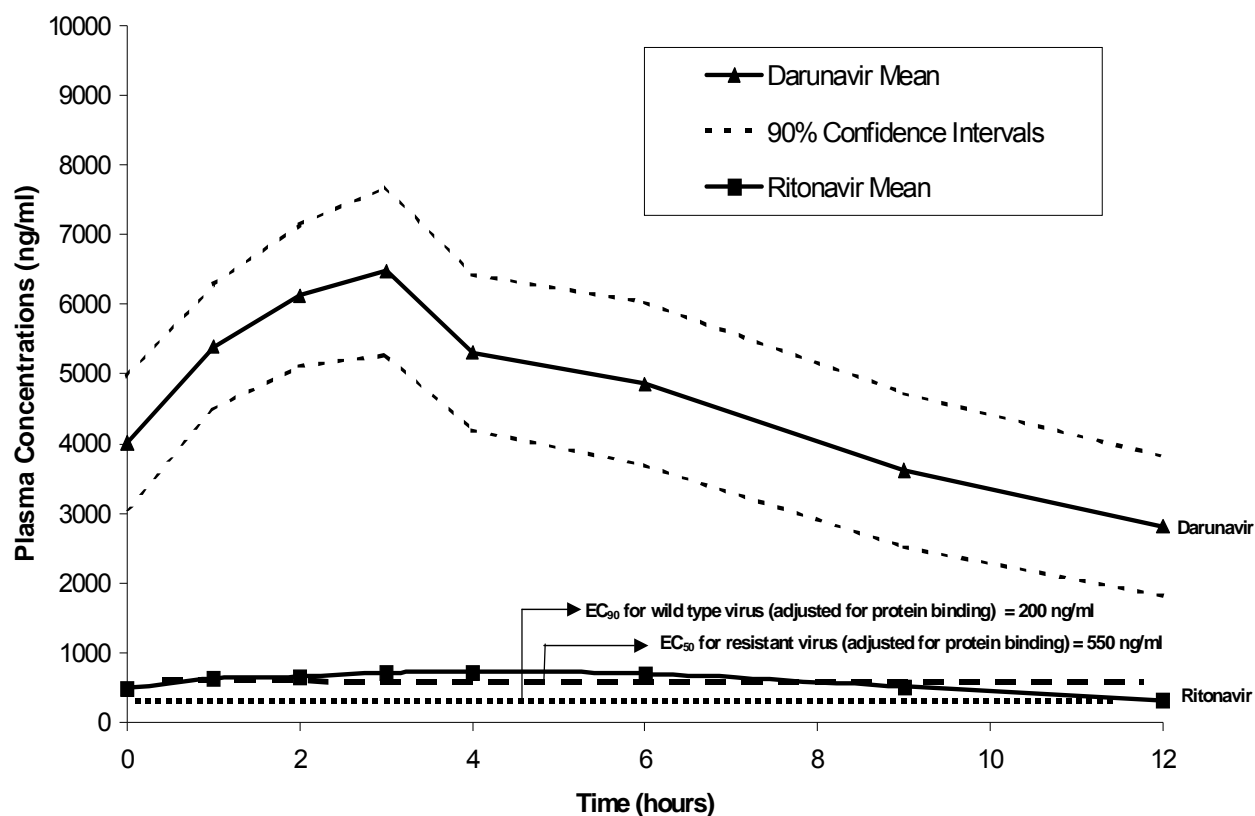
In deciding on a new regimen for patients who have failed an antiretroviral regimen, careful consideration should be given to treatment history of the individual patient and the patterns of mutations associated with different agents. When available, genotypic or phenotypic testing can be performed to guide the use of darunavir.

Pharmacokinetics

The pharmacokinetic properties of PREZISTA, 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 subjects.

Darunavir is primarily metabolized by cytochrome P₄₅₀ 3A (CYP3A). Ritonavir inhibits CYP3A, thereby increasing the plasma concentrations of darunavir considerably.

Figure 1: 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)[¶]



†† Mean plasma concentration-time profiles were derived from population pharmacokinetic analysis.

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 PREZISTA alone was approximately 37% and increased to approximately 82% in the presence of 100 mg b.i.d. ritonavir. 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 PREZISTA was given orally in combination with ritonavir at 100 mg b.i.d. (see PRECAUTIONS).

When administered without food, the relative bioavailability of PREZISTA in the presence of low-dose ritonavir is 30% lower, 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.

Distribution

Darunavir is approximately 95% bound to plasma protein. Darunavir binds primarily to plasma alpha-1-acid glycoprotein. The apparent volume of distribution of darunavir using population pharmacokinetic analysis was 122 l.

Metabolism

In vitro experiments with human liver microsomes indicate that darunavir primarily undergoes oxidative metabolism. Darunavir is extensively metabolised by the hepatic CYP system and primarily by isozyme CYP3A4. A ¹⁴C-darunavir trial in healthy volunteers showed that a majority of the radioactivity in plasma after a single 400/100 mg PREZISTA/rtv dose was due to the parent drug. 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 wildtype HIV.

Elimination

After a 400/100 mg ¹⁴C-darunavir/rtv dose, approximately 79.5% and 13.9% of the administered dose of ¹⁴C-darunavir could be retrieved in faeces and urine, respectively. Unchanged darunavir accounted for approximately 41.2% and 7.7% of the administered dose in faeces 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

Paediatrics

The pharmacokinetics of darunavir in combination with ritonavir in 74 treatment-experienced paediatric patients, aged 6 to < 18 years and weighing at least 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). Median (range) darunavir AUC_{12h} and C_{0h} values in this paediatric population were 63,670 (33,527; 115,360) ng.h/ml and 3,888 (1,836; 7,821) ng.h/ml, respectively.

Elderly

Population pharmacokinetic analysis in HIV-infected patients showed that PREZISTA pharmacokinetics are not considerably different in the age range (18 to 75 years) evaluated in HIV infected patients (n=12, age ≥ 65) (see PRECAUTIONS).

Gender

Population pharmacokinetic analysis showed a slightly higher darunavir exposure (16.8%) in HIV infected females compared to males. This difference is not clinically relevant.

Renal impairment

No pharmacokinetic data are available in subjects with severe renal impairment or end stage renal disease.

Results from a mass balance study with ¹⁴C-darunavir/rtv 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 PREZISTA has not been studied in patients with renal impairment, population pharmacokinetic analysis showed that the pharmacokinetics of PREZISTA were not significantly affected in HIV infected patients with moderate renal impairment (CrCl between 30-60 ml/min, n=20) (see DOSAGE AND ADMINISTRATION and PRECAUTIONS).

Hepatic impairment

Darunavir is primarily metabolised and eliminated by the liver. 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 subjects with mild (Child-Pugh Class A, n=8) and moderate (Child-Pugh Class B, n=8) hepatic impairment were comparable with those in healthy subjects. However, darunavir unbound concentrations were approximately 50% and 100% higher, respectively, in mild and moderate hepatic impairment, compared with those in healthy subjects. The clinical relevance of this increase in unbound darunavir concentrations is unknown. Therefore, PREZISTA should be used with caution in mild to moderate hepatic impairment. The effect of severe hepatic impairment on the pharmacokinetics of darunavir has not been studied (see DOSAGE AND ADMINISTRATION and PRECAUTIONS).

Drug Interactions:

Darunavir and ritonavir are both inhibitors of CYP3A. Co-administration of darunavir and ritonavir with drugs primarily metabolized by CYP3A may result in increased plasma concentrations of such drugs, which could increase or prolong their therapeutic effect and adverse events (see sections Contraindications, Precautions and Interactions with Other Drugs).

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 concentration of darunavir or drug are summarized in Table 8 (see Interactions with Other Drugs)

CLINICAL TRIALS

Efficacy of PREZISTA/rtv in treatment naïve adult patients

The evidence of efficacy of PREZISTA/rtv 800/100 mg once daily (q.d.) is based on the analyses of 96 week data from the randomised, controlled, open-label Phase III trial ARTEMIS (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. Randomisation 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.

Demographics and baseline characteristics were balanced between the PREZISTA/rtv arm and the lopinavir/rtv arm. 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 log₁₀ copies/ml and the median baseline CD4+ cell count was 228 x 10⁶ cells/l (range 4 – 750 x 10⁶ cells/l).

Table 3 shows the efficacy data of the 48 week analyses from the ARTEMIS trial.

Table 3: Efficacy data from the ARTEMIS trial (48 week analysis)

Outcomes	At week 48 ^a			At week 96 ^b		
	PREZISTA/rtv 800/100 mg q.d. N=343	lopinavir/rtv 800/200 mg per day N=346	Treatment difference (95% CI of difference)	PREZISTA/rtv 800/100 mg q.d. N=343	lopinavir/rtv 800/200 mg per day N=346	Treatment difference (95% CI of difference)
HIV-1 RNA < 50 copies/ml ^c	287 (83.7%)	271 (78.3%)	5.3 (-0.5; 11.2) ^d	271 (79.0%)	245 (70.8%)	8.2 (1.7; 14.7) ^d
HIV-1 RNA < 400 copies/ml ^c	301 (87.8%)	295 (85.3%)	2.5 (-2.6; 7.6) ^d	285 (83.1%)	268 (77.5%)	5.6 (-0.3; 11.6)
mean HIV-1 RNA log change from baseline (log ₁₀ copies/ml) ^e	-2.77	-2.65	-0.11 ^f (-0.30; 0.07) ^d	-2.64	-2.45	-0.20 ^f (-0.40; 0.01) ^d
median CD4+ cell count change from baseline (x 10 ⁶ /l) ^e	137	141		171	188	

- a) Data based on analysis at week 48
- b) Data based on analysis at week 96
- c) Imputations according to the TLOVR algorithm
- d) Based on normal approximation to the difference in % response
- e) Non-completer is failure imputation: patients who discontinued prematurely are imputed with a change equal to 0
- f) Difference in means

In the 48 week analysis, the virologic response (HIV-1 RNA < 50 copies/ml) for the PREZISTA/rtv arm was 83.7% and for the lopinavir/rtv arm 78.3% (Figure 2). Statistical comparisons between the treatment arms at week 48 confirmed non-inferiority of DRV/rvt versus lopinavir/rtv (p-value < 0.001) for both ITT (Intent-To-Treat) & OP (On Protocol) population.

