PRODUCT NAME
EDURANT® (rilpivirine) 25 mg film-coated tablets

DOSAGE FORMS AND STRENGTHS
White to off-white, film-coated, round, biconvex, tablet of 6.4 mm, debossed with "TMC" on one side and "25" on the other side.

Each film-coated tablet contains rilpivirine hydrochloride equivalent to 25 mg rilpivirine.

For a full list of excipients, see *List of Excipients*.

CLINICAL INFORMATION
Indications
Adult and Pediatric Patients (12 years old and above)
EDURANT®, in combination with other antiretroviral medicinal products, is indicated for the treatment of human immunodeficiency virus type 1 (HIV-1) infection in antiretroviral treatment-naïve patients with a viral load ≤ 100,000 HIV-1 RNA copies/mL at the start of therapy.

Dosage and Administration
EDURANT® must always be given in combination with other antiretroviral medicinal products.

Dosage – Adult and Pediatric patients (12 years old and above)
The recommended dose of EDURANT® is one 25 mg tablet taken orally once daily. EDURANT® **must be taken** with a meal (see *Pharmacokinetic Properties*).

Dose adjustment
For patients concomitantly receiving rifabutin, the EDURANT® dose should be increased to 50 mg (two tablets of 25 mg each) once daily, taken with a meal. When rifabutin co-administration is stopped, the EDURANT® dose should be decreased to 25 mg once daily, taken with a meal (see *Interaction with Other Medicinal Products and Other Forms of Interaction*).

Special populations
Pediatrics (less than 12 years of age)
The safety and efficacy of EDURANT® in children less than 12 years of age are under investigation (see *Pharmacokinetic Properties*). Treatment with EDURANT® is not recommended in children less than 12 years of age.

Pregnancy and Postpartum
Lower exposures of rilpivirine were observed during pregnancy, therefore viral load should be monitored closely (see *Pregnancy, Breastfeeding and Fertility and Pharmacokinetic Properties Special Populations – Pregnancy and Postpartum*).

Elderly (65 years of age and older)
There is limited information regarding the use of EDURANT® in patients >65 years of age. No dose adjustment of EDURANT® is required in elderly patients (see *Pharmacokinetic Properties*). EDURANT® should be used with caution in this population.

Renal Impairment
EDURANT® has mainly been studied in patients with normal renal function. No dose adjustment of EDURANT® is required in patients with mild or moderate renal impairment. In patients with severe renal impairment or end-stage renal disease, EDURANT® should be used with caution. In patients with severe renal impairment or end-stage renal disease, the combination of EDURANT® with a strong CYP3A inhibitor (e.g., ritonavir-boosted HIV protease inhibitor) should only be used if the benefit outweighs the risk (see *Pharmacokinetic Properties*).

Treatment with EDURANT® resulted in an early small increase of mean serum creatinine levels which remained stable over time and is not considered clinically relevant (see *Adverse Reactions*).

Hepatic Impairment
There is limited information regarding the use of EDURANT® in patients with mild or moderate hepatic impairment (Child-Pugh score A or B). No dose adjustment of EDURANT® is required in patients with mild or moderate hepatic impairment. EDURANT® should be used with caution in patients with moderate hepatic impairment. EDURANT® has not been studied in patients with severe hepatic impairment (Child-Pugh score C). Therefore, EDURANT® is not recommended in patients with severe hepatic impairment (see *Pharmacokinetic Properties*).

Timing of Dosing
If the patient misses a dose of EDURANT® within 12 hours of the time it is usually taken, the patient should take EDURANT® with a meal as soon as possible and then take the next dose of EDURANT® at the regularly scheduled time. If a patient misses a dose of EDURANT® by more than 12 hours, the patient should not take the missed dose, but resume the usual dosing schedule.

Contraindications
Hypersensitivity to rilpivirine or to any of the excipients.

EDURANT® should not be co-administered with the following medicinal products, as significant decreases in rilpivirine plasma concentrations may occur (due to CYP3A enzyme induction or gastric pH increase), which may result in loss of therapeutic effect of EDURANT® (see *Interactions*):
- the anticonvulsants carbamazepine, oxcarbazepine, phenobarbital, phenytoin
- the antimycobacterials rifampicin, rifapentine
- proton pump inhibitors, such as omeprazole, esomeprazole, lansoprazole, pantoprazole, rabeprazole
- the glucocorticoid systemic dexamethasone, except as a single dose treatment
- St John's wort (*Hypericum perforatum*)

Warnings and Precautions
Patients should be advised that current antiretroviral therapy does not cure HIV and has not been proven to prevent the transmission of HIV to others through blood or sexual contact. Appropriate precautions to prevent the transmission of HIV should continue to be employed.

Virologic Failure and Development of Resistance
EDURANT® has not been evaluated in patients with previous virologic failure to any other antiretroviral therapy. The list of rilpivirine resistance-associated mutations presented in section *Pharmacodynamic Properties* should only guide the use of EDURANT® in the

treatment-naïve population.

In the pooled analysis from the phase III trials in adults through 96 weeks, patients treated with EDURANT® with a baseline viral load > 100000 HIV-1 RNA copies/mL had a greater risk of virologic failure (18.2% with EDURANT® versus 7.9% with efavirenz) compared to patients with a baseline viral load ≤ 100000 HIV-1 RNA copies/mL. The greater risk of virologic failure for patients in the EDURANT® arm was observed in the first 48 weeks of these trials while low rates of virologic failure, similar between the treatment arms, were observed from week 48 to week 96 (see *Pharmacodynamic Properties*). Patients with a baseline viral load > 100000 HIV-1 RNA copies/mL who experienced virologic failure exhibited a higher rate of treatment-emergent resistance to the NNRTI class. More patients who failed virologically on EDURANT® than who failed virologically on efavirenz developed lamivudine/emtricitabine associated resistance (see *Pharmacodynamic Properties*).

No new information was identified in pediatric patients 12 to less than 18 years of age in trial C213.

This information should be taken into consideration when initiating therapy with EDURANT®.

Cardiovascular

At supra-therapeutic doses (75 and 300 mg once daily), rilpivirine has been associated with prolongation of the QTc interval of the electrocardiogram (ECG).

EDURANT® at the recommended dose of 25 mg once daily is not associated with a clinically relevant effect on QTc. EDURANT® should be used with caution when co-administered with medicinal products with a known risk of Torsade de Pointes.

Interactions with Medicinal Products

Caution should be given to prescribing EDURANT® with medicinal products that may reduce the exposure of rilpivirine.

For information on interactions with medicinal products, see *Contraindications* and *Interaction with Other Medicinal Products and Other Forms of Interaction*.

Depressive Disorders

The adverse reaction depressive disorders (depressed mood, depression, dysphoria, major depression, mood altered, negative thoughts, suicide attempt, suicidal ideation) has been reported with EDURANT®. During the Phase 3 trials (N = 1368) through 96 weeks, the incidence of depressive disorders (regardless of causality, severity) reported among EDURANT® (n = 686) or efavirenz (n = 682) was 9% and 8%, respectively. Most events were mild or moderate in severity. The incidence of Grade 3 and 4 depressive disorders (regardless of causality) was 1% for both EDURANT® and efavirenz. The incidence of discontinuation due to depressive disorders among EDURANT® or efavirenz was 1% in each arm. Suicide ideation was reported in 4 subjects in each arm while suicide attempt was reported in 2 subjects in the EDURANT® arm. Patients with severe depressive symptoms should seek immediate medical evaluation to assess the possibility that the symptoms are related to EDURANT®, and if so, to determine whether the risks of continued therapy outweigh the benefits.

Fat Redistribution

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

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

Immune Reconstitution Inflammatory Syndrome

Immune reconstitution inflammatory syndrome has been reported in patients treated with combination antiretroviral therapy, including EDURANT®. During the initial phase of combination antiretroviral treatment, patients whose immune system responds may develop an inflammatory response to indolent or residual opportunistic infections (such as *Mycobacterium avium* complex, cytomegalovirus, *Pneumocystis jiroveci* pneumonia, and tuberculosis), which may necessitate further evaluation and treatment. Autoimmune disorders such as Graves' disease and autoimmune hepatitis have also been reported to occur in the setting of immune reconstitution inflammatory syndrome; however, the time to onset is more variable, and these events can occur many months after initiation of treatment (see *Adverse Reactions*).

Interactions

Medicinal Products that Affect Rilpivirine Exposure

Rilpivirine is primarily metabolised by cytochrome P450 (CYP)3A, and medicinal products that induce or inhibit CYP3A may thus affect the clearance of rilpivirine (see *Pharmacokinetic Properties*). Co-administration of EDURANT® and medicinal products that induce CYP3A may result in decreased plasma concentrations of rilpivirine which could potentially reduce the therapeutic effect of EDURANT®. Co-administration of EDURANT® and medicinal products that inhibit CYP3A may result in increased plasma concentrations of rilpivirine.

