

Product Monograph

Pr **Atripla**[®]

**(Efavirenz/Emtricitabine/
Tenofovir Disoproxil Fumarate Tablets)**

600 mg/200 mg/300 mg

Antiretroviral Agent

Bristol-Myers Squibb & Gilead Sciences, LLC
Foster City, CA 94404
USA

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Gilead Sciences Canada, Inc.
Mississauga, ON L5N 2W3

Bristol-Myers Squibb Canada
Montreal, QC H4S 0A4

www.gilead.com

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PART I. HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form/Strength	Clinically Relevant Nonmedicinal Ingredients
Oral	Tablet Efavirenz 600 mg / Emtricitabine 200 mg / Tenofovir Disoproxil Fumarate 300 mg	None <i>For a complete listing, see the DOSAGE FORMS, COMPOSITION, AND PACKAGING section.</i>

INDICATIONS AND CLINICAL USE

ATRIPLA is indicated for use alone as a complete regimen or in combination with other antiretroviral agents for the treatment of HIV-1 infection in adults. (see **CLINICAL TRIALS, Description of Clinical Studies and ADVERSE REACTIONS**).

Geriatrics (>65 years of age):

Clinical studies of the components of ATRIPLA did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects.

Pediatrics (<18 years of age):

Safety and effectiveness in pediatric patients have not been established. ATRIPLA is not recommended for pediatric administration.

CONTRAINDICATIONS

ATRIPLA is contraindicated in patients with previously demonstrated hypersensitivity to any of the components of the product. For a complete listing, see the **DOSAGE FORMS, COMPOSITION AND PACKAGING** section of the Product Monograph.

Competition for CYP3A4 by efavirenz could result in inhibition of metabolism of some drugs and create the potential for serious and/or life-threatening adverse events. Drugs that are contraindicated with ATRIPLA are listed in Table 1. See also **DRUG INTERACTIONS, Drug-Drug Interactions**.

Table 1. Drugs That Should Not Be Coadministered with ATRIPLA

Drug Class: Drug Name	Clinical Comment
Antifungal: voriconazole	CONTRAINDICATED because efavirenz significantly decreases voriconazole plasma concentrations, and coadministration may decrease the therapeutic effectiveness of voriconazole. Also, voriconazole significantly increases efavirenz plasma concentrations, which may increase the risk of efavirenz-associated side effects. Because ATRIPLA is a fixed dose combination product, the dose of efavirenz cannot be altered; therefore, voriconazole and ATRIPLA cannot be coadministered.
Antimigraine: 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.
Benzodiazepines: midazolam, triazolam	CONTRAINDICATED due to potential for serious and/or life-threatening reactions such as prolonged or increased sedation or respiratory depression.
Neuroleptic: pimozide	CONTRAINDICATED due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.
GI motility agent: cisapride*	CONTRAINDICATED due to potential for serious and/or life-threatening reactions such as cardiac arrhythmias.

*Cisapride is not marketed in Canada.

WARNINGS AND PRECAUTIONS

Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of nucleoside analogs including VIREAD, a component of ATRIPLA, alone or in combination with other antiretrovirals (see **WARNINGS AND PRECAUTIONS: Hepatic/Biliary/Pancreatic**).

ATRIPLA is not approved for the treatment of chronic hepatitis B virus (HBV) infection and the safety and efficacy of ATRIPLA have not been established in patients coinfecting with HBV and HIV. Severe acute exacerbations of hepatitis B have been reported in patients who are coinfecting with HBV and HIV after the discontinuation of EMTRIVA[®] or VIREAD[®], two of the components of ATRIPLA. In some patients infected with HBV and treated with EMTRIVA, the exacerbations of hepatitis B were associated with liver decompensation and liver failure. Hepatic function should be closely monitored with both clinical and laboratory follow-up for at least several months in patients who discontinue ATRIPLA and are coinfecting with HIV and HBV. If appropriate, initiation of anti-hepatitis B therapy may be warranted (see **WARNINGS AND PRECAUTIONS: Special populations**).