Analyses of data at 96 weeks of treatment in the ARTEMIS trial demonstrated sustained antiretroviral efficacy and immunological benefit. In the 96 week analysis, virologic response (HIV-1 RNA < 50 copies/ml) was 79.0% and 70.8% for the PREZISTA/rtv arm and lopinavir/ritonavir arm, respectively. Non-inferiority in virologic response was demonstrated (p < 0.001) for both ITT and OP population, furthermore superiority of the PREZISTA/rtv arm over the lopinavir/rtv arm was demonstrated (p = 0.012 for the ITT population and p = 0.011 for the OP population).

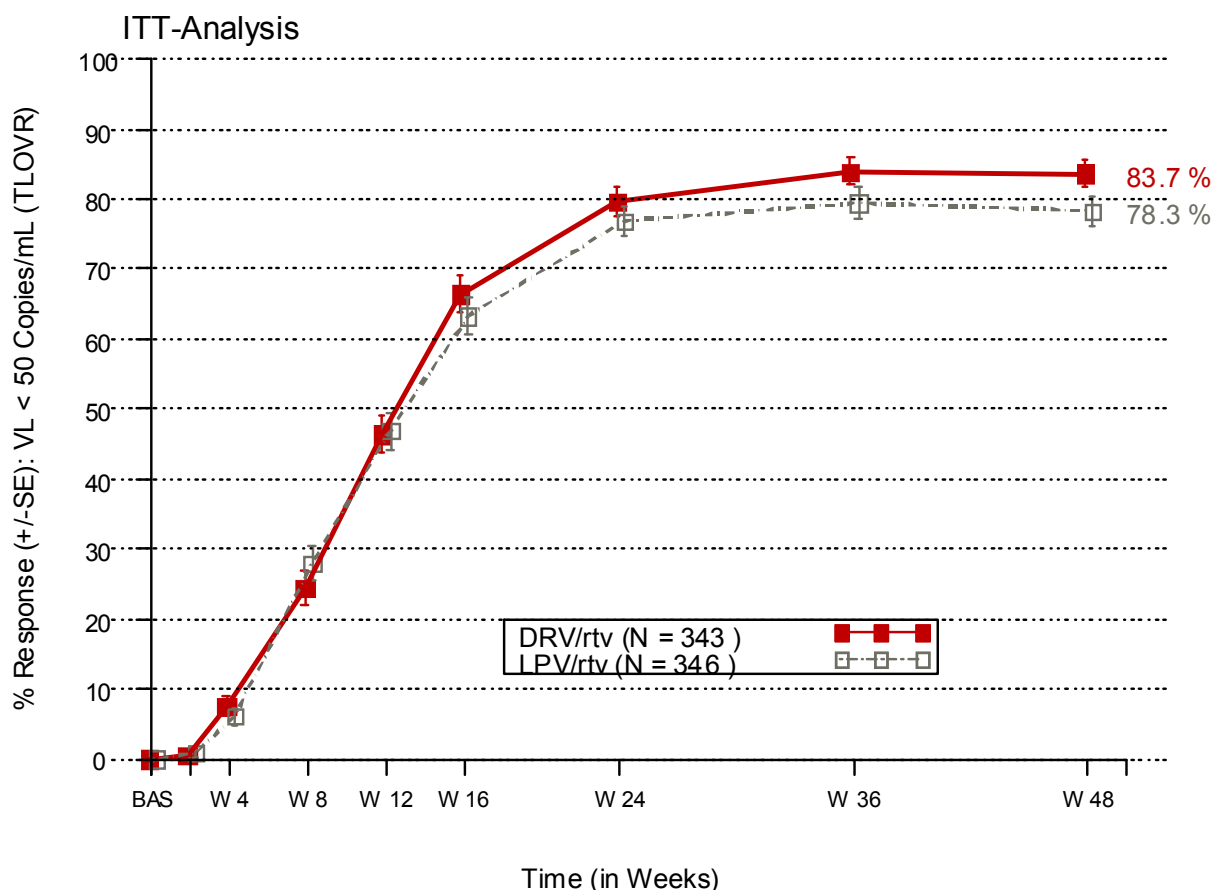


Figure 2. Virologic Response Over Time: Percentage of Subjects With Viral Load < 50 Copies/mL (ITT – TLOVR) – ARTEMIS

The virological response (< 50 copies/ml) at 96 weeks by baseline viral load and baseline CD4+ cell count is presented in Table 4.

Table 4. Virological response (HIV-1 RNA < 50 copies/ml) by baseline viral load

	PREZISTA/rtv 800/100 mg q.d. N=343	lopinavir/rtv 800/200 mg per day N=346	treatment difference

	N	number of responders at week 96 n (%)	N	number of responders at week 96 n (%)	Difference in % response (95% CI of difference in % response) ^{a)}
Baseline plasma viral load (copies/ml)					
< 100.000	226	182 (80.5%)	226	170 (75.2%)	5.3 (-2.3; 13.0)
≥ 100.000	117	89 (76.1%)	120	75 (62.5%)	13.6 (1.9; 25.3)
Baseline CD4+ cell count (x 10 ⁶ /l)					
< 200	141	111 (78.7%)	148	96 (64.9%)	13.9 (3.5; 24.2)
≥ 200	202	160 (79.2%)	198	149 (75.3%)	4.0 (-4.3; 12.2)

a) Based on a normal approximation to the difference in % response

Efficacy of PREZISTA/rtv (800/100 mg once daily) in treatment-experienced adult patients

The evidence of comparable efficacy of PREZISTA/rtv 800/100 mg once daily and PREZISTA/rtv 600/100 mg twice daily in treatment-experienced patients with no darunavir RAMs is based on the 48 week analysis of the Phase III trial ODIN.

ODIN is a randomised, open-label trial comparing PREZISTA/rtv 800/100 mg once daily to PREZISTA/rtv 600/100 mg twice daily in treatment-experienced HIV-1 infected patients with screening genotype resistance testing showing no darunavir RAMs (i.e. V11I, V32I, L33F, I47V, I50V, I54M, I54L, T74P, L76V, I84V, L89V) and a viral load of >1,000 HIV-1 RNA copies/ml. Both arms used an optimised background regimen consisting of ≥2 NRTIs selected by the investigator.

Demographics and baseline characteristics were balanced between the PREZISTA/rtv once daily arm and the PREZISTA/rtv twice daily arm. The 590 patients in total had a median age of 40 years (range 18-77), 64% were male, 36% white, 26% black, 18% hispanic, and 15% asian. The mean baseline plasma HIV-1 RNA was 4.16 log₁₀ copies/ml and the median baseline CD4+ cell count was 228 x 10⁶ cells/l (range 24 – 1306 x 10⁶ cells/l).

The table below shows the efficacy data of the 48 week analysis from the ODIN trial:

ODIN			
Outcomes	PREZISTA/rtv 800/100 mg once daily + OBR N=294	PREZISTA/rtv 600/100 mg twice daily + OBR N=296	Treatment difference (95% CI of difference)
HIV-1 RNA < 50 copies/ml ^a	212 (72.1%)	210 (70.9%)	1.2% (-6.1; 8.5) ^b
mean HIV-1 RNA log change from baseline (log ₁₀ copies/ml) ^c	-1.84	-1.80	-0.04 ^d (-0.24; 0.16)
mean CD4+ cell count change from	108	112	-5 ^d (-25; 16)

baseline (x 10 ⁶ /l) ^c			
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- a Imputations according to the TLOVR algorithm
- b Based on a normal approximation of the difference in % response
- c Last Observation Carried Forward imputation
- d Difference in means
- e NC=F

In the 48 week analysis, the virologic response defined as a confirmed plasma HIV-1 RNA viral load < 50 copies/ml, was 72.1% for the PREZISTA/rtv once daily arm and 70.9% for the PREZISTA/rtv twice daily arm. Statistical comparisons between the treatment arms at week 48 confirmed non-inferiority of PREZISTA/rtv once daily versus PREZISTA/rtv twice daily for both the ITT and OP population (p-value < 0.001).

Efficacy of PREZISTA/rtv (600/100 mg twice daily) in treatment experienced lopinavir-naïve adult patients

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 III trial TITAN (TMC114-C214) in treatment experienced, lopinavir/rtv naïve patients and on the analyses of 96 week data from the Phase IIb trials POWER 1, 2 and 3, in patients with high level of PI resistance.

TITAN is a randomised, controlled, open-label Phase III 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 optimised 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. The 298 patients on PREZISTA/rtv 600/100 mg b.i.d. had a median age of 40 years (range 18-68), 77% were male, 54% white, 18% black, 15% hispanic, and 9% asian. The mean baseline plasma HIV-1 RNA was 4.33 log₁₀ copies/ml and the median baseline CD4+ cell count was 235 x 10⁶ cells/l (range 3 – 831 x 10⁶ cells/l).

Table 5 shows the efficacy data of the 48 week and 96 week analyses from the TITAN trial.

Table 5: Efficacy data from the TITAN trial (48 week analysis)

Outcomes	At week 48 ^a			At week 96 ^b		
	PREZISTA/rtv 600/100 mg b.i.d. + OBR N=298	lopinavir/rtv 400/100 mg b.i.d. + OBR N=297	Treatment difference (95% CI of difference)	PREZISTA/rtv 600/100 mg b.i.d. + OBR N=298	lopinavir/rtv 400/100 mg b.i.d. + OBR N=297	Treatment difference (95% CI of difference)
HIV-1 RNA < 400 copies/ml ^c	228 (76.5%)	199 (67.0%)	9.5% (2.3; 16.7) ^d	199 (66.8%)	175 (58.9%)	7.9% (0.1; 15.6) ^d
HIV-1 RNA < 50 copies/ml ^c	211 (70.8%)	179 (60.3%)	10.5% (2.9; 18.1) ^d	180 (60.4%)	164 (55.2%)	5.2% (-2.8; 13.1) ^d

mean HIV-1 RNA log change from baseline (log ₁₀ copies/ml) ^e	-1.95	-1.72	-0.23 ^f (-0.44; -0.02) ^d	-1.71	-1.52	-0.19 ⁰ (-0.40; 0.03) ^d
median CD4+ cell count change from baseline (x 10 ⁶ /l) ^e	88	81		81	93	

- a) Data based on analyses at week 48
- b) Data based on analyses at week 96
- c) Imputations according to the TLOVR algorithm
- d) Based on a normal approximation of the difference in % response
- e) NC=F
- f) Difference in means

In the 48 week analysis, virologic response, defined as the percentage of subjects with plasma HIV-1 RNA level < 400 copies/ml, was 76.5% and 67.0% for the PREZISTA/rtv arm and lopinavir/rtv arm, respectively. Non-inferiority in virologic response was demonstrated (p < 0.001) for both the ITT (Figure 3) and OP population.