Co-administration of EDURANT® with medicinal products that increase gastric pH may result in decreased plasma concentrations of rilpivirine which could potentially reduce the therapeutic effect of EDURANT®.

Medicinal Products that are Affected by the Use of Rilpivirine

EDURANT® at a dose of 25 mg q.d. is not likely to have a clinically relevant effect on the exposure of medicinal products metabolised by CYP enzymes.

Established and theoretical interactions with selected antiretrovirals and non-antiretroviral medicinal products are listed in table 1 and table 2, respectively.

Interaction table

Interactions between rilpivirine and co-administered medicinal products are listed in the tables below (increase is indicated as "↑", decrease as "↓", no change as "↔", not applicable as "NA", once daily as "q.d." and twice daily as "b.i.d.").

Table 1: Drug interactions – Rilpivirine co-administered with antiretroviral and antiviral medicinal products

Co-administered medicinal product	Dose of co-administered medicinal product	Medicinal product assessed	C_{max}	AUC	C_{min}
HIV NUCLEOSIDE OR NUCLEOTIDE REVERSE TRANSCRIPTASE INHIBITORS (NRTIs/N[t]RTIs)					
Didanosine*#	400 mg q.d.	didanosine	↔	↑ 12%	NA
		rilpivirine	↔	↔	↔

	No dose adjustment is required when EDURANT® is co-administered with didanosine. Didanosine should be administered on an empty stomach and at least two hours before or at least four hours after EDURANT® (which should be administered with a meal).				
Tenofovir disoproxil fumarate*#	300 mg q.d.	tenofovir	↑ 19%	↑ 23%	↑ 24%
		rilpivirine	↔	↔	↔
	No dose adjustment is required when EDURANT® is co-administered with tenofovir disoproxil fumarate.				
Other NRTIs (abacavir, emtricitabine, lamivudine, stavudine and zidovudine)	Based on the different elimination routes for rilpivirine and these other NRTIs, no clinically relevant drug-drug interactions are expected between these medicinal products and EDURANT®.				
HIV NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS (NNRTIs)					
NNRTIs (delavirdine, efavirenz, etravirine, nevirapine)	It is not recommended to co-administer EDURANT® with NNRTIs.				
HIV PROTEASE INHIBITORS (PIs) - with co-administration of low dose ritonavir					
Darunavir/ritonavir*#	800/100 mg q.d.	darunavir	↔	↔	↓ 11%
		rilpivirine	↑ 79%	↑ 130%	↑ 178%
	Concomitant use of EDURANT® with darunavir/ritonavir may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). No dose adjustment is required when EDURANT® is co-administered with darunavir/ritonavir.				
Lopinavir/ritonavir (soft gel capsules)*#	400/100 mg b.i.d.	lopinavir	↔	↔	↓ 11%
		rilpivirine	↑ 29%	↑ 52%	↑ 74%
	Concomitant use of EDURANT® with lopinavir/ritonavir may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). No dose adjustment is required when EDURANT® is co-administered with lopinavir/ritonavir.				
Other boosted PIs (atazanavir/ritonavir, fosamprenavir/ritonavir, saquinavir/ritonavir, tipranavir/ritonavir)	Concomitant use of EDURANT® with boosted PIs may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). EDURANT® is not expected to affect the plasma concentrations of co-administered PIs.				
HIV PROTEASE INHIBITORS (PIs) - without co-administration of low dose ritonavir					
Unboosted PIs (atazanavir, fosamprenavir, indinavir, nelfinavir)	Concomitant use of EDURANT® with unboosted PIs may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). EDURANT® is not expected to affect the plasma concentrations of co-administered PIs.				
CCR5 ANTAGONISTS					
Maraviroc	No clinically relevant drug-drug interaction is expected when EDURANT® is co-administered with maraviroc.				
HIV INTEGRASE STRAND TRANSFER INHIBITORS					
Raltegravir*	400 mg b.i.d.	raltegravir	↑ 10%	↑ 9%	↑ 27%
		rilpivirine	↔	↔	↔
	No dose adjustment is required when EDURANT® is co-administered with raltegravir.				
OTHER ANTIVIRAL AGENTS					
Ribavirin	No clinically relevant drug-drug interaction is expected when EDURANT® is co-administered with ribavirin.				
Simeprevir*	150 mg once daily	simeprevir	↑ 10%	↔	↔
		rilpivirine	↔	↔	↑ 25%
	No dose adjustment is required for either drug when EDURANT® is co-administered with simeprevir.				

* The interaction between EDURANT® and the drug was evaluated in a clinical study. All other drug interactions shown are predicted.

\# This interaction study has been performed with a dose higher than the recommended dose for EDURANT® assessing the maximal effect on the co-administered drug. The dosing recommendation is applicable to the recommended dose of EDURANT® 25 mg q.d.

Table 2: Drug interactions – Rilpivirine co-administered with non-antiretroviral medicinal products

Co-administered medicinal product	Dose of co-administered medicinal product	Medicinal product assessed	C_{max}	AUC	C_{min}
ANTIARRHYTHMICS					
Digoxin*	0.5 mg single dose	digoxin	↔	↔	NA
	No dose adjustment is required when EDURANT® is co-administered with digoxin.				
ANTIDIABETICS					
Metformin*	850 mg single dose	metformin	↔	↔	NA
	No dose adjustment is required when EDURANT® is co-administered with metformin.				
ANTICONVULSANTS					
Carbamazepine Oxcarbazepine Phenobarbital Phenytoin	EDURANT® should not be used in combination with these anticonvulsants as co-administration may cause significant decreases in rilpivirine plasma concentrations (induction of CYP3A enzymes). This may result in loss of therapeutic effect of EDURANT®.				
AZOLE ANTIFUNGAL AGENTS					
Ketoconazole*#	400 mg q.d.	ketoconazole	↔	↓ 24%	↓ 66%
		rilpivirine	↑ 30%	↑ 49%	↑ 76%
Fluconazole Itraconazole Posaconazole Voriconazole	Concomitant use of EDURANT® with azole antifungal agents may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). No rilpivirine dose adjustment is required when EDURANT® is co-administered with azole antifungal agents. Clinically monitor for breakthrough fungal injections when azole antifungals are co-administered with EDURANT®.				
ANTIMYCOBACTERIALS					
Rifabutin*	300 mg q.d. †	rifabutin	↔	↔	↔
		25-*O*-desacetyl-rifabutin	↔	↔	↔
	300 mg q.d.	rilpivirine (25 mg q.d.)	↓ 31%	↓ 42%	↓ 48%
	300 mg q.d.	rilpivirine (50 mg q.d.)	↑ 43%	↑ 16%	↔
			(as compared to 25 mg q.d. rilpivirine alone)		
	Concomitant use of EDURANT® with rifabutin may cause significant decreases in rilpivirine plasma concentrations (induction of CYP3A enzymes). This may result in loss of therapeutic effect of EDURANT®. Throughout co-administration of EDURANT® with rifabutin, the EDURANT® dose should be increased from 25 mg once daily to 50 mg once daily. When rifabutin co-administration is stopped, the EDURANT® dose should be decreased to 25 mg once daily.				
Rifampicin*#	600 mg q.d.	rifampicin	↔	↔	NA
		25-desacetyl-rifampicin	↔	↓ 9%	NA
		rilpivirine	↓ 69%	↓ 80%	↓ 89%
Rifapentine	EDURANT® should not be used in combination with rifampicin or rifapentine as co-administration may cause significant decreases in rilpivirine plasma concentrations (induction of CYP3A enzymes). This may result in loss of therapeutic effect of EDURANT®.				
ANTICOAGULANTS					
Dabigatran	A risk for increases in dabigatran plasma concentrations cannot be excluded (inhibition of intestinal P-gp). The combination of EDURANT® and dabigatran etexilate should be used with caution.				