Renal failure, renal insufficiency, elevated creatinine, hypophosphatemia, and Fanconi syndrome have been reported with the use of tenofovir DF during clinical practice (see **WARNINGS AND PRECAUTIONS**).

General

As a fixed-dose combination of efavirenz, emtricitabine and tenofovir DF, ATRIPLA should not be administered with SUSTIVA[®], EMTRIVA, VIREAD, or TRUVADA[®]. Due to similarities between emtricitabine and lamivudine, ATRIPLA should not be administered with drugs containing lamivudine including Combivir[®], 3TC[®], Heptovir[®], Kivexa[®], and Trizivir[®].

ATRIPLA should not be administered with HEPSERA[®] (adefovir dipivoxil).

Clinical trial data of patients who were switched to ATRIPLA from a prior PI-containing regimen suggest that they may have lower response rates than patients who remain on their baseline regimens (see CLINICAL TRIALS section). Therefore, patients who are switched to ATRIPLA from a prior PI-containing regimen should be monitored for both treatment-emergent adverse events and rises in viral load.

Concomitant use of St. John's Wort (*Hypericum perforatum*) or St. John's Wort-containing products with ATRIPLA is not recommended. Coadministration of non-nucleoside reverse transcriptase inhibitors (NNRTIs), including efavirenz, with St. John's Wort is expected to substantially decrease NNRTI concentrations. Decreased concentrations may result in suboptimal levels of efavirenz and lead to loss of virologic response and possible resistance to efavirenz or to the class of NNRTIs.

Efavirenz may cause dizziness, impaired concentration, and/or drowsiness. Patients should be instructed that, if they experience these symptoms, they should avoid potentially hazardous tasks such as driving or operating machinery. (See **WARNINGS AND PRECAUTIONS: Neurologic**).

Carcinogenesis, Mutagenesis, Impairment of Fertility

Efavirenz:

Oral carcinogenicity studies in mice and rats were carried out with efavirenz. Mice were dosed with 0, 25, 75, 150, or 300 mg/kg/day for 2 years. Incidences of hepatocellular adenomas and carcinomas and pulmonary alveolar/bronchiolar adenomas at all doses were increased above background in females. No increases in tumor incidence above background were seen in males. The systemic exposure (based on AUCs) in mice was approximately 1.7-fold that in humans receiving the 600-mg/day dose. Rats were administered efavirenz at doses of 0, 25, 50, or 100 mg/kg/day for 2 years; no increases in tumor incidence above background were observed. The exposure in rats was lower than that in humans.

The mechanism of the carcinogenic potential is unknown. However, in genetic toxicology assays, efavirenz showed no evidence of mutagenic or clastogenic activity in a battery of in vitro and in vivo studies. These included bacterial mutation assays in *S. typhimurium* and *E. coli*, mammalian mutation assays in Chinese hamster ovary cells, chromosome aberration assays in human peripheral blood lymphocytes or Chinese hamster ovary cells, and an

in vivo mouse bone marrow micronucleus assay. Given the lack of genotoxic activity of efavirenz, the relevance to humans of neoplasms in efavirenz-treated mice is not known.

Efavirenz did not impair mating or fertility of male or female rats, and did not affect sperm of treated male rats. The reproductive performance of offspring born to female rats given efavirenz was not affected. As a result of the rapid clearance of efavirenz in rats, systemic drug exposures achieved in these studies were equivalent to or below those achieved in humans given therapeutic doses of efavirenz.

Emtricitabine:

In long-term oral carcinogenicity studies of emtricitabine, no drug-related increase in tumor incidence was found in mice at doses up to 750 mg/kg/day (26 times the human systemic exposure at the therapeutic dose of 200 mg/day) or in rats at doses up to 600 mg/kg/day (31 times the human systemic exposure at the therapeutic dose).

Emtricitabine was not genotoxic in the in vitro reverse mutation bacterial test (Ames test), or in vivo mouse lymphoma or mouse micronucleus assays.

Emtricitabine did not affect fertility in male rats at approximately 140-fold or in male and female mice at approximately 60-fold higher exposures (AUC) than in humans given the recommended 200 mg daily dose. Fertility was normal in the offspring of mice exposed daily from before birth (in utero) through sexual maturity at daily exposures (AUC) of approximately 60-fold higher than human exposures at the recommended 200 mg daily dose.