Analyses of data at 96 weeks of treatment in the TITAN trial demonstrated sustained antiretroviral efficacy and immunological benefit. In the 96 week analysis, virologic response, defined as the percentage of subjects with plasma HIV-1 RNA level < 400 copies/ml, was 66.8% and 58.9% for the PREZISTA/rtv arm and lopinavir/rtv arm, respectively. Non-inferiority in virologic response was demonstrated (p < 0.001) for both ITT and OP population, furthermore superiority of PREZISTA/rtv over the lopinavir/rtv arm was demonstrated (p = 0.034 for the ITT population and p = 0.033 for the OP population). 60.4% of patients on PREZISTA/rtv reached HIV-1 RNA less than 50 copies/ml versus 55.2% in the lopinavir/rtv arm.

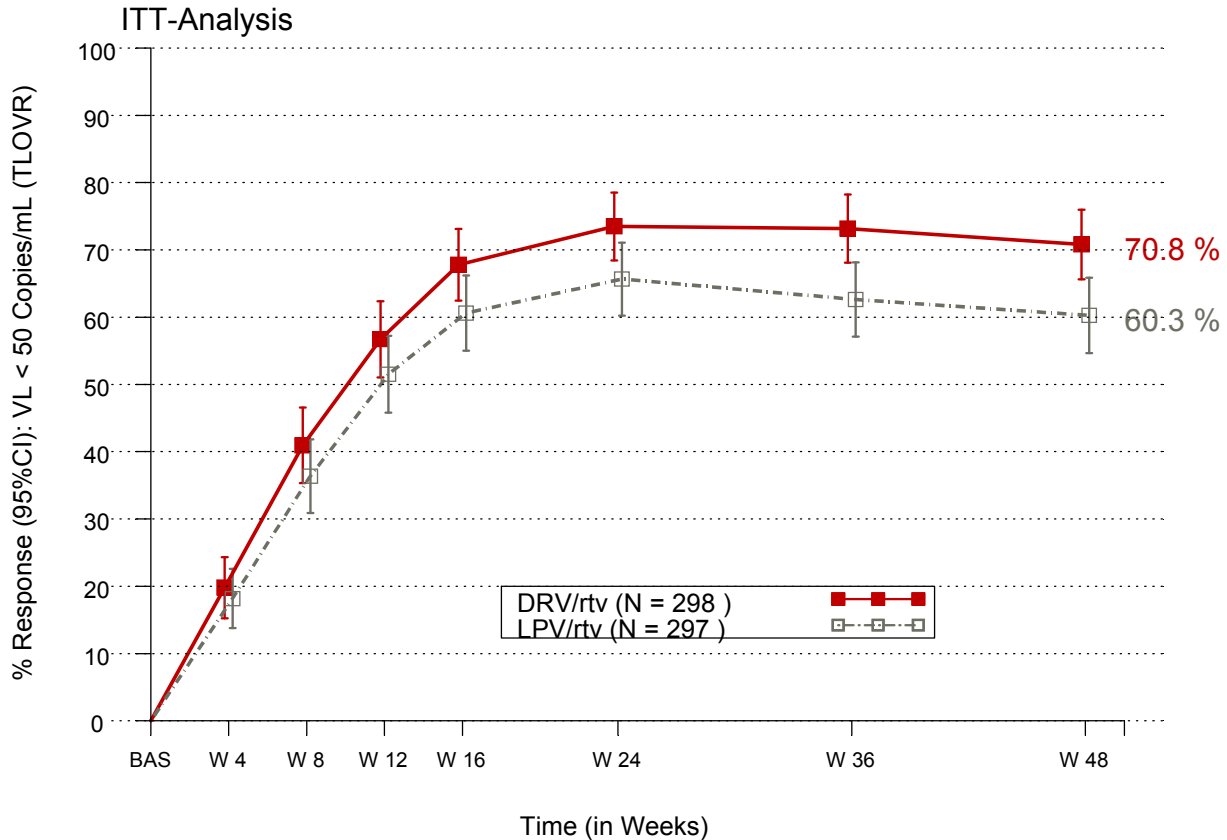


Figure 3. Virologic Response Over Time: Percentage of Subjects With Viral Load < 50 Copies/mL (ITT – TLOVR) – TITAN

Efficacy of PREZISTA/rtv in treatment experienced adult patients who failed more than one PI-containing regimen

POWER 1 (TMC114-C213) and POWER 2 (TMC114-C202) are randomised, 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 randomised 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 ribonucleic acid (RNA) > 1000 copies/ml, had prior treatment with PI(s), NNRTI(s) and NRTI(s), had at least 1 primary (i.e. major) PI mutation at screening and were on a stable PI-containing regimen at screening for at least 8 weeks. Randomisation was stratified by the number of PI mutations, screening viral load and the use of enfuvirtide.

Demographics and baseline characteristics were balanced between the PREZISTA/rtv arm and the comparator PI arm. In both trials combined, the 131 patients on PREZISTA/rtv 600/100 mg b.i.d. had a median age of 43 years (range 27-73), 89% were male, 81% white, 10% black and 7% hispanic. The mean baseline plasma HIV-1 RNA was 4.61 log₁₀ copies/ml and the median baseline CD4+ cell count was 153 x 10⁶ cells/l (range 3 – 776 x 10⁶ cells/l). The median darunavir FC was 4.3. In the PREZISTA/rtv 600/100 mg b.i.d. arm patients had prior exposure to a mean of 4 PIs, 5 NRTIs and 1 NNRTI versus 4 PIs, 6 NRTIs and 1 NNRTI in the comparator arm. Twenty percent of the patients in the PREZISTA/rtv arm had prior use of enfuvirtide versus 17% in the comparator arm.

The virologic response, defined as a decrease in plasma HIV-1 RNA viral load of at least 1.0 log₁₀ versus baseline, was evaluated in patients receiving PREZISTA/rtv plus an optimised background regimen (OBR) versus a control arm receiving an investigator-selected PI(s) regimen plus an OBR. The OBR consisted of at least 2 NRTIs with or without enfuvirtide (ENF). 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%. Twenty-three percent of the control patients used dual-boosted PIs. Approximately 47% of all patients used enfuvirtide and 35% of the use was in patients who were ENF-naïve.

Table 6 below shows the efficacy data of the 48 week and 96 week analyses from the pooled POWER 1 and POWER 2 trials

Table 6: Efficacy Outcomes at Weeks 48 and 96 of the Studies POWER 1 and POWER 2 (Pooled Analysis)

	Randomized Studies POWER 1 and POWER 2		
	PREZISTA/rtv 600 mg b.i.d. + OBR n=131	Comparator PI + OBR n=124	Treatment difference (95% CI of difference)
Week 48 time point			
HIV-1 RNA log ₁₀ mean change from baseline (log ₁₀ copies/ml) ^{a)}	-1.69	-0.37	-1.32 (-1.58; -1.05)
HIV-1 RNA ≥ 1 log ₁₀ below baseline ^{d)}	81 (61.8%)	20 (16.1%)	45.7% (35.0%; 56.4%) ^{e)}
HIV-1 RNA < 400 copies/ml ^{d)}	72 (55.0%)	18 (14.5%)	40.4% (29.8%; 51.1%) ^{e)}
HIV-1 RNA < 50 copies/ml ^{d)}	59 (45.0%)	14 (11.3%)	33.7% (23.4%; 44.1%) ^{e)}
CD4+ cell count mean change from baseline (x 10 ⁶ /l) ^{c)}	103	17	86 ^{b)} (57; 114)

Week 96 time point			
HIV-1 RNA log ₁₀ mean change from baseline (log ₁₀ copies/ml) ^{a)}	-1.58	-0.25	-1.33 (-1.59; -1.07)
HIV-1 RNA ≥ 1 log ₁₀ below baseline ^{d)}	74 (56.5%)	12 (9.7%)	46.8% (36.9%; 56.8%) ^{e)}
HIV-1 RNA < 400 copies/ml ^{d)}	65 (49.6%)	12 (9.7%)	39.9% (29.9%; 50.0%) ^{e)}
HIV-1 RNA < 50 copies/ml ^{d)}	51 (38.9%)	11 (8.9%)	30.1% (20.3%; 39.8%) ^{e)}
CD4+ cell count mean change from baseline (x 10 ⁶ /l) ^{c)}	133	15	118 ^{b)} (84; 152)

- a) Non-completer is failure imputation: patients who discontinued prematurely are imputed with a change equal to 0
- b) P-values < 0.001, based on the ANOVA model
- c) Last Observation Carried Forward imputation
- d) Imputations according to the TLOVR algorithm
- e) Confidence interval around observed differences of response rates; P-values < 0.001, based on the logistic regression model.