Table 2: Drug interactions – Rilpivirine co-administered with non-antiretroviral medicinal products					
MACROLIDE ANTIBIOTICS					
Clarithromycin Erythromycin	Concomitant use of EDURANT® with clarithromycin or erythromycin may cause an increase in the plasma concentrations of rilpivirine (inhibition of CYP3A enzymes). Where possible, alternatives such as azithromycin should be considered.				
GLUCOCORTICOIDS					
Dexamethasone (systemic)	EDURANT® should not be used in combination with systemic dexamethasone as co-administration may cause significant decreases in rilpivirine plasma concentrations (induction of CYP3A enzymes). This may result in loss of therapeutic effect of EDURANT®. Alternatives should be considered, particularly for long-term use.				
PROTON PUMP INHIBITORS					
Omeprazole*#	20 mg q.d.	omeprazole	↓ 14%	↓ 14%	NA
		rilpivirine	↓ 40%	↓ 40%	↓ 33%
Lansoprazole Rabeprazole Pantoprazole Esomeprazole	EDURANT® should not be used in combination with proton pump inhibitors as co-administration may cause significant decreases in rilpivirine plasma concentrations (gastric pH increase). This may result in loss of therapeutic effect of EDURANT®.				
H_2-RECEPTOR ANTAGONISTS					
Famotidine*#	40 mg single dose taken 12 hours before rilpivirine	rilpivirine	↔	↓ 9%	NA
	40 mg single dose taken 2 hours before rilpivirine	rilpivirine	↓ 85%	↓ 76%	NA
	40 mg single dose taken 4 hours after rilpivirine	rilpivirine	↑ 21%	↑ 13%	NA
Cimetidine Nizatidine Ranitidine	The combination of EDURANT® and H_2-receptor antagonists should be used with caution as co-administration may cause significant decreases in rilpivirine plasma concentrations (gastric pH increase). H_2-receptor antagonists should only be administered at least 12 hours before or at least 4 hours after EDURANT®.				
ANTACIDS					
Antacids (e.g., aluminium or magnesium hydroxide, calcium carbonate)	The combination of EDURANT® and antacids should be used with caution as co-administration may cause significant decreases in rilpivirine plasma concentrations (gastric pH increase). Antacids should only be administered either at least 2 hours before or at least 4 hours after EDURANT®.				
NARCOTIC ANALGESICS					
Methadone*	60-100 mg q.d., individualised dose	R(-) methadone	↓ 14%	↓ 16%	↓ 22%
		S(+) methadone	↓ 13%	↓ 16%	↓ 21%
	No dose adjustments are required when initiating co-administration of methadone with EDURANT®. However, clinical monitoring is recommended as methadone maintenance therapy may need to be adjusted in some patients.				
HERBAL PRODUCTS					
St John's wort (*Hypericum perforatum*)	EDURANT® should not be used in combination with products containing St John's wort as co-administration may cause significant decreases in rilpivirine plasma concentrations (induction of CYP3A enzymes). This may result in loss of therapeutic effect of EDURANT®.				
ANALGESICS					
Acetaminophen*# (paracetamol)	500 mg single dose	acetaminophen	↔	↔	NA
		rilpivirine	↔	↔	↑ 26%
	No dose adjustment is required when EDURANT® is co-administered with acetaminophen (paracetamol).				
ESTROGEN-BASED CONTRACEPTIVES					
Ethinylestradiol*	0.035 mg q.d.	ethinylestradiol	↑ 17%	↔	↔
Norethindrone*	1 mg q.d.	norethindrone	↔	↔	↔

Table 2: Drug interactions – Rilpivirine co-administered with non-antiretroviral medicinal products					
	No dose adjustment is required for the concomitant use of EDURANT® and estrogen- and/or progesterone-based contraceptives.				
HMG CO-A REDUCTASE INHIBITORS					
Atorvastatin*#	40 mg q.d.	atorvastatin	↑ 35%	↔	↓ 15%
		rilpivirine	↓ 9%	↔	↔
Fluvastatin Lovastatin Pitavastatin Pravastatin Rosuvastatin Simvastatin	No dose adjustment is required when EDURANT® is co-administered with an HMG Co-A reductase inhibitor.				
PHOSPHODIESTERASE TYPE 5 (PDE-5) INHIBITOR					
Sildenafil*#	50 mg single dose	sildenafil	↔	↔	NA
		rilpivirine	↔	↔	↔
Vardenafil Tadalafil	No dose adjustment is required when EDURANT® is co-administered with a PDE-5 inhibitor.				

* The interaction between EDURANT® and the drug was evaluated in a clinical study. All other drug-drug interactions shown are predicted.

\# This interaction study has been performed with a dose higher than the recommended dose for EDURANT® assessing the maximal effect on the co-administered drug. The dosing recommendation is applicable to the recommended dose of EDURANT® 25 mg q.d.

† This interaction study has been performed with a dose higher than the recommended dose for EDURANT®.

QT prolonging drugs

There is limited information available on the potential for a pharmacodynamic interaction between rilpivirine and medicinal products that prolong the QTc interval of the electrocardiogram. In a study of healthy subjects, supratherapeutic doses of rilpivirine (75 mg q.d. and 300 mg q.d.) have been shown to prolong the QTc interval of the electrocardiogram (see *Pharmacodynamic Properties*). EDURANT® should be used with caution when co-administered with a medicinal product with a known risk of Torsade de Pointes.

Pregnancy, Breast-feeding and Fertility

Women of Childbearing Potential

Since there are no well controlled clinical studies with EDURANT® in pregnant women, adequate contraception is recommended for women of childbearing potential when taking EDURANT®.

Contraception in Males and Females

A trial to investigate the effect of EDURANT® when co-administered with oral contraceptives demonstrated that EDURANT® is unlikely to decrease the effectiveness of oral contraceptives. EDURANT® and estrogen- and/or progesterone-based contraceptives can be used together without dose adjustments (see *Interactions*).

Pregnancy

There are no well controlled clinical studies with EDURANT® in pregnant women. Studies in animals have shown no evidence of relevant embryonic or foetal toxicity or an effect on reproductive function (see *Non-Clinical Information*). There was no teratogenicity with rilpivirine in rats and rabbits. The exposures at the embryo-fetal No Observed Adverse Effects Levels (NOAELs) in rats and rabbits were respectively 15 and 70 times higher than the exposure in humans at the recommended dose of 25 mg q.d. (see *Non-Clinical Information*).

To monitor maternal-fetal outcomes of pregnant women, an Antiretroviral Pregnancy Registry has been established (http://www.apregistry.com). This is a voluntary prospective, exposure-registration, observational study designed to collect and evaluate data on the outcomes of pregnancy exposures to antiretroviral products. For rilpivirine, sufficient first trimester exposures are available to allow detection of at least a two-fold increase in risk of overall birth defects. No such increases have been detected to date.

Rilpivirine in combination with a background regimen was evaluated in a clinical trial of 19 pregnant women during the second and third trimesters, and postpartum. The pharmacokinetic data demonstrate that total exposure (AUC) to rilpivirine as a part of an antiretroviral regimen was approximately 30% lower during pregnancy compared with postpartum (6-12 weeks). Virologic response was preserved throughout the trial period. No mother to child transmission occurred in all 10 infants born to the mothers who completed the trial and for whom the HIV status was available. Rilpivirine was well tolerated during pregnancy and postpartum. There were no new safety findings compared with the known safety profile of rilpivirine in HIV 1 infected adults (see *Pharmacokinetic Properties Special Populations – Pregnancy and Postpartum*).

EDURANT® should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Breast-feeding
It is not known whether rilpivirine is secreted in human milk. EDURANT® is excreted in the milk of rats. Because of both the potential for HIV transmission and the potential for adverse events in nursing infants, mothers should be instructed not to breastfeed if they are receiving EDURANT®.

Fertility
No human data on the effect of rilpivirine on fertility are available. In a study conducted in rats, there were no effects on mating or fertility with rilpivirine up to 400 mg/kg/day, a dose of rilpivirine that showed maternal toxicity (see *Non-Clinical Information*). This dose is associated with an exposure that is approximately 40 times higher than the exposure in humans at the recommended dose of 25 mg q.d.

Effects on Ability to Drive and Use Machines
EDURANT® has no or negligible influence on the ability to drive and use machines. No studies on the effects of EDURANT® on the ability to drive and use machines have been performed. Fatigue, dizziness and somnolence have been reported in some patients taking EDURANT® and should be considered when assessing a patient's ability to drive or operate machinery.

Adverse Reactions
Throughout this section, adverse reactions are reported. Adverse reactions are adverse events that were considered to be reasonably associated with the use of EDURANT® based on the comprehensive assessment of the available adverse event information. A causal relationship with EDURANT® usually cannot be reliably established in individual cases. Further, because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

Adverse reactions from clinical trials in adult patients

The safety assessment is based on the week 96 pooled data from 1368 patients in the phase III controlled trials TMC278-C209 (ECHO) and TMC278-C215 (THRIVE) in antiretroviral treatment-naïve HIV-1 infected adult patients, 686 of whom received EDURANT® (25 mg q.d.) (see *Pharmacodynamic Properties*). The median duration of exposure for patients in the EDURANT® and efavirenz arms was 104.3 and 104.1 weeks, respectively. Most ADRs occurred in the first 48 weeks of treatment.

In the phase III controlled trials ECHO and THRIVE through 96 weeks, the most frequently reported adverse drug reactions (ADRs) (> 2%) to EDURANT® that were at least grade 2 in severity were depression, headache, insomnia, transaminases increased and rash (see table 3 for the complete list of ADRs).