Tenofovir disoproxil fumarate:

Tenofovir disoproxil fumarate (tenofovir DF) did not show any carcinogenic potential in a long-term oral carcinogenicity study in rats. A long-term oral carcinogenicity study in mice showed a low incidence of duodenal tumors, considered likely related to high local concentrations in the gastrointestinal tract at the high dose of 600 mg/kg/day. The mechanism of tumor formation in mice and potential relevance for humans is uncertain.

Tenofovir DF was mutagenic in the in vitro mouse lymphoma assay and negative in an in vitro bacterial mutagenicity test (Ames test). In an in vivo mouse micronucleus assay, tenofovir DF was negative at doses up to 2000 mg/kg when administered orally to male mice.

There were no effects on fertility, mating performance or early embryonic development when tenofovir disoproxil fumarate was administered at 600 mg/kg/day to male rats for 28 days prior to mating and to female rats for 15 days prior to mating through day seven of gestation. There was, however, an alteration of the estrous cycle in female rats. A dose of 600 mg/kg/day is equivalent to 19 times the human dose based on body surface area comparisons.

Endocrine and Metabolism

Fat Redistribution:

Redistribution/accumulation of body fat (lipodystrophy) 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.

Hepatic/ Biliary/Pancreatic

Lactic Acidosis/Severe Hepatomegaly with Steatosis:

Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of antiretroviral nucleoside analogs including the tenofovir DF component of ATRIPLA, alone or in combination with other antiretrovirals in the treatment of HIV infection. A majority of these cases have been in women. Obesity and prolonged nucleoside exposure may be risk factors. Particular caution should be exercised when administering nucleoside analogs to any patient with known risk factors for liver disease; however, cases have also been reported in patients with no known risk factors. Treatment with ATRIPLA should be suspended in any patient who develops clinical or laboratory findings suggestive of lactic acidosis or pronounced hepatotoxicity (which may include hepatomegaly and steatosis even in the absence of marked transaminase elevations).

Hepatic Impairment:

The pharmacokinetics of efavirenz has not been adequately studied in patients with hepatic impairment. Because of the extensive cytochrome P450-mediated metabolism of efavirenz and limited clinical experience in patients with hepatic impairment, caution should be exercised in administering ATRIPLA to these patients. Patients should be monitored carefully for adverse events, and laboratory tests to evaluate the liver disease should be performed at periodic intervals.

Tenofovir and tenofovir disoproxil are not metabolized by liver enzymes. Clinically relevant pharmacokinetic changes in patients with hepatic impairment are not observed.

Emtricitabine has not been evaluated in patients with hepatic impairment; however, emtricitabine has not been shown to be metabolized by liver enzymes, so the impact of liver impairment is likely to be limited.

The safety and efficacy of ATRIPLA have not been established or specifically studied in patients with underlying liver disorders. Patients with chronic hepatitis B or C and treated with antiretroviral therapy are at increased risk for severe and potentially fatal hepatic adverse events. In case of concomitant antiviral therapy for hepatitis B or C, please refer also to the relevant product information for these medicinal products (**see WARNINGS AND PRECAUTIONS: Special Populations**).

Pancreatitis:

Pancreatitis has occurred during therapy with combination regimens that included tenofovir DF. Caution should be used when administering nucleoside analogues (including ATRIPLA) to patients with a history of pancreatitis or risk factors for the development of pancreatitis. Therapy should be suspended in patients with suspected pancreatitis.

In controlled clinical studies the rate of clinical pancreatitis was similar in patients receiving [1/1008 (0.1%)] and not receiving [2/635 (0.3%)] efavirenz. Asymptomatic increases in serum amylase levels were observed in a significantly higher number of patients treated with efavirenz 600 mg than in control patients.

Elevated triglycerides have been reported in patients receiving efavirenz, in some cases to levels which can predispose a patient to pancreatitis. Among patients with elevated triglycerides, there have been no cases of pancreatitis. Because these triglyceride levels were not obtained in a fasting state, the exact clinical relevance of these measurements is not known.