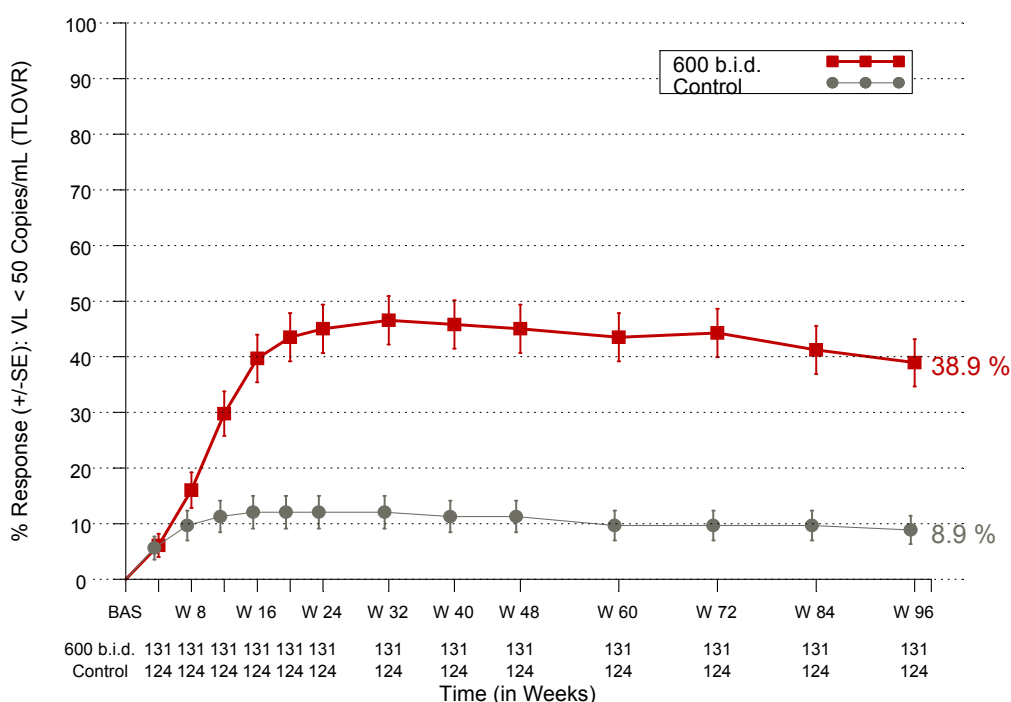


Figure 4: Virologic Response Over Time: Percentage of Subjects With Plasma Viral Load < 50 Copies/mL (ITT – TLOVR) in the pooled POWER 1 and POWER 2 trials

POWER 3

Additional data on the efficacy of PREZISTA/rtv 600/100 mg b.i.d. have been obtained in treatment-experienced adult subjects participating in the non-randomized trial TMC114-C215 (POWER 3). At week 48, 334 patients were included in the POWER 3 efficacy analysis who had 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 the TMC114-C215 analysis were the same as those for studies POWER 1 and POWER 2.

Baseline characteristics of the subjects included in the TMC114-C215/C208 analysis were comparable to those subjects in Studies POWER 1 and POWER 2.

The POWER 3 48-week efficacy analysis supported the viral load reduction and CD4+ cell count increases observed in the Studies POWER 1 and POWER 2. Of the 334 subjects at Week 48, 59% had a virologic response defined as a decrease of at least 1.0 log₁₀ in plasma viral load versus baseline and 46% of the subjects reached less than 50 HIV-1 RNA copies/mL.

Analyses of data through 96 weeks of treatment with PREZISTA/rtv (600/100 mg b.i.d.) in the POWER 3 study supported the sustained antiretroviral efficacy and immunological benefit as demonstrated in the studies POWER 1 and POWER 2. Of the 336 subjects at Week 96 in study POWER 3, 52.2% of patients had a virologic response defined as a decrease of at least 1 log₁₀ in HIV-1 RNA from baseline. 42.1% of the patients reached an HIV-1 RNA level < 50 copies/ml and 50.0% of patients reached less than 400 HIV-1 RNA copies/ml. The mean decrease in HIV-1 RNA level compared to baseline was 1.43 log₁₀ copies/ml and a mean increase in CD4+ cell count of 103 x 10⁶ cells/l was observed. Out of the 206 patients who responded with complete viral suppression (< 50 copies/ml) at week 48 in Studies POWER 1, POWER 2 and POWER 3, 177 patients (86% of the responders at week 48) remained responders at week 96.

Description of the clinical study in paediatric patients

DELPHI (TMC114-C212) is an open-label, Phase II trial evaluating the pharmacokinetics, safety, tolerability, and efficacy of PREZISTA/rtv in 80 antiretroviral treatment-experienced HIV-1 infected paediatric patients aged 6 to < 18 years and weighing at least 20 kg. At week 24, the virologic response rate was evaluated in paediatric patients receiving PREZISTA/rtv in combination with other antiretroviral agents (see DOSAGE AND ADMINISTRATION for dosage recommendations by body weight). Virologic response was defined as a decrease in plasma HIV-1 RNA viral load of at least 1.0 log₁₀ versus baseline. The mean baseline plasma HIV-1 RNA was 4.64 log₁₀ copies/ml, and the median baseline CD4+ cell count was 330 x 10⁶ cells/l (range: 6 to 1505 x 10⁶ cells/l).

In the study, 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 patients taking ritonavir oral solution, 23 switched to the 100 mg capsule formulation and exceeded the weight-based ritonavir dose without changes in observed safety.

At week 24, 73.8% of the paediatric patients had at least 1.0 log₁₀ HIV-1 RNA decrease from baseline. The proportion of paediatric patients reaching undetectable viral load (< 50 HIV-1 RNA copies/ml) was 50.0%, and the proportion of paediatric patients with < 400 HIV-1 RNA copies/ml was 63.8%. The mean change in plasma HIV-1 RNA from baseline was -1.98 log₁₀ copies/ml. The mean CD4+ cell count increase from baseline was 117 x 10⁶ cells/l.

INDICATIONS

Adult patients

PREZISTA (with low dose ritonavir as a pharmacokinetic enhancer) is indicated in combination with other antiretroviral agents for the treatment of human immunodeficiency virus-1 (HIV-1) infection in adult patients.

Paediatric patients

PREZISTA (with low dose ritonavir as a pharmacokinetic enhancer) is indicated in combination with other antiretroviral agents for the treatment of human immunodeficiency virus (HIV) infection in treatment-experienced paediatric patients aged 6 years and older.

CONTRAINDICATIONS

Hypersensitivity to darunavir or to any of the excipients.

Darunavir and ritonavir are both inhibitors of the CYP3A isoform. PREZISTA/rtv should not be co-administered with medicinal products that are highly dependent on CYP3A for clearance and for which increased plasma concentrations are associated with serious and/or life-threatening events (narrow therapeutic index). These medicinal products include astemizole, alfuzosin, sildenafil (when used for treatment of pulmonary arterial hypertension), terfenadine, midazolam, triazolam, cisapride, pimozide and the ergot alkaloids (e.g., ergotamine, dihydroergotamine, ergonovine and methylergonovine), antiarrhythmic drugs (e.g. amiodarone, bepridil, flecainide, systemic lidocaine, quinidine) (see Interactions with Other Drugs).

Due to the need for co-administration of PREZISTA with low dose ritonavir, please refer to ritonavir prescribing information for a description of ritonavir contraindications and precautions.

PRECAUTIONS

PREZISTA (darunavir) must be co-administered with ritonavir and food to exert its therapeutic effect (see DOSAGE and ADMINISTRATION). Failure to correctly administer PREZISTA with ritonavir and food will result in reduced plasma concentrations of darunavir that will be insufficient to achieve the desired therapeutic effect.

Patients should be advised that current antiretroviral therapy does not cure HIV and has not been proven to prevent the transmission of HIV. Appropriate precautions should continue to be employed.

Hepatotoxicity

Drug-induced hepatitis (e.g., acute hepatitis, cytolytic hepatitis) has been reported with PREZISTA/rtv. During the clinical development program (N=3063), hepatitis has been reported in 0.5% of patients receiving combination therapy with PREZISTA/rtv. Patients with pre-existing liver dysfunction, including chronic active hepatitis B or C, have an increased risk for liver function abnormalities including severe hepatic adverse events.

Post-marketing cases of liver injury, including some fatalities, have been reported. These have generally occurred in patients with advanced HIV-1 disease taking multiple concomitant medications, having co-morbidities including hepatitis B or C co-infection, and/or developing immune reconstitution syndrome. A causal relationship with PREZISTA/rtv therapy has not been established.

Appropriate laboratory testing should be conducted prior to initiating therapy with PREZISTA/rtv and patients should be monitored during treatment. Increased AST/ALT monitoring should be considered in patients with underlying chronic hepatitis, cirrhosis, or in patients who have pre-treatment elevations of transaminases, especially during the first several months of PREZISTA/rtv treatment.

If there is 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, hepatomegaly) in patients on PREZISTA/rtv, interruption or discontinuation of treatment must be considered.

Severe skin reactions

During the clinical development program (N=3,063), severe skin reactions, which may be accompanied with fever and/or elevations of transaminases, have been reported in 0.4% of patients. Stevens-Johnson Syndrome has been rarely (<0.1%) reported; and during post-marketing experience toxic epidermal necrolysis has 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 Effects). 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%.

Rash occurred more commonly in treatment-experienced subjects 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.

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.

Haemophilic patients

There have been reports of increased bleeding, including spontaneous skin haematomas and haemarthrosis in patients with haemophilia type A and B treated with PIs. In some patients additional factor VIII was given. In more than half of the reported cases, treatment with PIs was continued or reintroduced if treatment had been discontinued. A causal relationship has been suggested, although the mechanism of action has not been elucidated. Haemophilic patients should therefore be made aware of the possibility of increased bleeding.

Diabetes Mellitus/Hyperglycaemia

New onset diabetes mellitus, hyperglycaemia, or exacerbation of existing diabetes mellitus has been reported in patients receiving antiretroviral therapy, including PIs. In some of these patients the hyperglycaemia was severe and in some cases also associated with ketoacidosis. Many patients had confounding medical conditions some of which required therapy with agents that have been associated with the development of diabetes mellitus or hyperglycaemia.

Fat redistribution & metabolic disorders

Combination antiretroviral therapy has been associated with redistribution of body fat (lipodystrophy) in HIV infected patients. The long-term consequences of these events are currently unknown. Knowledge about the mechanism is incomplete. A connection between visceral lipomatosis and PIs and lipoatrophy and NRTIs has been hypothesized. A higher risk of lipodystrophy has been associated with individual factors such as older age, and with drug related factors such as longer duration of antiretroviral treatment and associated metabolic disturbances. Clinical examination should include evaluation for physical signs of fat redistribution. Consideration should be given to measurement of fasting serum lipids and blood glucose. Lipid disorders should be managed as clinically appropriate (see ADVERSE REACTIONS).

Immune reconstitution inflammatory syndrome

In HIV infected patients with severe immune deficiency at the time of institution of combination antiretroviral therapy (CART), an inflammatory reaction to asymptomatic or residual opportunistic pathogens may arise and cause serious clinical conditions, or aggravation of symptoms. Typically, such reactions have been observed within the first weeks or months of initiation of CART. Relevant examples are cytomegalovirus retinitis, generalised and/or focal mycobacterial infections, and *Pneumocystis carinii* pneumonia. Any inflammatory symptoms should be evaluated and treatment instituted when necessary.

Elderly

As limited information is available on the use of PREZISTA/rtv in patients aged 65 and over, caution should be exercised in the administration of PREZISTA in elderly patients, reflecting the greater frequency of decreased hepatic function and of concomitant disease or other therapy (see Pharmacokinetics).