The majority of the ADRs reported during treatment with EDURANT® 25 mg once daily were grade 1 to 2 in severity. Grade 3 or 4 ADRs were reported in 3.6% and 5.9% of the EDURANT® and efavirenz treated patients, respectively. The most common (reported in more than 1 patient in the EDURANT® arm) grade 3 or 4 ADRs were transaminases increased (1.6% in the EDURANT® arm and 2.9% in the efavirenz arm), depression (0.7% and 0.7%, respectively), abdominal pain (0.4% and 0.1%, respectively), dizziness (0.3% and 0.4%, respectively) and rash (0.3% and 0.6%, respectively). 1.7% of patients in the EDURANT® arm discontinued treatment due to ADRs compared to 4.0% of patients in the efavirenz arm. In the EDURANT® arm, all ADRs leading to discontinuation had an incidence < 0.5%. In the efavirenz arm, the most common ADRs leading to discontinuation were rash (1.5%), transaminases increased (0.7%), depression (0.6%) and abnormal dreams (0.6%).

ADRs of at least moderate intensity (≥ grade 2) reported in adult patients treated with EDURANT® are summarised in table 3. The ADRs are listed by system organ class (SOC) and frequency. Selected treatment emergent laboratory abnormalities, considered as ADRs, are included in table 4.

Table 3: ADRs of at least moderate intensity (≥ grade 2) reported in antiretroviral treatment-naïve HIV-1 infected adult patients treated with EDURANT®

	Pooled data from the week 96 analysis of the phase III ECHO and THRIVE trials		
System Organ Class (SOC) Adverse drug reaction, %	**EDURANT® + BR N=686**	**Efavirenz + BR N=682**	**Treatment Difference (95%CI)**
Metabolism and nutrition disorders			
Decreased appetite	1.2%	0.6%	0.6 (-0.4; 1.6)
Psychiatric disorders			
Depression	4.1%	3.2%	0.9 (-1.1; 2.8)
Insomnia	3.5%	3.5%	0 (-2.0; 1.9)
Abnormal dreams*†	1.6%	4.0%	-2.4 (-4.1; -0.6)
Sleep disorders	1.3%	0.9%	0.4 (-0.7; 1.5)
Depressed mood	0.4%	0.3%	0.1 (-0.5; 0.8)
Nervous system disorders			
Headache*	3.5%	3.8%	-0.3 (-2.3; 1.7)
Dizziness*#	1.0%	6.7%	-5.7 (-7.7; -3.7)
Somnolence	0.7%	1.3%	-0.6 (-1.7; 0.5)
Gastrointestinal disorders			
Abdominal pain	2.0%	1.9%	0.1 (-1.3; 1.6)
Nausea*	1.3%	2.8%	-1.5 (-3.0; 0)
Vomiting	1.0%	2.1%	-1.0 (-2.3; 0.3)
Abdominal discomfort	0.4%	0.1%	0.3 (-0.3; 0.9)
Skin and subcutaneous tissue disorders			
Rash*#	2.3%	9.5%	-7.2 (-9.7; -4.7)
General disorders and administration site conditions			
Fatigue	1.6%	2.1%	-0.4 (-1.9; 1.0)

Table 3: ADRs of at least moderate intensity (≥ grade 2) reported in antiretroviral treatment-naïve HIV-1 infected adult patients treated with EDURANT®			
Investigations			
Transaminases increased	2.8%	4.0%	-1.2 (-3.1; 0.7)

BR=background regimen; CI=confidence interval
N=total number of subjects per treatment group
* Treatment comparison was pre-specified for these ADRs (Fisher's Exact Test)
† p-value < 0.01
# p-value < 0.0001

No new ADR terms were identified in adult patients in the phase III ECHO and THRIVE trials between 48 weeks and 96 weeks nor in the phase IIb TMC278-C204 trial through 240 weeks.

Laboratory abnormalities

Selected treatment emergent clinical laboratory abnormalities (grade 3 or grade 4), considered as ADRs, reported in EDURANT®-treated patients are shown in table 4.

Table 4: Selected treatment emergent laboratory abnormalities (grade 3 or grade 4) observed in antiretroviral treatment-naïve HIV-1 infected adult patients			
Laboratory parameter abnormality, %	**DAIDS toxicity range**	**Pooled data from the week 96 analysis of the phase III ECHO and THRIVE trials**	
		EDURANT® + BR N=686	**Efavirenz + BR N=682**
HEMATOLOGY			
Decreased hemoglobin	< 4.5 mmol/L < 7.4 g/dL	0.1%	0.6%
Decreased platelet count	< 49999/mm^3 < 49999 x 10^9/L	0.1%	0.3%
Decreased white blood cell count	< 1499/mm^3 < 1.499 giga/L	1.2%	1.0%
BIOCHEMISTRY			
Increased creatinine	> 1.8 x ULN	0.1%	0.1%
Increased AST	> 5.0 x ULN	2.3%	3.3%
Increased ALT	> 5.0 x ULN	1.6%	3.7%
Increased bilirubin	> 2.5 x ULN	0.7%	0.3%
Increased pancreatic amylase	> 2 x ULN	3.8%	4.8%
Increased lipase	> 3 x ULN	0.9%	1.6%
Increased total cholesterol (fasted)*	> 7.77 mmol/L > 300 mg/dL	0.1%	3.3%
Increased LDL cholesterol (fasted)*	≥ 4.91 mmol/L ≥ 191 mg/dL	1.5%	5.3%
Increased triglycerides (fasted)*	≥ 8.49 mmol/L ≥ 751 mg/dL	0.6%	3.3%

BR=background regimen; ULN=upper limit of normal
N=number of subjects per treatment group
* $p \leq 0.001$ according to Fisher's Exact test (difference in grade 3 plus 4 abnormalities between the two treatment groups).
Note: Percentages were calculated for the number of subjects with results for the analyte.

Adrenal Function

In the pooled phase III trials, at week 96, there was an overall mean change from baseline in basal cortisol of -19.1 (-30.85; -7.37) nmol/L in the EDURANT® group and of -0.6 (-13.29; 12.17) nmol/L in the efavirenz group. At week 96, the mean change from baseline in ACTH-stimulated cortisol levels was lower in the EDURANT® group (+18.4 ±

8.36 nmol/L) than in the efavirenz group (+54.1 ± 7.24 nmol/L). Mean values for both basal and ACTH-stimulated cortisol values at week 96 were within the normal range. Overall, there were no serious adverse events, deaths, or treatment discontinuations that could clearly be attributed to adrenal insufficiency.

Serum Creatinine

In the pooled Phase III trials, an increase in serum creatinine was observed over the 96 weeks of treatment with EDURANT®. Most of this increase occurred within the first four weeks of treatment, with a mean change of 0.1 mg/dL (range: -0.3 mg/dL to 0.6 mg/dL) observed after 96 weeks of treatment. In subjects who entered the trial with mild or moderate renal impairment, the serum creatinine increase observed was similar to that seen in subjects with normal renal function. These changes are not considered to be clinically relevant and no subject discontinued treatment due to increases in serum creatinine. Serum creatinine increases occurred regardless of the background N(t)RTI regimen.

Serum Lipids

Changes from baseline in total cholesterol, LDL-cholesterol, HDL-cholesterol and triglycerides are presented in table 5. The mean changes from baseline were smaller in the EDURANT® arm versus the efavirenz arm. The impact of such findings has not been demonstrated.

Table 5: Lipid values, mean change from baseline

		Pooled data from the week 96 analysis of the phase III ECHO and THRIVE Trials				
		EDURANT® + BR N=686		**Efavirenz + BR N=682**		
		Baseline	**Week 96**	**Baseline**	**Week 96**	
Mean (95% CI)	**Mean (mg/dL)**	**Mean (mg/dL)**	**Mean change* (mg/dL)**	**Mean (mg/dL)**	**Mean (mg/dL)**	**Mean change* (mg/dL)**
Total cholesterol (fasted)†	161	167	5	161	190	28
HDL-cholesterol (fasted)†	41	46	4	40	51	11
LDL-cholesterol (fasted)†	96	98	1	96	110	14
Triglycerides (fasted)†	124	117	-7	133	148	12

N=number of subjects per treatment group

* The change from baseline is the mean of within-patient changes from baseline for patients with both baseline and week 96 values.

† p-value < 0.001, Wilcoxon rank-sum test for treatment comparison of change from baseline

Adverse drug reactions from a clinical trial in pediatric patients (12 to less than 18 years of age)

The safety assessment is based on the Week 48 analysis of the single-arm, open-label Phase 2 trial, TMC278-C213, in which 36 antiretroviral treatment-naïve HIV-1 infected patients 12 to less than 18 years of age and weighing at least 32 kg received EDURANT® (25 mg once daily) in combination with other antiretroviral medicinal products (see *Pharmacological properties – Clinical Studies*). The median duration of exposure for patients was 63.5 weeks. There were no patients who discontinued treatment due to ADRs. No new ADRs were identified compared to those seen in adults.