Immune

Hypersensitivity Reactions:

In clinical trials, hypersensitivity reactions were uncommon (<1%) in patients treated with efavirenz.

Immune Reconstitution:

Immune reconstitution syndrome has been reported in patients treated with combination antiretroviral therapy, including the components of ATRIPLA. During the initial phase of treatment, patients responding to antiretroviral therapy may develop an inflammatory response to indolent or residual opportunistic infections (such as *Mycobacterium avium* infections, cytomegalovirus, *Pneumocystis jiroveci* pneumonia (PCP), and tuberculosis), which may necessitate further evaluation and treatment.

Musculoskeletal

Bone Effects:

Bone monitoring should be considered for HIV infected patients who have a history of pathologic bone fracture or are at risk for osteopenia. Although the effect of supplementation with calcium and vitamin D was not studied, such supplementation may be beneficial for all patients. If bone abnormalities are suspected then appropriate consultation should be obtained.

In a 144 week study of treatment-naïve patients, decreases from baseline in bone mineral density (BMD) were seen at the lumbar spine and hip in both arms of the study. At Week 144, there was a significantly greater mean percentage decrease from baseline in BMD at the lumbar spine in patients in the tenofovir DF + lamivudine + efavirenz group compared with patients in the stavudine + lamivudine + efavirenz group. Changes in BMD at the hip were similar in the two treatment groups. In both groups, the majority of the reduction in

BMD occurred in the first 24–48 weeks of the study and this reduction was sustained through Week 144. Twenty-eight percent of tenofovir DF-treated patients vs. 21% of the stavudine-treated patients lost at least 5% of BMD at the spine or 7% of BMD at the hip. Clinically relevant fractures (excluding fingers and toes) were reported in 4 patients in the tenofovir DF group and 6 patients in the stavudine group. In addition, there were significant increases in biochemical markers of bone metabolism (serum bone-specific alkaline phosphatase, serum osteocalcin, serum C-telopeptide, and urinary N-telopeptide) in the tenofovir DF group relative to the stavudine group, suggesting increased bone turnover. Serum parathyroid hormone levels and 1,25 Vitamin D levels were also higher in the tenofovir DF group. Except for bone specific alkaline phosphatase, these changes resulted in values that remained within the normal range. The effects of tenofovir DF-associated changes in BMD and biochemical markers on long-term bone health and future fracture risk are unknown.

Cases of osteomalacia (associated with proximal renal tubulopathy and infrequently contributing to fractures) have been reported in association with the use of tenofovir DF (see ADVERSE REACTIONS, Post Market Adverse Drug Reactions: Tenofovir DF).

For additional information, please consult the VIREAD Product Monograph.

Neurologic

Nervous System Symptoms:

Fifty-three percent (531/1008) of patients receiving efavirenz in controlled clinical trials reported central nervous system symptoms (any grade, regardless of causality) compared to 25% (156/635) of patients receiving control regimens. These symptoms included, but were not limited to, dizziness (28.1% of the 1008 patients), insomnia (16.3%), impaired concentration (8.3%), somnolence (7.0%), abnormal dreams (6.2%), and hallucinations (1.2%). Other reported symptoms were euphoria, confusion, agitation, amnesia, stupor, abnormal thinking, and depersonalization. The majority of these symptoms were mild-moderate (50.7%); symptoms were severe in 2.0% of patients. Overall, 2.1% of patients discontinued therapy as a result. These symptoms usually begin during the first or second day of therapy and generally resolve after the first 2–4 weeks. After 4 weeks of therapy the prevalence of nervous system symptoms of at least moderate severity ranged from 5–9% in patients treated with regimens containing efavirenz and from 3–5% in patients treated with a control regimen. Patients should be informed that these symptoms are likely to improve with continued therapy. Dosing at bedtime improves tolerability of these symptoms.

Analysis of long-term data from a clinical study, (median follow-up 180 weeks, 102 weeks, and 76 weeks for patients treated with efavirenz + zidovudine + lamivudine, efavirenz + indinavir, and indinavir + zidovudine + lamivudine, respectively) showed that, beyond 24 weeks of therapy, the incidences of new-onset nervous system symptoms among efavirenz-treated patients were generally similar to those in the indinavir-containing control arm.