Use in children

PREZISTA/rtv should not be used in children below 3 years of age in view of toxicity 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 Use in Lactation).

The safety and efficacy of PREZISTA/rtv in antiretroviral treatment-experienced children aged 3 to less than 6 years and in antiretroviral treatment naïve paediatric patients have not been established.

Patients with Co-existing Conditions

Patients with Hepatic Impairment

The safety and efficacy of PREZISTA have not been established in patients with severe hepatic impairment. Therefore, PREZISTA should not be used in patients with severe hepatic impairment. Due to an increase in unbound darunavir plasma concentrations, PREZISTA should be used with caution in patients with mild or moderate hepatic impairment (see Pharmacokinetics and DOSAGE AND ADMINISTRATION).

Patients with Renal Impairment

There are no pharmacokinetic data available in HIV-1 infected patients with severe renal impairment or end stage renal disease. 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 haemodialysis or peritoneal dialysis (see Pharmacokinetics and DOSAGE AND ADMINISTRATION).

Carcinogenicity

PREZISTA 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. Systemic exposures at the highest dose (based on plasma AUC) were approximately 0.5-fold (mice) and 0.75-fold (rats) relative to humans at the recommended therapeutic dose of darunavir/ritonavir (600/100 mg b.i.d).

The incidences of hepatocellular adenomas were statistically significantly increased at all doses in male mice; at the mid and high dose in female mice and male rats; and at the high dose in female rats. The incidence of hepatocellular carcinomas was significantly increased at the high dose in male mice; and male and female rats. The relevance of these findings for humans is limited. An increase in the incidence of thyroid follicular cell adenomas was noted in male rats. This is considered rodent specific and of no relevance to humans.

Genotoxicity

PREZISTA 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.

Effects on fertility

In a study conducted in rats, there were no effects on mating with PREZISTA treatment up to 1000 mg/kg/day, but exposure levels were below (AUC - 0.5 fold) that in humans at the clinically recommended dose. The number of corpora lutea and hence the number of live young was lower for females at 1000 mg/kg/day PREZISTA, and correlated with lower maternal body weight; the NOEL for effects on fertility was 200 mg/kg/day PREZISTA (corresponding to an exposure level 0.3 – fold that in humans at the recommended clinical dose).

Use in Pregnancy

Category B2.

There are no adequate and well controlled studies with darunavir in pregnant women. PREZISTA/ritonavir should be used during pregnancy only if the potential benefit justifies the potential risk.

In animal studies with PREZISTA treatment up to 1000 mg/kg/day, there was no teratogenicity with darunavir in mice, rats and rabbits when treated alone nor in mice when treated in combination with ritonavir. However, the exposure levels in mice and rats were about half those with the recommended clinical dose in humans, and only 5% in rabbits. In a pre- and post-natal rat study the pups had lower birth weight following maternal treatment with 1000 mg/kg/day darunavir.

Use in Lactation

It is not known whether darunavir is excreted in human milk. Studies in rats have demonstrated that darunavir is excreted in milk. Because of both the potential for HIV transmission and the potential for serious adverse events in breast-feeding infants, mothers should be instructed not to breastfeed if they are receiving PREZISTA.

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-metabolising enzymes approaches adult values by 3 years of age.

Interactions with Other Drugs

Darunavir and ritonavir are both inhibitors of the CYP3A isoform. Co-administration of PREZISTA and ritonavir and drugs primarily metabolized by CYP3A may result in increased plasma concentrations of such drugs, which could increase or prolong their therapeutic effect and adverse events (see Tables 7 and 8).

Darunavir is metabolised by CYP3A. Medicinal products that induce CYP3A activity would be expected to increase the clearance of darunavir, resulting in lower plasma concentrations of darunavir. Co-administration of darunavir and other medicinal products that inhibit CYP3A may decrease the clearance of darunavir and may result in increased plasma concentrations of darunavir.

Drugs that are contraindicated and not recommended for concomitant administration with PREZISTA/rtv are included in Table 7. 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 7: Drugs that should not be concomitantly administered with PREZISTA/rtv

Drug class: drug name	Clinical comment
Alpha blocker: alfuzosin	Concentrations of alfuzosin may be increased when coadministered with PREZISTA/rtv.
Antiarrhythmics: bepridil, flecainide, lidocaine (systemic), quinidine, amiodarone	Concentrations of bepridil, flecainide, lidocaine, quinidine and amiodarone may be increased when coadministered with PREZISTA/rtv. Caution is warranted and therapeutic concentration monitoring, if available, is recommended for antiarrhythmics when coadministered with PREZISTA/rtv.
Antihistamines: astemizole, terfenadine	Contraindicated due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
Antimycotics: rifampicin	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 to PREZISTA.
Ergot Derivatives: dihydroergotamine, ergonovine, ergotamine, methylergonovine	Contraindicated 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 Agent: cisapride	Contraindicated due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
Herbal Products: St. John's wort (<i>Hypericum perforatum</i>)	PREZISTA/rtv should not be used concomitantly with products containing St. John's wort (<i>Hypericum perforatum</i>) because coadministration may cause significant decreases in darunavir plasma concentrations. This may result in loss of therapeutic effect to PREZISTA.
Neuroleptic: pimozide	Contraindicated due to the potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
Sedative/Hypnotics: midazolam, triazolam	Contraindicated due to potential for serious and/or life-threatening reactions such as prolonged or increased sedation or respiratory depression.
HIV-Protease Inhibitor: lopinavir/ritonavir	Results of interaction trials with PREZISTA with or without ritonavir and lopinavir/ritonavir (1200 mg PREZISTA 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%. This may significantly affect the therapeutic effect of PREZISTA in HIV-1 infected patients. The appropriate doses of the combination have not been established. Hence, it is not recommended to co-administer PREZISTA/rtv with lopinavir/ritonavir.
HIV-Protease Inhibitor: saquinavir	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 coadministered with saquinavir and ritonavir; saquinavir exposure was not affected when administered concomitantly with darunavir/ritonavir. It is not recommended to coadminister saquinavir and PREZISTA, with or without low-dose ritonavir.

Drug class: drug name	Clinical comment
Anticonvulsants: phenobarbital, phenytoin	Phenobarbital and phenytoin are inducers of CYP450 enzymes. PREZISTA/rtv should not be used in combination with these medicines, as coadministration may cause significant decreases in darunavir plasma concentrations. This may result in loss of therapeutic effect to PREZISTA.
HMG-CoA Reductase Inhibitors: lovastatin, simvastatin	HMG-CoA reductase inhibitors, such as lovastatin and simvastatin, which are highly dependent on CYP3A metabolism, are expected to have markedly increased plasma concentrations when coadministered with darunavir/ritonavir. Increased concentrations of HMGCoA reductase inhibitors may cause myopathy, including rhabdomyolysis. Concomitant use of PREZISTA/rtv with lovastatin or simvastatin is not recommended. For information regarding atorvastatin and pravastatin, see Table 8.
PDE-5 inhibitors for pulmonary arterial hypertension: sildenafil	A safe and effective dose of sildenafil for the treatment of pulmonary arterial hypertension has not been established. There is an increased potential for sildenafil-associated adverse events (including visual disturbances, hypotension, prolonged erection and syncope). Therefore, co-administration of PREZISTA/rtv with sildenafil when used for pulmonary arterial hypertension is contraindicated. For information regarding tadalafil, see Table 8.

Table 8: Established and other potentially significant drug interactions: Alteration in dose or regimen may be recommended based on drug interaction studies or predicted interactions

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
<i>HIV-Antiviral agents: Non-nucleoside Reverse Transcriptase Inhibitors (NNRTIs)</i>		
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.

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
Etravirine	↔ darunavir ↓ etravirine	A pharmacokinetic interaction study between darunavir/rtv and etravirine in healthy subjects indicated that etravirine has no significant effect on the pharmacokinetics of darunavir. In this study, when 100 mg b.i.d. of etravirine was co-administered with 600/100 mg b.i.d. darunavir/rtv a 37 % decrease in etravirine plasma levels was observed. However, when 200 mg b.i.d. etravirine was co-administered, exposure was increased by 80% compared with etravirine alone. Based on the results of this study, a dose adjustment for darunavir is not considered necessary when co-administered with 200 mg etravirine and 100 mg rtv.
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. Since this difference is not considered to be clinically relevant, the combination of PREZISTA/rtv and nevirapine can be used without dose adjustments.
<i>HIV-Antiviral Agents: Nucleoside Reverse Transcriptase Inhibitors (NRTIs)</i>		
Didanosine	↔ 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. As it is recommended that didanosine be administered on an empty stomach, didanosine should be administered one hour before or two hours after PREZISTA/rtv (which are administered with food).

Drug class: drug name		Effect on concentration of darunavir or drug	Clinical comment
Tenofovir fumarate	disoproxil	↔ 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.
<i>HIV-Antiviral agents: Protease Inhibitors (PIs)</i>			
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 low dose ritonavir as a pharmacokinetic enhancer (see CLINICAL PHARMACOLOGY and PRECAUTIONS).
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 coadministered. Atazanavir can be coadministered 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 coadministered 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 from 800 mg b.i.d. to 600 mg b.i.d. may be warranted in case of intolerance.
<i>Other Agents</i>			

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
Antiarrhythmic: 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 (LSM) was 1.77 with a 90% CI of 0.90 to 3.50). The pharmacokinetics of digoxin were significantly influenced by PREZISTA/rtv. Therefore, it is recommended that the lowest possible dose of digoxin should initially be prescribed in case digoxin is given to patients on PREZISTA/rtv therapy. Given that digoxin has a narrow therapeutic index, the digoxin dose should be carefully titrated to obtain the desired clinical effect while assessing the overall clinical state of the subject.
Anticoagulant: warfarin	↓ warfarin ↔ darunavir	Warfarin concentrations may be affected when coadministered with PREZISTA/rtv. It is recommended that the international normalized ratio (INR) be monitored when warfarin is combined with PREZISTA/rtv.
Anticonvulsant: carbamazepine	↑ carbamazepine ↔ darunavir	An interaction trial between PREZISTA/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 carbamazepine dose may need to be reduced by 25% to 50% in the presence of PREZISTA/rtv.
Anti-infective: clarithromycin	↑ 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, a dose reduction of clarithromycin should be considered.