ADRs were reported in nineteen pediatric subjects (52.8%). Most ADRs were Grade 1 or 2. The most common ADRs reported in at least 2 subjects (regardless of severity) include

headache (19.4%), depression (19.4%), somnolence (13.9%), nausea (11.1%), dizziness (8.3%), abdominal pain (8.3%), vomiting (5.6%) and rash (5.6%). Observed laboratory abnormalities were comparable to those in adults.

Adrenal Function

In trial TMC278 C213, at Week 48, the overall mean change from baseline in basal cortisol showed an increase of 1.59 (0.24, 2.93) mcg/dL.

Six of 30 (20%) subjects with a normal 250 mcg ACTH stimulation test at baseline developed an abnormal 250 mcg ACTH stimulation test (peak cortisol level < 18.1 mcg/dL) during the trial. Three of these subjects had an abnormal 250 mcg ACTH stimulation test at Week 48. Overall there were no serious adverse events, deaths, or treatment discontinuations that could clearly be attributed to adrenal insufficiency. The clinical significance of the abnormal 250 mcg ACTH stimulation tests is not known.

Lipodystrophy

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

Immune reconstitution inflammatory syndrome

In HIV infected patients with severe immune deficiency at the time of initiation of combination antiretroviral therapy (CART), an inflammatory reaction to asymptomatic or residual opportunistic infections may arise (immune reconstitution inflammatory syndrome). Autoimmune disorders such as Graves' disease and autoimmune hepatitis have also been reported in the context of immune reconstitution inflammatory syndrome (see *Warnings and Precautions*).

Postmarketing Experience

Adverse reactions have been identified during post-marketing in patients receiving a rilpivirine containing regimen. Because these reactions are reported voluntarily from a population of unknown size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Renal and Genitourinary Disorders: nephrotic syndrome

Additional Information on Special Populations

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

In patients co-infected with hepatitis B or C virus receiving EDURANT®, the incidence of hepatic enzyme elevation was higher than in patients receiving EDURANT® who were not co-infected. This observation was the same in the efavirenz arm. The pharmacokinetic exposure of rilpivirine in co-infected patients was comparable to that in patients without co-infection.

Overdose

There is no specific antidote for overdose with EDURANT®. Human experience of overdose with EDURANT® is limited. Treatment of overdose with EDURANT® consists of general supportive measures including monitoring of vital signs and ECG (QT interval) as well as observation of the clinical status of the patient. It is advisable to contact a poison control center to obtain the latest recommendations for the management of an overdose. Since

rilpivirine is highly bound to plasma protein, dialysis is unlikely to result in significant removal of the active substance.

PHARMACOLOGICAL PROPERTIES

Pharmacodynamic Properties

Pharmacotherapeutic group: Antiviral for systemic use, NNRTI (non-nucleoside reverse transcriptase inhibitor), ATC code: J05AG05.

Mechanism of action

Rilpivirine is a diarylpyrimidine NNRTI of HIV-1. Rilpivirine activity is mediated by non-competitive inhibition of HIV-1 reverse transcriptase (RT). Rilpivirine does not inhibit the human cellular DNA polymerases α, β and γ.

Pharmacodynamic effects

Antiviral activity *in vitro*

Rilpivirine exhibited activity against laboratory strains of wild-type HIV-1 in an acutely infected T-cell line with a median EC_{50} value for HIV-1/IIIB of 0.73 nM (0.27 ng/mL). Although rilpivirine demonstrated limited *in vitro* activity against HIV-2 with EC_{50} values ranging from 2510 to 10830 nM (920 to 3970 ng/mL), treatment of HIV-2 infection with EDURANT® is not recommended in the absence of clinical data.
Rilpivirine also demonstrated antiviral activity against a broad panel of HIV-1 group M (subtype A, B, C, D, F, G, H) primary isolates with EC_{50} values ranging from 0.07 to 1.01 nM (0.03 to 0.37 ng/mL) and group O primary isolates with EC_{50} values ranging from 2.88 to 8.45 nM (1.06 to 3.10 ng/mL).

The antiviral activity of rilpivirine was not antagonistic when combined with the N(t)RTIs abacavir, didanosine, emtricitabine, lamivudine, stavudine, zidovudine and tenofovir; the PIs amprenavir, atazanavir, darunavir, indinavir, lopinavir, nelfinavir, ritonavir, saquinavir and tipranavir; the NNRTIs efavirenz, etravirine and nevirapine; the fusion inhibitor enfuvirtide; the CCR5 co-receptor antagonist maraviroc; and the integrase strand transfer inhibitor raltegravir.

Resistance

In cell culture

Rilpivirine-resistant strains were selected in cell culture starting from wild-type HIV-1 of different origins and subtypes as well as NNRTI resistant HIV-1. The most commonly observed amino acid substitutions that emerged included: L100I, K101E, V108I, E138K, V179F, Y181C, H221Y, F227C and M230I.

Resistance to rilpivirine was determined as a fold change in EC_{50} value (FC) above the biological cut-off (BCO) of the assay.

In treatment-naïve adult subjects

For the resistance analysis, a broader definition of virologic failure was used than in the primary efficacy analysis. In the week 48 pooled resistance analysis from the phase III trials, 62 (of a total of 72) virologic failures in the EDURANT® arm had resistance data at baseline and time of failure. In this analysis, the amino acid substitutions associated with NNRTI resistance that developed in at least 2 rilpivirine virologic failures were: V90I, K101E, E138K, E138Q, V179I, Y181C, V189I, H221Y, and F227C. In the trials, the presence of the substitutions V90I and V189I, at baseline, did not affect response. The E138K substitution emerged most frequently during rilpivirine treatment, commonly in combination with the M184I substitution. In the week 48 analysis, 31 out of 62 of rilpivirine virologic failures

had concomitant NNRTI and NRTI RAMs; 17 of those 31 had the combination of E138K and M184I. The most common mutations were the same in the week 48 and week 96 analyses.

More patients who failed virologically on EDURANT® than who failed virologically on efavirenz developed lamivudine/emtricitabine associated resistance.

In the week 96 pooled resistance analysis, low rates of virologic failure, similar between the treatment arms, were observed from week 48 to week 96 (3.2% in the EDURANT® arm and 2.3% in the efavirenz arm).

Considering all of the available *in vitro* and *in vivo* data, the following amino acid substitutions, when present at baseline, are likely to affect the activity of rilpivirine: K101E, K101P, E138A, E138G, E138K, E138R, E138Q, V179L, Y181C, Y181I, Y181V, Y188L, H221Y, F227C, M230I, and M230L. These rilpivirine resistance-associated mutations should only guide the use of EDURANT® in the treatment-naïve population. These resistance-associated mutations were derived from *in vivo* data involving treatment-naïve subjects only and therefore cannot be used to predict the activity of rilpivirine in subjects who have virologically failed an antiretroviral-containing regimen.

Cross-resistance

Site-directed NNRTI mutant virus

In a panel of 67 HIV-1 recombinant laboratory strains with one resistance-associated mutation at RT positions associated with NNRTI resistance, including the most commonly found K103N and Y181C, rilpivirine showed antiviral activity against 64 (96%) of these strains. The single resistance-associated mutations associated with a loss of susceptibility to rilpivirine were: K101P, Y181I and Y181V. The K103N substitution did not result in reduced susceptibility to rilpivirine by itself, but the combination of K103N and L100I resulted in a 7-fold reduced susceptibility to rilpivirine.

Recombinant clinical isolates

Rilpivirine retained sensitivity (FC ≤ BCO) against 62% of 4786 HIV-1 recombinant clinical isolates resistant to efavirenz and/or nevirapine.

Treatment-naïve HIV-1 infected adult patients

In the week 48 pooled analysis of the phase III trials ECHO and THRIVE, 31 of the 62 subjects with virologic failure on EDURANT with phenotypic resistance data lost susceptibility to rilpivirine. Of these, 28 were resistant to etravirine, 27 to efavirenz, and 14 to nevirapine. These cross-resistance findings were confirmed in the week 96 pooled analyses of the phase III clinical trials.

In the week 96 pooled analyses, among virologic failures in the EDURANT® arm with baseline viral load ≤ 100000 copies/mL and with resistance to rilpivirine, there were fewer patients with phenotypic cross-resistance than among those in the EDURANT® arm with baseline viral load > 100000 copies/mL. 3, 4 and 1 rilpivirine virologic failures with baseline viral load ≤ 100000 copies/mL and with resistance to rilpivirine (N = 5) had cross-resistance to efavirenz, etravirine and nevirapine, respectively, compared to 27, 28, and 15 rilpivirine virologic failures with baseline viral load > 100000 copies/mL (N = 30), respectively.

Effects on electrocardiogram

The effect of EDURANT® at the recommended dose of 25 mg q.d. on the QTcF interval was evaluated in a randomised, placebo and active (moxifloxacin 400 mg once daily) controlled

crossover study in 60 healthy adults, with 13 measurements over 24 hours at steady-state. EDURANT® at the recommended dose of 25 mg q.d. is not associated with a clinically relevant effect on QTc.