Patients receiving ATRIPLA should be alerted to the potential for additive central nervous system effects when ATRIPLA is used concomitantly with alcohol or psychoactive drugs. Patients should be informed that ATRIPLA may cause dizziness, impaired concentration,

and/or drowsiness. Patients should be instructed that if they experience these symptoms they should avoid potentially hazardous tasks such as driving or operating machinery. (See **WARNINGS AND PRECAUTIONS: General**).

Seizures:

Caution should be taken in any patient with any history of seizures. Convulsions have been observed infrequently in patients receiving efavirenz, generally in the presence of known medical history of seizures. Overall, the rate of seizure in controlled clinical trials has been 0.89% in efavirenz treated patients and 0.63% in the control patients. Patients who are receiving concomitant anticonvulsant medications primarily metabolized by the liver such as phenytoin, carbamazepine, and phenobarbital, may require periodic monitoring of plasma levels (See **DRUG INTERACTIONS**).

Psychiatric

Serious psychiatric adverse reactions have been reported in patients treated with efavirenz. In controlled trials of 1008 patients treated with regimens containing efavirenz for an average of 2.1 years and 635 patients treated with control regimens for an average of 1.5 years, the frequency (regardless of causality) of specific serious psychiatric events among patients who received efavirenz or control regimens respectively, were: severe depression (2.4%, 0.9%), suicidal ideation (0.7%, 0.3%), non-fatal suicide attempts (0.5%, 0%), aggressive behavior (0.4%, 0.5%), paranoid reactions (0.4%, 0.3%), and manic reactions (0.2%, 0.3%). When psychiatric symptoms similar to those noted above were combined and evaluated as a group in a multifactorial analysis of data from a clinical study, treatment with efavirenz was associated with an increase in the occurrence of these selected psychiatric symptoms. Other factors associated with an increase in the occurrence of these psychiatric symptoms were history of injection drug use, psychiatric history, and receipt of psychiatric medication at study entry; similar associations were observed in both the efavirenz and control treatment groups. In Study 006, onset of new serious psychiatric symptoms occurred throughout the study for both efavirenz-treated and control-treated patients. One percent of efavirenz-treated patients discontinued or interrupted treatment because of one or more of these selected psychiatric symptoms. There have also been occasional postmarketing reports of death by suicide, delusions, and psychosis-like behavior, although a causal relationship to the use of efavirenz cannot be determined from these reports. Patients with serious psychiatric adverse experiences should seek immediate medical evaluation to assess the probability that the symptoms may be related to the use of ATRIPLA, and if so, to determine whether the risks of continued therapy outweigh the benefits.

Renal

Nephrotoxicity:

Emtricitabine and tenofovir are principally eliminated by the kidney, however efavirenz is not. Since ATRIPLA is a combination product and the dose of the individual components cannot be altered, patients with creatinine clearance <50 mL/min should not receive ATRIPLA.

Renal impairment, including cases of acute renal failure and Fanconi syndrome (renal tubular injury with severe hypophosphatemia), has been reported in association with the use of tenofovir DF (**see ADVERSE REACTIONS, Post Marketing Experience**). The majority of these cases occurred in patients with underlying systemic or renal disease, or in patients taking nephrotoxic agents, however, some cases occurred in patients without identified risk factors.

It is recommended that creatine clearance be calculated in all patients prior to initiating therapy and as clinically appropriate during therapy with ATRIPLA. Routine monitoring of calculated creatine clearance and serum phosphorus should be performed in patients at risk for renal impairment, including patients who have previously experienced renal events while receiving HEPSERA.

ATRIPLA should be avoided with concurrent or recent use of a nephrotoxic agent. Examples of nephrotoxic agents include but are not limited to aminoglycosides, amphotericin B, foscarnet, ganciclovir, pentamidine, vancomycin, cidofovir, or interleukin-2. Particular caution should be exercised when administering ATRIPLA to patients with known risk factors for renal disease and a history of renal dysfunction; however, cases of renal failure have also been reported in patients with no known risk factors.