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
Antifungals: ketoconazole, itraconazole, voriconazole	↑ ketoconazole ↑ darunavir ↑ itraconazole (not studied) ↑ voriconazole (not studied)	<p>Ketoconazole, itraconazole, and voriconazole are potent inhibitors as well as substrates of CYP3A. Concomitant systemic use of ketoconazole, itraconazole, or voriconazole and darunavir and ritonavir may increase plasma concentrations of darunavir. Simultaneously, plasma concentrations of ketoconazole, itraconazole, or voriconazole may be increased by darunavir and ritonavir. This was confirmed in an interaction trial where the 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 coadministration is required, the daily dose of ketoconazole or itraconazole should not exceed 200 mg. Plasma concentrations of voriconazole may be decreased in the presence of darunavir/ritonavir. Voriconazole should not be administered to patients receiving PREZISTA/rtv unless an assessment of the benefit/risk ratio justifies the use of voriconazole.</p>
Antimycobacterial: rifabutin	↑ rifabutin ↓ darunavir	<p>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-desacetylrifabutin. A dosage reduction of rifabutin by 75% of the usual dose of 300 mg/day (i.e. rifabutin 150 mg q.o.d.) and increased monitoring for rifabutin-related adverse events is warranted in patients receiving the combination.</p>
Beta agonists: salmeterol	↑ salmeterol	<p>Concomitant use of salmeterol and PREZISTA/rtv is not recommended. The combination may result in increased risk of cardiovascular adverse events with salmeterol, including QT prolongation, palpitations and sinus tachycardia.</p>

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
Calcium Channel Blockers: felodipine, nifedipine, nicardipine	↑ calcium channel blockers	Plasma concentrations of calcium channel blockers (e.g. felodipine, nifedipine, nicardipine) may increase when PREZISTA/rtv are coadministered. Caution is warranted and clinical monitoring of patients is recommended.
Corticosteroid: dexamethasone, fluticasone propionate	↓ darunavir ↑ fluticasone propionate	Use with caution. Systemic dexamethasone induces CYP3A and can thereby decrease darunavir plasma concentrations. This may result in loss of therapeutic effect to PREZISTA. Concomitant use of inhaled fluticasone propionate and PREZISTA/rtv may increase plasma concentrations of fluticasone propionate. Alternatives should be considered, particularly for long term use.
Endothelin receptor antagonist: bosentan	↑ bosentan	Concomitant use of bosentan and PREZISTA/rtv may increase plasma concentrations of bosentan. In patients who have been receiving PREZISTA/rtv for at least 10 days, start bosentan at 62.5 mg q.d. or q.o.d. based upon individual tolerability. For patients on bosentan and initiating PREZISTA/rtv, discontinue the 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 q.d. or q.o.d. based upon individual tolerability.
Gout therapy: colchicine	↑ colchicine	Concomitant use of colchicine and PREZISTA/rtv may increase the exposure to colchicine. The following dose adjustments are recommended for colchicine. For the treatment of gout-flares in patients on PREZISTA/rtv, the recommended dose of colchicine is 0.6 mg (1 tablet), followed by 0.3 mg (half tablet) 1 hour later. Treatment course to be repeated no earlier than 3 days. For the prophylaxis of gout-flares in patients on PREZISTA/rtv, the recommended dose of colchicine is 0.3 mg q.d. or q.o.d. For the treatment of familial Mediterranean fever in patients on PREZISTA/rtv, the maximum dose of colchicine is 0.6 mg q.d. (may be given as 0.3 mg b.i.d.). Patients with renal or hepatic impairment should not be given colchicine with PREZISTA/rtv.
Oestrogen-based Contraceptive: ethinyl oestradiol and norethindrone	↓ ethinyl oestradiol and norethindrone	The results of an interaction trial between PREZISTA/rtv (600/100 mg b.i.d.) and ethinyl oestradiol and norethindrone demonstrated that at steady-state systemic exposures to ethinyl oestradiol and norethindrone are decreased by 44% and 14%, respectively. Therefore, alternative methods of non-hormonal contraception are recommended.

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
HMG-CoA Reductase Inhibitors: atorvastatin, pravastatin	↑ HMG-CoA reductase inhibitors	<p>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 coadministered 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. An interaction trial between darunavir/ritonavir (600/100mg b.i.d.) and pravastatin (40mg, single dose) demonstrated that darunavir/ritonavir did not increase exposure to a single dose of pravastatin in most subjects but up to 5-fold in a limited subset of subjects. When administration of pravastatin and PREZISTA co-administered with low dose ritonavir is required, it is recommended to start with the lowest possible dose of pravastatin and titrate it up to the desired clinical effect while monitoring for safety.</p> <p>An interaction study evaluating PREZISTA/rtv (600/100 mg b.i.d.) in combination with rosuvastatin (10 mg q.d.) resulted in an increase in rosuvastatin exposure. It is recommended to start with the lowest possible dose of rosuvastatin and titrate up to the desired clinical effect while monitoring for safety.</p>
H ₂ -Receptor Antagonists and Proton Pump Inhibitors: omeprazole, ranitidine	↔ darunavir	<p>Coadministration 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 coadministered with H₂-receptor antagonists and proton pump inhibitors without dose adjustments.</p>
Immunosuppressants: cyclosporin, tacrolimus, sirolimus	↑ immuno-suppressants	<p>Plasma concentrations of cyclosporine, tacrolimus or sirolimus may be increased when coadministered with PREZISTA/rtv. Therapeutic concentration monitoring of the immunosuppressive agent is recommended for immunosuppressant agents when coadministered with PREZISTA/rtv.</p>

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
Narcotic Analgesic: methadone, buprenorphine/naloxone	↔ methadone, ↔ buprenorphine /naloxone	<p>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.</p> <p>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.</p>

Drug class: drug name	Effect on concentration of darunavir or drug	Clinical comment
PDE-5 inhibitors: sildenafil, vardenafil, tadalafil	↑ PDE-5 inhibitors	<p>Treatment of erectile dysfunction: 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 coadministered with darunavir (400 mg b.i.d.) and low-dose ritonavir (100 mg b.i.d.). Concomitant use of PDE-5 inhibitors with PREZISTA/rtv should be done with caution. If concomitant use of PREZISTA/rtv with sildenafil, vardenafil, or tadalafil is indicated, sildenafil at a single dose not exceeding 25 mg in 48 hours, vardenafil at a single dose not exceeding 2.5 mg dose in 72 hours, or tadalafil at a single dose not exceeding 10 mg dose in 72 hours, is recommended.</p> <p>Treatment of pulmonary arterial hypertension: co-administration of PREZISTA/rtv with sildenafil when used for pulmonary arterial hypertension is contraindicated (see Table 7). For the treatment of pulmonary arterial hypertension with tadalafil co-administered with PREZISTA/rtv, a dose adjustment for tadalafil is warranted. In patients who have been receiving PREZISTA/rtv for at least 1 week, start tadalafil at 20 mg q.d., and increase to 40 mg q.d. based upon individual tolerability. For patients on tadalafil and initiating PREZISTA/rtv, discontinue the use of tadalafil at least 24 hours prior to initiating PREZISTA/rtv and avoid the use of tadalafil during the initiation of PREZISTA/rtv. After at least 1 week following the initiation of PREZISTA/rtv, resume tadalafil at 20 mg q.d. and increase to 40 mg q.d. based upon individual tolerability.</p>
Selective Serotonin Reuptake Inhibitors (SSRIs): sertraline, paroxetine	↔ darunavir ↓ sertraline ↓ paroxetine	<p>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 coadministered with darunavir and ritonavir. If sertraline or paroxetine is coadministered 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.</p>

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

Effects on Laboratory Tests

None known.

Effect on Ability to Drive or Operate Machinery

No trials on the effects of PREZISTA in combination with ritonavir on the ability to drive or use machines have been performed. However, dizziness has been reported in some patients during treatment with regimens containing PREZISTA/rtv and should be borne in mind when considering a patient's ability to drive or operate machinery (see ADVERSE REACTIONS).

ADVERSE EFFECTS

The overall safety profile of PREZISTA is based on all available clinical trial and post-marketing data, and is consistent with the data presented below. Adverse drug reactions to PREZISTA/rtv identified in the safety assessment of the clinical trials in adults.

Adverse drug reactions to PREZISTA/rtv (800/100 mg once daily) identified in antiretroviral treatment naïve patients:

The safety assessment is based on all safety data from the Phase III trial ARTEMIS comparing PREZISTA/rtv 800/100 mg once daily versus lopinavir/ritonavir 800/200 mg per day in antiretroviral naïve HIV-1 infected adult patients. The total patient years exposure in the PREZISTA/rtv arm and the lopinavir/rtv arm was 626.6 and 608.1, respectively.

The majority of the ADRs reported during treatment with PREZISTA/rtv were mild in severity.

The most frequent ($\geq 5\%$) ADRs of moderate to severe (grade 2-4) intensity were diarrhoea and headache and abdominal pain.

The most frequent ($> 1\%$) ADRs of severe (grade 3 or 4) intensity were related to laboratory abnormalities. All other grade 3 or 4 ADRs were reported in less than 1% of the patients.

2.3% percent of the patients in the PREZISTA/rtv arm discontinued treatment due to ADRs.