When supratherapeutic doses of 75 mg q.d. and 300 mg q.d. of EDURANT® were studied in healthy adults, the maximum mean time-matched (95% upper confidence bound) differences in QTcF interval from placebo after baseline correction were 10.7 (15.3) and 23.3 (28.4) ms, respectively. Steady-state administration of EDURANT® 75 mg q.d. and 300 mg q.d. resulted in a mean C_{max} approximately 2.6-fold and 6.7-fold, respectively, higher than the mean steady-state C_{max} observed with the recommended 25 mg q.d. dose of EDURANT®.

Clinical experience

Treatment-naïve HIV-1 infected adult patients

The evidence of efficacy of EDURANT® is based on the analyses of 96 week data from 2 randomised, double-blinded, active-controlled, phase III trials TMC278-C209 (ECHO) and TMC278-C215 (THRIVE). The trials were identical in design, with the exception of the background regimen (BR). At 96 weeks, the virologic response rate confirmed undetectable viral load (< 50 HIV-1 RNA copies/mL)] was evaluated in patients receiving EDURANT® 25 mg q.d. in addition to a BR versus patients receiving efavirenz 600 mg q.d. in addition to a BR. Similar efficacy for EDURANT® was seen in each trial demonstrating non-inferiority to efavirenz.

Antiretroviral treatment-naïve HIV-1 infected patients were enrolled who had a plasma HIV-1 RNA ≥ 5000 copies/mL and were screened for susceptibility to N(t)RTIs and for absence of specific NNRTI RAMs. In ECHO, the BR was fixed to the N(t)RTIs, tenofovir disoproxil fumarate plus emtricitabine. In THRIVE, the BR consisted of 2 investigator-selected N(t)RTIs: tenofovir disoproxil fumarate plus emtricitabine or zidovudine plus lamivudine or abacavir plus lamivudine. In ECHO, randomisation was stratified by screening viral load. In THRIVE, randomisation was stratified by screening viral load and by N(t)RTI BR.

This analysis included 690 patients in ECHO and 678 patients in THRIVE who had completed 96 weeks of treatment or discontinued earlier.

In the pooled analysis for ECHO and THRIVE, demographics and baseline characteristics were balanced between the EDURANT® arm and the efavirenz arm. Table 6 displays selected demographic and baseline disease characteristics of the patients in the EDURANT® and efavirenz arms.

Table 6: Demographic and baseline disease characteristics of antiretroviral treatment-naïve HIV-1 infected adult subjects in the ECHO and THRIVE trials (pooled analysis)

	Pooled data from the ECHO and THRIVE trials	
	EDURANT® + BR N=686	**Efavirenz + BR N=682**
Demographic characteristics		
Median Age, years (range)	36 (18-78)	36 (19-69)
Sex		
Male	76%	76%
Female	24%	24%
Race		
White	61%	60%
Black/African American	24%	23%
Asian	11%	14%
Other	2%	2%

Not allowed to ask per local regulations	1%	1%
Baseline disease characteristics		
Median baseline plasma HIV-1 RNA (range), $\log_{10}$ copies/mL	5.0 (2-7)	5.0 (3-7)
Median baseline CD4+ cell count (range), x 10^6 cells/L	249 (1-888)	260 (1-1137)
Percentage of subjects with: hepatitis B/C virus co-infection	7.3%	9.5%
Percentage of patients with the following background regimens: tenofovir disoproxil fumarate plus emtricitabine	80.2%	80.1%
zidovudine plus lamivudine	14.7%	15.1%
abacavir plus lamivudine	5.1%	4.8%

BR=background regimen

Table 7 below shows the efficacy results at 48 weeks and at 96 weeks for patients treated with EDURANT® and patients treated with efavirenz from the pooled data from the ECHO and THRIVE trials. The response rate (confirmed undetectable viral load < 50 HIV-1 RNA copies/mL) at week 96 was comparable between the EDURANT® arm and the efavirenz arm. The incidence of virologic failure was higher in the EDURANT® arm than the efavirenz arm at week 96; however, most of the virologic failures occurred within the first 48 weeks of treatment. Discontinuations due to adverse events were higher in the efavirenz arm at week 96 than the EDURANT® arm. Most of these discontinuations occurred in the first 48 weeks of treatment.

Table 7: Virologic Outcome of Randomised Treatment in the ECHO and THRIVE Trials in adults (Pooled Analysis at Week 48 (primary) and Week 96; ITT-TLOVR*)

	Outcome at Week 48		***Outcome at Week 96***	
%	**EDURANT® + BR N=686**	**Efavirenz + BR N=682**	**EDURANT® + BR N=686**	**Efavirenz + BR N=682**
Confirmed Undetectable Viral Load (< 50 HIV-1 RNA copies/mL) §#	84.3	82.3	77.6	77.6
Virologic Failure†	9.0	4.8	11.5	5.9
Death	0.1	0.4	0.1	0.9
Discontinued due to adverse event (AE)	2.0	6.7	3.8	7.6
Discontinued for non-AE reason¶	4.5	5.7	7.0	8.1

N = number of subjects per treatment group

* intent-to-treat time to loss of virologic response

§ Subjects achieved virologic response (two consecutive viral loads < 50 copies/mL) and maintained it through week 48/96.

Predicted difference of response rates (95% CI) at week 48: 1.6% (-2.2%; 5.3%) and at week 96: -0.4% (-4.6%; 3.8%); both p-values < 0.0001 (non-inferiority at 12% margin) from logistic regression model, including stratification factors and study.

† Includes subjects who were rebounder (confirmed viral load ≥ 50 copies/mL after being responder) or who were never suppressed (no confirmed viral load < 50 copies/mL, either ongoing or discontinued due to lack or loss of efficacy).

¶ e.g. lost to follow-up, non-compliance, withdrew consent

At week 96, the mean change from baseline in CD4+ cell count was +228 x 10^6 cells/L in the EDURANT® arm and +219 x 10^6 cells/L in the efavirenz arm in the pooled analysis of the ECHO and THRIVE trials [estimated treatment difference (95% CI): 11.3 (-6.8; 29.4)].

From the week 96 pooled resistance analysis, the resistance outcome for patients with protocol defined virological failure, and paired genotypes (baseline and failure) is shown in table 8.

Table 8: Resistance outcome by background NRTI regimen used (pooled data from ECHO and THRIVE trials in the week 96 resistance analysis)

	tenofovir/ emtricitabine	zidovudine/ lamivudine	abacavir/ lamivudine	All*
EDURANT®-treated				
Resistance[#] to emtricitabine/lamivudine % (n/N)	6.9 (38/550)	3.0 (3/101)	8.6 (3/35)	6.4 (44/686)
Resistance to rilpivirine % (n/N)	6.5 (36/550)	3.0 (3/101)	8.6 (3/35)	6.1 (42/686)
Efavirenz-treated				
Resistance to emtricitabine/lamivudine % (n/N)	1.1 (6/546)	1.9 (2/103)	3.0 (1/33)	1.3 (9/682)
Resistance to efavirenz % (n/N)	2.4 (13/546)	2.9 (3/103)	3.0 (1/33)	2.5 (17/682)

***** The number of patients with virologic failure and paired genotypes (baseline and failure) were 71, 11, and 4 for EDURANT® and 30, 10, and 2 for efavirenz, for the tenofovir/emtricitabine, zidovudine/lamivudine, and abacavir/lamivudine regimens, respectively.

[#] Resistance was defined as the emergence of any resistance-associated mutation at failure.

A subgroup analysis of the virologic response (< 50 HIV-1 RNA copies/mL) at 48 and 96 weeks and virologic failure by baseline viral load, CD4 count and by background NRTIs (pooled data from the ECHO and THRIVE trials) is presented in table 9.