Adverse Drug Reactions to PREZISTA/rtv 800/100 mg once daily of at least moderate intensity (grade 2-4) in antiretroviral treatment naïve HIV-1 infected adult patients are presented in the table below*:

System Organ Class Adverse Drug Reaction	PREZISTA/rtv 800/100 mg once daily. TDF/FTC# N=343	lopinavir/rtv 800/200 mg per day + TDF/FTC# N=346
Nervous system disorders Headache	5.8%	4.6%

System Organ Class Adverse Drug Reaction	PREZISTA/rtv 800/100 mg once daily. + TDF/FTC# N=343	lopinavir/rtv 800/200 mg per day + TDF/FTC# N=346
Gastrointestinal disorders		
Abdominal pain	5.2%	5.8%
Acute pancreatitis	0.3%	0.3%
Diarrhoea	7.6%	14.7%
Dyspepsia	0.3%	0%
Flatulence	0.9%	0.9%
Nausea	2.6%	3.5%
Vomiting	1.5%	3.2%
Skin and subcutaneous tissue disorders		
Angioedema [†]	0.3%	0%
Lipodystrophy (lipohypertrophy, lipodystrophy and lipoatrophy)	0.3%	0.6%
Pruritus	0.9%	0.6%
Rash	1.7%	4.0%
Stevens-Johnson Syndrome	0.3%	0%
Urticaria [†]	0.9%	0.3%
Musculoskeletal and connective tissue disorders		
Myalgia	0.6%	1.2%
Metabolism and nutrition disorders		
Anorexia	1.5%	0.9%
Diabetes mellitus	0.6%	0.6%
General disorders and administration site conditions		
Asthenia	0.9%	0%
Fatigue	0.3%	2.6%
Immune system disorders		
(Drug) Hypersensitivity [†]	0.6%	1.4%
Immune reconstitution syndrome	0.3%	0.3%
Hepatobiliary disorders		
Hepatitis acute, cytolytic hepatitis, hepatotoxicity	0.3%	0.6%
Psychiatric disorders		
Abnormal dreams	0.3%	0.3%

* Excluding laboratory abnormalities reported as ADRs

Tenofovir disoproxil fumarate/emtricitabine

† Adverse drug reactions identified from post-marketing experience

Laboratory abnormalities, grade 2-4, considered ADRs, in antiretroviral treatment naïve HIV-1 infected adult patients are shown in the table below*:

Laboratory parameter	Limit	PREZISTA/rtv 800/100 mg once daily. + TDF/FTC# N=343	lopinavir/rtv 800/200 mg per day + TDF/FTC# N=346

Laboratory parameter	Limit	PREZISTA/rtv 800/100 mg once daily. TDF/FTC [#] N=343	lopinavir/rtv 800/200 mg per day + TDF/FTC [#] N=346
ALT			
Grade 2	> 2.5 to ≤ 5.0 x ULN	7.3%	6.1%
Grade 3	> 5.0 to ≤ 10.0 x ULN	2.9%	2.6%
Grade 4	> 10.0 x ULN	0.9%	2.9%
AST			
Grade 2	> 2.5 to ≤ 5.0 x ULN	6.1%	6.1%
Grade 3	> 5.0 to ≤ 10.0 x ULN	4.1%	1.8%
Grade 4	> 10.0 x ULN	1.2%	2.3%
ALP			
Grade 2	> 2.5 to ≤ 5.0 x ULN	1.5%	1.2%
Grade 3	> 5.0 to ≤ 10.0 x ULN	0%	0.3%
Grade 4	> 10.0 x ULN	0%	0%
Triglycerides			
Grade 2	500-750 mg/dl	2.6%	7.9%
Grade 3	751-1200 mg/dl	1.2%	4.7%
Grade 4	> 1200 mg/dl	0.6%	0.9%
Total cholesterol*			
Grade 2	240-300 mg/dl	16.4%	23.0%
Grade 3	> 300 mg/dl	1.2%	4.7%
LDL cholesterol*			
Grade 2	160-190 mg/dl	13.5%	9.6%
Grade 3	≥ 191 mg/dl	4.7%	5.0%
Elevated glucose levels			
Grade 2	126-250 mg/dl	7.3%	7.6%
Grade 3	251-500 mg/dl	0.9%	0%
Grade 4	> 500 mg/dl	0%	0%
Pancreatic lipase			
Grade 2	> 1.5 to ≤ 3.0 x ULN	1.8%	1.2%
Grade 3	> 3.0 to ≤ 5.0 x ULN	0.6%	0.6%
Grade 4	> 5.0 x ULN	0%	0.6%
Pancreatic amylase			
Grade 2	> 1.5 to ≤ 2.0 x ULN	4.7%	1.7%
Grade 3	> 2.0 to ≤ 5.0 x ULN	2.6%	2.9%
Grade 4	> 5.0 x ULN	0%	0.6%

* Grade 4 data not applicable in Division of AIDS grading scale

Tenofovir disoproxil fumarate/emtricitabine

Adverse drug reactions to PREZISTA/rtv(600/100 mg twice daily) identified in antiretroviral treatment experienced adult patients – TITAN study

The safety assessment is based on all safety data from the Phase III trial TITAN 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 patient years of exposure in the PREZISTA/rtv arm and the lopinavir/rtv arm was 462.5 and 436.1, respectively.

The majority of the ADRs reported during treatment with PREZISTA/rtv were mild in severity. The most frequent ($\geq 5\%$) ADRs of moderate to severe (grade 2-4) intensity were diarrhoea, hypertriglyceridaemia, hypercholesterolaemia, nausea, abdominal pain, vomiting, lipodystrophy, hepatic enzymes increased and rash. The most frequent ($> 1\%$) severe (grade 3 or 4) ADRs were lipodystrophy or related to laboratory abnormalities. All other grade 3 or 4 ADRs were reported in less than 1% of the patients. 4.7 percent of the patients discontinued treatment due to ADRs.

Adverse Drug Reactions to PREZISTA/rtv 600/100 mg b.i.d. of at least moderate intensity (grade 2-4) in antiretroviral treatment experienced HIV-1 infected adult patients in the TITAN trial are mentioned in the table below*:

System Organ Class Adverse Drug Reaction	PREZISTA/rtv 600/100 mg b.i.d. + OBR[#] N=298	lopinavir/rtv 400/100 mg b.i.d. + OBR[#] N=297
Nervous system disorders		
Headache	2.7%	3.0%
Gastrointestinal disorders		
Abdominal distension	2.0%	0.3%
Abdominal pain	5.7%	2.7%
Acute pancreatitis	0.3%	0.3%
Diarrhoea	14.4%	19.9%
Dyspepsia	2.0%	1.0%
Flatulence	0.3%	1.0%
Nausea	7.0%	6.4%
Vomiting	5.4%	2.7%
Skin and subcutaneous tissue disorders		
Lipodystrophy (lipohypertrophy, lipodystrophy, and lipoatrophy)	5.4%	4.4%
Pruritus	1.0%	1.0%
Rash	5.0%	2.0%
Urticaria [†]	0.3%	0%
Musculoskeletal and connective tissue disorders		
Myalgia	1.0%	0.7%
Metabolism and nutrition disorders		
Anorexia	1.7%	2.0%
Diabetes mellitus	1.7%	0.3%
General disorders and administration site conditions		
Asthenia	3.4%	1.0%
Fatigue	2.0%	1.3%
Immune system disorders		
Immune reconstitution syndrome	0.3%	0%
Reproductive system and breast disorders		
Gynaecomastia	0.3%	0.3%
Psychiatric disorders		
Abnormal dreams	0.7%	0%

* Excluding laboratory abnormalities reported as ADRs

Optimised Background Regimen

† Adverse drug reactions identified from post-marketing experience

Laboratory abnormalities, grade 2-4, considered ADRs, in antiretroviral treatment experienced HIV-1 infected adult patients in the TITAN trial are shown in the table below*:

Laboratory parameter	Limit	PREZISTA/rtv 600/100 mg b.i.d. + OBR# N=298	lopinavir/rtv 400/100 mg b.i.d. + OBR# N=297
ALT			
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%
AST			
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%
ALP			
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%
Triglycerides			
Grade 2	500-750 mg/dl	10.4%	11.4%
Grade 3	751-1200 mg/dl	6.9%	9.7%
Grade 4	> 1200 mg/dl	3.1%	6.2%
Total cholesterol*			
Grade 2	240-300 mg/dl	24.9%	23.2%
Grade 3	> 300 mg/dl	9.7%	13.5%
LDL cholesterol*			
Grade 2	160-190 mg/dl	114.4%	13.5%
Grade 3	≥ 191 mg/dl	7.7%	9.3%
Elevated glucose levels			
Grade 2	126-250 mg/dl	10.0%	11.4%
Grade 3	251-500 mg/dl	1.4%	0.3%
Grade 4	> 500 mg/dl	0.3%	0%
Pancreatic lipase			
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%

* Grade 4 data not applicable in Division of AIDS grading scale

Optimised Background Regimen

Adverse Drug Reactions to PREZISTA/rtv (600/100 mg twice daily) identified in adults in the pooled trials POWER 1, 2 and 3

In the pooled POWER trials, the total patient years of exposure was 812.4 in patients who immediately started treatment on PREZISTA/rtv 600/100 mg b.i.d. (See CLINICAL TRIALS)

The majority of the ADRs reported during treatment with PREZISTA/rtv were mild in severity. The most frequent (≥5%) moderate to severe (grade 2 – 4) ADRs were diarrhoea, headache, abdominal pain, nausea and vomiting. The most frequent grade 3 or 4 ADRs were increased hepatic and pancreatic enzymes, hypertriglyceridaemia, diarrhoea, hypercholesterolaemia, headache, abdominal pain and vomiting. All other grade 3 or 4 ADRs were reported in less than 1% of the patients.

2.1 percent of the patients discontinued treatment due to ADRs.

Adverse Drug Reactions to PREZISTA/rtv 600/100 mg b.i.d. of at least moderate intensity (grade 2-4) in antiretroviral treatment experienced HIV-1 infected adult patients in the pooled trials POWER 1, 2 and 3, are mentioned in the table below¹:

System Organ Class Adverse Drug Reaction	PREZISTA/rtv 600/100 mg b.i.d. + OBR² (n=467)
Nervous system disorders Headache	8.8%
Gastrointestinal disorders Abdominal distension Abdominal pain Acute pancreatitis Diarrhoea Dyspepsia Flatulence Nausea Vomiting	1.9% 6.4% 0.4% 13.7% 1.5% 1.5% 6.2% 5.6%
Skin and subcutaneous tissue disorders Lipodystrophy (lipohypertrophy, lipodystrophy, and lipoatrophy) Pruritus Rash Urticaria [†]	3.0% 2.6% 3.4% 0.6%
Musculoskeletal and connective tissue disorders Myalgia Osteonecrosis [†]	3.2% 0.6%
Metabolism and nutrition disorders Anorexia Diabetes mellitus	2.4% 1.3%
General disorders and administration site conditions Asthenia Fatigue	3.6% 3.9%
Immune system disorders (Drug) hypersensitivity[†] Immune reconstitution syndrome	0.6% 0.2%
Hepatobiliary disorders Hepatitis acute, cytolytic hepatitis, hepatotoxicity	0.4%
Reproductive system and breast disorders Gynaecomastia	0.9%
Psychiatric disorders Abnormal dreams	0.4%

¹Excluding laboratory abnormalities reported as ADRs.

²Optimised Background Regimen.