Table 9: Virologic response (< 50 HIV-1 RNA copies/mL, ITT-TLOVR) and virologic failure by baseline viral load and by background NRTIs (Pooled analysis at Week 48 [primary] and Week 96 from the ECHO and THRIVE trials in adults)

	Outcome at Week 48				***Outcome at Week 96***			
	EDURANT® + BR N=686		**Efavirenz + BR N=682**		**EDURANT® + BR N=686**		**Efavirenz + BR N=682**	
	N	**n (%)**	**N**	**n (%)**	**N**	**n (%)**	**N**	**n (%)**
Proportion of patients with HIV-1 RNA < 50 copies/mL at week 48* and at week 96*by baseline plasma viral load (copies/mL)								
≤ 100000	368	332 (90.2%)	330	276 (83.6%)	368	309 (84.0%)	329	263 (79.9%)
> 100000	318	246 (77.4%)	352	285 (81.0%)	318	223 (70.1%)	353	266 (75.4%)
> 100000 to ≤ 500000	249	198 (79.5%)	270	223 (82.6%)	249	178 (71.5%)	270	205 (75.9%)
> 500000	69	48 (69.6%)	82	62 (75.6%)	69	45 (65.2%)	83	61 (73.5%)
Virologic Failure[†] by baseline plasma viral load (copies/mL)								
≤ 100000	368	14 (3.8%)	330	11 (3.3%)	368	21 (5.7%)	329	12 (3.6%)
> 100000	318	48 (15.1%)	352	22 (6.3%)	318	58 (18.2%)	353	28 (7.9%)
> 100000 to ≤ 500000	249	33 (13.3%)	270	13 (4.8%)	249	43 (17.3%)	270	18 (6.7%)
> 500000	69	15 (21.7%)	82	9 (11.0%)	69	15 (21.7%)	83	10 (12.0%)
Proportion of patients with HIV-1 RNA < 50 copies/mL at week 48* and at week 96* by baseline CD4 count (x 10^6 cells/L)								
< 50	34	20 (58.8%)	36	29 (80.6%)	34	19 (55.9%)	36	25 (69.4%)

Table 9: Virologic response (< 50 HIV-1 RNA copies/mL, ITT-TLOVR) and virologic failure by baseline viral load and by background NRTIs (Pooled analysis at Week 48 [primary] and Week 96 from the ECHO and THRIVE trials in adults)								
≥ 50-< 200	194	156 (80.4%)	175	143 (81.7%)	194	138 (71.1%)	175	131 (74.9%)
≥ 200-< 350	313	272 (86.9%)	307	253 (82.4%)	313	252 (80.5%)	307	244 (79.5%)
≥ 350	144	130 (90.3%)	164	136 (82.9%)	144	123 (85.4%)	164	129 (78.7%)
Virologic Failure† by baseline CD4 count (x 10^6 cells/L)								
< 50	34	6 (17.6%)	36	1 (2.8%)	34	6 (17.6%)	36	4 (11.1%)
≥ 50-< 200	194	27 (13.9%)	175	14 (8.0%)	194	37 (19.1%)	175	14 (8.0%)
≥ 200-< 350	313	21 (6.7%)	307	14 (4.6%)	313	26 (8.3%)	307	15 (4.9%)
≥ 350	144	8 (5.6%)	164	4 (2.4%)	144	10 (6.9%)	164	7 (4.3%)
Proportion of patients with HIV-1 RNA < 50 copies/mL at week 48* and at week 96* by background N(t)RTI								
tenofovir disoproxil fumarate plus emtricitabine	550	459 (83.5%)	546	450 (82.4%)	550	423 (76.9%)	546	422 (77.3%)
zidovudine plus lamivudine	101	88 (87.1%)	103	83 (80.6%)	101	82 (81.2%)	103	79 (76.7%)
abacavir plus lamivudine	35	31 (88.6%)	33	28 (84.8%)	35	27 (77.1%)	33	28 (84.8%)

N=number of subjects per treatment group
n=number of observations

* Imputations according to the TLOVR algorithm.

† Includes subjects who were rebounder (confirmed viral load ≥ 50 copies/mL after being responder) or who were never suppressed (no confirmed viral load < 50 copies/mL, either ongoing or discontinued due to lack or loss of efficacy).

Study TMC278-C204 was a randomised, active-controlled, phase IIb trial in antiretroviral treatment-naïve HIV-1 infected adult patients consisting of 2 parts: an initial partially blinded dose-finding part [EDURANT® doses blinded] up to 96 weeks, followed by a long-term, open label part. In the open label part of the trial, patients originally randomised to one of the 3 doses of EDURANT® were all treated with EDURANT® 25 mg once daily in addition to a BR, once the dose for the phase III studies was selected. Patients in the control arm received efavirenz 600 mg once daily in addition to a BR in both parts of the study. The BR consisted of 2 investigator-selected N(t)RTIs: zidovudine plus lamivudine or tenofovir disoproxil fumarate plus emtricitabine.

Study TMC278-C204 enrolled 368 HIV-1 infected treatment-naïve adult patients who had a plasma HIV-1 RNA ≥ 5000 copies/mL, previously received ≤ 2 weeks of treatment with an N(t)RTI or protease inhibitor, had no prior use of NNRTIs, and were screened for susceptibility to N(t)RTI and for absence of specific NNRTI RAMs.

At 96 weeks, the proportion of patients with < 50 HIV-1 RNA copies/mL receiving EDURANT® 25 mg (N=93) compared to patients receiving efavirenz (N=89) was 76% and 71%, respectively. The mean increase from baseline in CD4+ counts was 146 x 10^6 cells/L in patients receiving EDURANT® 25 mg and 160 x 10^6 cells/L in patients receiving efavirenz.

Of those patients who were responders at week 96, 74% of patients receiving EDURANT® remained with undetectable viral load (< 50 HIV-1 RNA copies/mL) at week 240 compared to 81% of patients receiving efavirenz. There were no safety concerns identified in the week 240 analyses.

Pregnancy
Rilpivirine in combination with a background regimen was evaluated in a clinical trial of 19 pregnant women during the second and third trimesters, and postpartum. The pharmacokinetic data demonstrate that total exposure (AUC) to rilpivirine as a part of an antiretroviral regimen was approximately 30% lower during pregnancy compared with postpartum (6-12 weeks). Virologic response was preserved throughout the trial period. No mother to child transmission occurred in all 10 infants born to the mothers who completed the trial and for whom the HIV status was available. Rilpivirine was well tolerated during pregnancy and postpartum. There were no new safety findings compared with the known safety profile of rilpivirine in HIV 1 infected adults.

Treatment-naïve HIV-1 infected pediatric patients (12 years to less than 18 years of age)
The pharmacokinetics, safety, tolerability and efficacy of EDURANT® 25 mg once daily, in combination with an investigator-selected BR containing two NRTIs, was evaluated in trial TMC278-C213, a single-arm, open-label Phase 2 trial in antiretroviral treatment-naive HIV-1 infected pediatric subjects 12 to less than 18 years of age and weighing at least 32 kg. This analysis included 36 patients who had completed at least 48 weeks of treatment or discontinued earlier.

The 36 subjects had a median age of 14.5 years (range: 12 to less than 18 years of age), and were 55.6% female, 88.9% Black and 11.1% Asian. The median baseline plasma HIV-1 RNA was 4.8 $\log_{10}$ copies/mL, and the median baseline CD4+ cell count was 414 x 10^6 cells/L (range: 25 to 983 x 10^6 cells/L).

The proportion of subjects with HIV-1 RNA < 50 copies/mL at Week 48 (TLOVR) was 72.2% (26/36). The proportion of responders was higher in subjects with a baseline viral load ≤100000 copies/mL (78.6%, 22/28) as compared to those with a baseline viral load >100000 copies/mL (50.0%, 4/8). The proportion of virological failures was 22.2% (8/36). The proportion of virologic failures was lower in subjects with a baseline viral load ≤100000 copies/mL (17.9%, 5/28) as compared to those with a baseline viral load >100000 copies/mL (37.5%, 3/8). One subject discontinued due to an adverse event and 1 subject discontinued due to reasons other than an adverse event or virology failure. At Week 48, the mean increase in CD4+ cell count from baseline was 201.2 x 10^6 cells/L.

Pharmacokinetic Properties
The pharmacokinetic properties of rilpivirine have been evaluated in adult healthy subjects and in antiretroviral treatment-naïve HIV-1 infected patients 12 years of age and older. Exposure to rilpivirine was generally lower in HIV-1 infected patients than in healthy subjects.

Absorption
After oral administration, the maximum plasma concentration of rilpivirine is generally achieved within 4-5 hours. The absolute bioavailability of EDURANT® is unknown.

Effect of food on absorption
The exposure to rilpivirine was approximately 40% lower when EDURANT® was taken in a fasted condition as compared to a normal caloric meal (533 kcal) or high-fat high-caloric meal (928 kcal). When EDURANT® was taken with only a protein-rich nutritional drink, exposures were 50% lower than when taken with a meal. EDURANT® **must be taken with a meal** to obtain optimal absorption. Taking EDURANT® in fasted condition or with only a

nutritional drink may result in decreased plasma concentrations of rilpivirine, which could potentially reduce the therapeutic effect of EDURANT® (see *Dosage and Administration*).

Distribution
Rilpivirine is approximately 99.7% bound to plasma proteins *in vitro*, primarily to albumin. The distribution of rilpivirine into compartments other than plasma (e.g., cerebrospinal fluid, genital tract secretions) has not been evaluated in humans.

Metabolism
In vitro experiments indicate that rilpivirine primarily undergoes oxidative metabolism mediated by the cytochrome P450 (CYP) 3A system.

Elimination
The terminal elimination half-life of rilpivirine is approximately 45 hours. After single dose oral administration of ^{14}C-rilpivirine, on average 85% and 6.1% of the radioactivity could be retrieved in faeces and urine, respectively. In faeces, unchanged rilpivirine accounted for on average 25% of the administered dose. Only trace amounts of unchanged rilpivirine (< 1% of dose) were detected in urine.

Special Populations
Pediatrics (less than 18 years of age)
The pharmacokinetics of rilpivirine in antiretroviral treatment-naïve HIV-1 infected pediatric subjects 12 to less than 18 years of age receiving EDURANT® 25 mg once daily were comparable to those in treatment-naïve HIV-1 infected adults receiving EDURANT® 25 mg once daily. There was no impact of body weight on rilpivirine pharmacokinetics in pediatric subjects in trial C213 (33 to 93 kg), similar to what was observed in adults.

The pharmacokinetics of rilpivirine in pediatric patients less than 12 years of age are under investigation. Dosing recommendations for pediatric patients less than 12 years of age cannot be made due to insufficient data (see *Dosage and Administration*).

Elderly (65 years of age and older)
Population pharmacokinetic analysis in HIV infected patients showed that rilpivirine pharmacokinetics are not different across the age range (18 to 78 years) evaluated, with only 3 subjects aged 65 years or older. No dose adjustment of EDURANT® is required in elderly patients. EDURANT® should be used with caution in this population (see *Dosage and Administration*).

Renal impairment
The pharmacokinetics of rilpivirine have not been studied in patients with renal insufficiency. Renal elimination of rilpivirine is negligible. No dose adjustment is needed for patients with mild or moderate renal impairment. In patients with severe renal impairment or end-stage renal disease, EDURANT® should be used with caution, as plasma concentrations may be increased due to alteration of drug absorption, distribution and/or metabolism secondary to renal dysfunction. In patients with severe renal impairment or end-stage renal disease, the combination of EDURANT® with a strong CYP3A inhibitor should only be used if the benefit outweighs the risk. As rilpivirine is highly bound to plasma proteins, it is unlikely that it will be significantly removed by haemodialysis or peritoneal dialysis (see *Dosage and Administration*).

Hepatic impairment

Rilpivirine is primarily metabolised and eliminated by the liver. In a study comparing 8 patients with mild hepatic impairment (Child-Pugh score A) to 8 matched controls, and 8 patients with moderate hepatic impairment (Child-Pugh score B) to 8 matched controls, the multiple dose exposure of rilpivirine was 47% higher in patients with mild hepatic impairment and 5% higher in patients with moderate hepatic impairment. However, it may not be excluded that the pharmacologically active, unbound, rilpivirine exposure is significantly increased in moderate hepatic impairment. No dose adjustment is suggested but caution is advised in patients with moderate hepatic impairment. EDURANT® has not been studied in patients with severe hepatic impairment (Child-Pugh score C). Therefore, EDURANT® is not recommended in patients with severe hepatic impairment (see *Dosage and Administration*).

Hepatitis B and/or hepatitis C virus co-infection

Population pharmacokinetic analysis indicated that hepatitis B and/or C virus co-infection had no clinically relevant effect on the exposure to rilpivirine.

Pregnancy and Postpartum

The exposure to total rilpivirine after intake of rilpivirine 25 mg once daily as part of an antiretroviral regimen was lower during pregnancy (similar for the 2nd and 3rd trimester) compared with postpartum (see Table 10). The decrease in unbound (i.e., active) rilpivirine pharmacokinetic parameters during pregnancy compared to postpartum was less pronounced than for total rilpivirine.

In women receiving rilpivirine 25 mg once daily during the 2nd trimester of pregnancy, mean intra-individual values for total rilpivirine C_{max}, AUC_{24h} and C_{min} values were, respectively, 21%, 29% and 35% lower as compared to postpartum; during the 3rd trimester of pregnancy, C_{max}, AUC_{24h} and C_{min} values were, respectively, 20%, 31% and 42% lower as compared to postpartum.

Table 10: Pharmacokinetic Results of Total Rilpivirine After Administration of Rilpivirine 25 mg Once Daily as Part of an Antiretroviral Regimen, During the 2nd Trimester of Pregnancy, the 3rd Trimester of Pregnancy and Postpartum

Pharmacokinetics of total rilpivirine (mean ± SD, t_{max}: median [range])	**Postpartum (6-12 Weeks) (n=11)**	**2nd Trimester of pregnancy (n=15)**	**3rd Trimester of pregnancy (n=13)**
C_{min}, ng/mL	84.0 ± 58.8	54.3 ± 25.8	52.9 ± 24.4
C_{max}, ng/mL	167 ± 101	121 ±45.9	123 ± 47.5
t_{max}, h	4.00 (2.03-25.08)	4.00 (1.00-9.00)	4.00 (2.00-24.93)
AUC_{24h}, ng.h/mL	2714 ± 1535	1792 ± 711	1762 ± 662

Other populations

Gender

No clinically relevant differences in the pharmacokinetics of rilpivirine have been observed between men and women.

Race

Population pharmacokinetic analysis of rilpivirine in HIV infected patients indicated that race had no clinically relevant effect on the exposure to rilpivirine.

NON-CLINICAL INFORMATION

General toxicology studies

Animal toxicology studies have been conducted with rilpivirine in mice, rats, rabbits, dogs and cynomolgus monkeys. The target organs and systems of toxicity were the adrenal cortex and the associated steroid biosynthesis (mouse, rat, dog, cynomolgus monkey), the reproductive organs (female mouse, male and female dog), the liver (mouse, rat, dog), the thyroid and pituitary gland (rat), the kidney (mouse, dog), the hematopoietic system (mouse, rat, dog), and the coagulation system (rat).

Repeated dose toxicity

Liver toxicity associated with liver enzyme induction was observed in rodents. In dogs, cholestasis- like effects were noted.

Reproductive toxicology studies

In a study conducted in rats, there were no effects on mating or fertility with rilpivirine up to 400 mg/kg/day, a dose of rilpivirine that showed maternal toxicity. This dose is associated with an exposure that is approximately 40 times higher than the exposure in humans at the recommended dose of 25 mg q.d. Studies in animals have shown no evidence of relevant embryonic or foetal toxicity or an effect on reproductive function. There was no teratogenicity with rilpivirine in rats and rabbits. The exposures at the embryo-fetal No Observed Adverse Effects Levels (NOAELs) in rats and rabbits were respectively 15 and 70 times higher than the exposure in humans at the recommended dose of 25 mg q.d. In a pre- and postnatal development assessment in rats, rilpivirine had no effect on development of offspring during lactation or post weaning when the mothers were dosed up to 400 mg/kg/day.

Carcinogenesis and mutagenesis

Rilpivirine was evaluated for carcinogenic potential by oral gavage administration to mice and rats up to 104 weeks. Daily doses of 20, 60 and 160 mg/kg/day were administered to mice and doses of 40, 200, 500 and 1500 mg/kg/day were administered to rats. An increase in the incidences of hepatocellular adenomas and carcinomas was observed in mice and rats. An increase in the incidences of follicular cell adenomas and/or carcinomas in the thyroid gland was observed in rats. Administration of rilpivirine did not cause a statistically significant increase in the incidence of any other benign or malignant neoplasm in mice or rats. The observed hepatocellular findings in mice and rats may be rodent-specific, associated with liver enzyme induction. The follicular cell findings may be rat-specific, associated with increased clearance of thyroxine. At the lowest tested doses in the carcinogenicity studies, the systemic exposures (based on AUC) to rilpivirine were 21-fold (mice) and 3-fold (rats), relative to those observed in humans at the recommended dose (25 mg q.d.).

Rilpivirine has tested negative in the *in vitro* Ames reverse mutation assay, *in vitro* chromosomal aberration assay in human lymphocyte and *in vitro* clastogenicity mouse lymphoma assay, tested in the absence and presence of a metabolic activation system. Rilpivirine did not induce chromosomal damage in the *in vivo* micronucleus test in mice.

PHARMACEUTICAL INFORMATION

List of Excipients

Tablet core

Croscarmellose sodium
Lactose monohydrate
Magnesium stearate
Polysorbate 20

Povidone K30
Silicified microcrystalline cellulose

Tablet coating
Hypromellose 2910 6mPa.s
Lactose monohydrate
Polyethylene glycol 3000/Macrogol 3000
Titanium dioxide
Triacetin

Incompatibilities
Not applicable.

Shelf Life
Observe expiry date on the outer pack.

Storage Conditions
Do not store above 30ºC.
Store in the original bottle in order to protect from light. Keep out of the sight and reach of children.

Nature and Contents of Container
75 mL high density polyethylene (HDPE) bottle with a polypropylene (PP) child resistant closure and induction seal liner. Each carton contains one bottle of 30 tablets.

Instructions for Use and Handling and Disposal
No special requirements.

PRODUCT REGISTRANT
Johnson & Johnson Pte Ltd
2 Science Park Drive,
#07-13, Ascent
Singapore Science Park 1
Singapore 118222

BATCH RELEASER
Janssen-Cilag S.p.A
Via C. Janssen
Borgo S. Michele
04100 Latina
Italy

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