[†] Adverse drug reactions identified from post-marketing experience

Laboratory abnormalities, considered ADRs, in antiretroviral treatment experienced HIV-1 infected adult patients in the pooled trials POWER 1, 2 and 3 are shown in the table below:

Laboratory parameter Preferred Term	Limit	PREZISTA/rtv 600/100 mg b.i.d. + OBR¹ N=467
ALT		

Laboratory parameter Preferred Term	Limit	PREZISTA/rtv 600/100 mg b.i.d. + OBR ¹ N=467
Grade 2	> 2.5 to ≤ 5.0 x ULN	6.1%
Grade 3	> 5.0 to ≤ 10.0 x ULN	2.4%
Grade 4	> 10.0 x ULN	0.9%
AST		
Grade 2	> 2.5 to ≤ 5.0 x ULN	6.9%
Grade 3	> 5.0 to ≤ 10.0 x ULN	3.0%
Grade 4	> 10.0 x ULN	0.6%
ALP		
Grade 2	> 2.5 to ≤ 5.0 x ULN	3.9%
Grade 3	> 5.0 to ≤ 10.0 x ULN	0.9%
Grade 4	> 10.0 x ULN	0%
Triglycerides		
Grade 2	500 – 750 mg/dl	9.3%
Grade 3	751 – 1200 mg/dl	8.2%
Grade 4	> 1200 mg/dl	3.9%
Total cholesterol ²		
Grade 2	240 – 300 mg/dl	17.7%
Grade 3	> 300 mg/dl	7.1%
LDL cholesterol ²		
Grade 2	160 – 190 mg/dl	13.2%
Grade 3	≥ 191 mg/dl	9.1%
Elevated glucose levels		
Grade 2	126 – 250 mg/dl	15.4%
Grade 3	251 – 500 mg/dl	1.7%
Grade 4	> 500 mg/dl	0.2%
Pancreatic lipase		
Grade 2	> 1.5 to ≤ 3.0 x ULN	5.2%
Grade 3	> 3.0 to ≤ 5.0 x ULN	2.6%
Grade 4	> 5.0 x ULN	0.9%
Pancreatic amylase		
Grade 2	> 1.5 to ≤ 2.0 x ULN	7.4%
Grade 3	> 2.0 to ≤ 5.0 x ULN	7.8%
Grade 4	> 5.0 x ULN	1.1%

¹ Optimised Background Regimen.

² Grade 4 data not applicable in Division of AIDS grading scale.

Additional adverse drug reactions to PREZISTA/rtv identified in adult patients in other clinical studies

System Organ Class	Adverse Drug Reaction	Incidence*
Musculoskeletal and connective tissue disorders	Osteonecrosis ⁺	0.4%

* Incidence of at least grade 2 ADRs, calculated on pooled data of phase IIb and III trials (N=3,063)

⁺ Adverse drug reactions identified from post-marketing experience

Adverse drug reactions to PREZISTA/rtv identified in paediatric patients

The safety assessment in children and adolescents is based on the safety data from the Phase II trial DELPHI in which 80 antiretroviral treatment-experienced HIV-1 infected paediatric patients aged from 6 to < 18 years and weighing at least 20 kg received PREZISTA/rtv in combination with other antiretroviral agents (see CLINICAL TRIALS). Frequency, type and severity of adverse drug reactions in children and adolescents were comparable to those observed in adults.

Postmarketing experience

Adverse drug reactions identified during post-marketing experience.

System Organ Class	Adverse Drug Reaction	Incidence
Skin and subcutaneous tissue disorders	Toxic Epidermal Necrolysis	Very rare

Rarely, events of rhabdomyolysis (associated with co-administration with HMG-CoA reductase inhibitors and PREZISTA) have been reported.

Effects of combination antiretroviral therapy

Combination antiretroviral therapy has been associated with redistribution of body fat (lipodystrophy) in HIV patients, including loss of peripheral and facial subcutaneous fat, increased intra-abdominal and visceral fat, breast hypertrophy and dorsocervical fat accumulation (buffalo hump).

Combination antiretroviral therapy has also been associated with metabolic abnormalities such as hypertriglyceridaemia, hypercholesterolaemia, insulin resistance, hyperglycaemia and hyperlactataemia.

In HIV infected patients with severe immune deficiency at the time of initiation of combination antiretroviral therapy, an inflammatory reaction to asymptomatic or residual opportunistic infections may arise.

There have been reports of increased spontaneous bleeding in haemophilia patients receiving PIs.

Increased CPK, myalgia, myositis and rarely, rhabdomyolysis have been reported with the use of protease inhibitors, particularly in combination with NRTIs.

Special Populations

Patients co-infected with hepatitis B and/or hepatitis C virus

In patients co-infected with hepatitis B or C virus receiving PREZISTA/rtv, the incidence of adverse events and clinical chemistry abnormalities was not higher than in patients receiving PREZISTA/rtv who were not co-infected, except for increased hepatic enzymes. The pharmacokinetic exposure in co-infected patients was comparable to that in patients without co-infection. Increased AST/ALT monitoring should be considered in patients with hepatitis co-infection, especially during the first months of PREZISTA/rtv therapy.

DOSAGE AND ADMINISTRATION

PREZISTA must always be given with low dose 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.

Do not halve tablets. Dose equivalence when tablets are divided has not been established.

After therapy with PREZISTA has been initiated, patients should be advised not to alter the dosage or discontinue therapy without consulting their physician.

Adults

Antiretroviral treatment-naïve patients

The recommended dosage of PREZISTA is 800 mg once daily (q.d.) taken with ritonavir 100 mg q.d. and with food.

Antiretroviral treatment-experienced patients

For antiretroviral treatment experienced patients, genotypic testing is recommended. However, when genotypic testing is not feasible, the PREZISTA/rtv once daily dosing regimen is recommended in protease inhibitor naïve patients and the twice daily dosing regimen (see below) is recommended in protease inhibitor experienced patients.

When genotypic testing results are available, the recommended dosing is:

Antiretroviral treatment-experienced patients	
with no darunavir resistance associated mutations (DRV-RAMs)*	with at least one darunavir resistance associated mutation (DRV-RAM)*
800 mg PREZISTA once daily (q.d.) taken with 100 mg ritonavir and with food	600 mg PREZISTA twice daily (b.i.d.) taken with 100 mg ritonavir and with food

* DRV-RAMs: V11I, V32I, L33F, I47V, I50V, I54M, I54L, T74P, L76V, I84V and L89V

The type of food does not affect the exposure to darunavir. Ritonavir (100 mg) is used as a pharmacokinetic enhancer of darunavir (see Interactions with Other Drugs and Pharmacokinetics).

Paediatric patients

Antiretroviral treatment-experienced paediatric patients (6 to < 18 years of age)

The recommended dose of PREZISTA/rtv for paediatric patients (6 to < 18 years of age and weighing at least 20 kg) is based on body weight (see Table 11) and should not exceed the recommended adult dose (600/100 mg b.i.d.). PREZISTA tablets should be taken with ritonavir twice daily and with food. The type of food does not affect exposure to darunavir.

Table 11. Recommended dose for treatment-experienced paediatric patients (6 to < 18 years of age) for PREZISTA tablets and ritonavir (See Clinical Trials)

Body weight (kg)	Dose
≥ 20 kg–< 30 kg	375 mg PREZISTA/50 mg ritonavir b.i.d.
≥ 30 kg–< 40 kg	450 mg PREZISTA/60 mg ritonavir b.i.d.
≥ 40 kg	600 mg PREZISTA/100 mg ritonavir b.i.d.

Antiretroviral treatment-experienced children less than 6 years of age and antiretroviral treatment naïve paediatric patients

The safety and efficacy of PREZISTA/rtv in antiretroviral treatment-experienced children aged 3 to less than 6 years and in antiretroviral treatment naïve paediatric patients have not been established.

PREZISTA/rtv should not be used in children below 3 years of age (see PRECAUTIONS).

Hepatic impairment: No dose adjustment is required in patients with mild (Child-Pugh Class A) or moderate (Child-Pugh Class B) hepatic impairment, however PREZISTA should be used with caution in these patients due to increased darunavir exposure and adverse events. No pharmacokinetic data are available in patients with severe hepatic impairment. Therefore, PREZISTA should not be used in patients with severe hepatic impairment (see PRECAUTIONS).

Renal impairment: No dose adjustment is required in patients with renal impairment (see PRECAUTIONS).

This product is not able to deliver all approved dose regimens.

OVERDOSAGE

Human experience of acute overdose with PREZISTA/rtv is limited. Single doses up to 3200 mg of the oral solution of PREZISTA alone and up to 1600 mg of the tablet formulation of PREZISTA in combination 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. Since darunavir is highly protein bound, dialysis is unlikely to be beneficial in significant removal of the active substance. Contact the Poisons Information Centre (telephone 0800 POISON or 0800 764 766) for advice on management of overdosage.

PRESENTATION

PREZISTA 75 mg film-coated tablet (not currently marketed): White caplet -shaped tablet, debossed with 75 on one side and TMC on the other side.

PREZISTA 150 mg film-coated tablet: White oval-shaped tablet, debossed with 150 on one side and TMC on the other side.

PREZISTA 300 mg film-coated tablet: Orange oval shaped tablet, debossed with 300MG on one side and TMC114 on the other side.

PREZISTA 400 mg film-coated tablet: Light orange oval shaped tablet, debossed with 400MG on one side and TMC on the other side.

PREZISTA 600 mg film-coated tablet: Orange oval shaped tablet, debossed with 600MG on one side and TMC on the other side.

PREZISTA 75 mg film-coated tablets (not currently marketed) are provided in high density polyethylene (HDPE) plastic bottles containing 480 tablets, fitted with polypropylene (PP) child resistant closures.

PREZISTA 150 mg film-coated tablets are provided in high density polyethylene (HDPE) plastic bottles containing 240 tablets, fitted with polypropylene (PP) child resistant closures.

PREZISTA 300 mg film-coated tablets are provided in high density polyethylene (HDPE) plastic bottles containing 120 tablets, fitted with polypropylene (PP) child resistant closures.

PREZISTA 400 and 600 mg film-coated tablets are provided in high density polyethylene (HDPE) plastic bottles containing 60 tablets, fitted with polypropylene (PP) child resistant closures.

This product is not able to deliver all approved dose regimens.

Storage

Store below 30°C.

Shelf Life

2 years.

MEDICINE CLASSIFICATION

Prescription Medicine

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