Skin

Rash:

In controlled clinical trials, 26% (266/1008) of patients treated with 600 mg efavirenz experienced new onset skin rash compared with 17% (111/635) of patients treated in control groups. Rash associated with blistering, moist desquamation or ulceration occurred in 0.9% (9/1008) of patients treated with efavirenz. Rashes are usually mild-to-moderate maculopapular skin eruptions that occur within the first 2 weeks of initiating therapy with efavirenz (median time to onset of rash in adults was 11 days) and in most patients continuing therapy with efavirenz, rash resolves within 1 month (median duration, 16 days). The discontinuation rate for rash in clinical trials was 6.4% (17/266) among patients with rash and 1.7% (17/1008) overall.

Grade 4 rash (including erythema multiforme, Stevens-Johnson syndrome, toxic epidermal necrolysis, and exfoliative dermatitis) was uncommon (<1%) in patients treated with efavirenz in clinical studies.

Experience with efavirenz in patients who discontinued other antiretroviral agents of the NNRTI class is limited. Nineteen patients who discontinued nevirapine because of rash have been treated with efavirenz. Nine of these patients developed mild-to-moderate rash while receiving therapy with efavirenz, and two of these patients discontinued because of rash.

ATRIPLA can be reinitiated in patients interrupting therapy because of Grades 1 and 2 rash. ATRIPLA should be discontinued in patients developing severe rash associated with blistering, desquamation, mucosal involvement or fever. Appropriate antihistamines and /or corticosteroids may improve the tolerability and hasten the resolution of rash.

Special Populations

Patients with HIV and Hepatitis B Virus Coinfection:

It is recommended that all patients with HIV be tested for the presence of HBV before initiating antiretroviral therapy. ATRIPLA is not approved for the treatment of chronic HBV infection and the safety and efficacy of ATRIPLA have not been established in patients co-infected with HBV and HIV. Severe acute exacerbations of hepatitis B have been reported in patients who are coinfected with HBV and HIV after the discontinuation of EMTRIVA and VIREAD, two of the components of ATRIPLA. In some patients infected with HBV and treated with EMTRIVA, the exacerbations of hepatitis B were associated with liver decompensation and liver failure. Hepatic function should be closely monitored with both clinical and laboratory follow-up for at least several months in patients who discontinue ATRIPLA and are co-infected with HIV and HBV. If appropriate, initiation of anti-hepatitis B therapy may be warranted. In patients with advanced liver disease or cirrhosis, post-treatment exacerbation of hepatitis may lead to hepatic decompensation. Therefore, in these patients, discontinuation of treatment without initiation of alternative anti-hepatitis B therapy is not recommended.

Pregnant Women:

Efavirenz may cause fetal harm when administered during the first trimester to a pregnant woman. Pregnancy should be avoided in women receiving ATRIPLA and for 12 weeks after discontinuation. Barrier contraception must always be used in combination with other methods of contraception (e.g., oral or other hormonal contraceptives) (see **DRUG INTERACTIONS** and **TOXICOLOGY**). Because of the long half-life of efavirenz, use of adequate contraceptive measures for 12 weeks after discontinuation of ATRIPLA is recommended. Women of childbearing potential should undergo pregnancy testing prior to initiation of ATRIPLA.

There are no adequate and well-controlled studies in pregnant women. ATRIPLA should be used in pregnant women only if the potential benefits outweigh the potential risks to the fetus, such as in pregnant women without other therapeutic options.

Efavirenz: As of July 2008, the Antiretroviral Pregnancy Registry has received prospective reports of 526 pregnancies exposed to efavirenz-containing regimens, nearly all of which were first-trimester exposures (507 pregnancies). Birth defects occurred in 13 of 407 live births (first-trimester exposure) and 2 of 37 live births (second-/third-trimester exposure). One of these prospectively reported defects with first-trimester exposure was a neural tube defect. A single case of anophthalmia with first-trimester exposure to efavirenz has also been prospectively reported; however, this case included severe oblique facial clefts and amniotic bonding, a known association with anophthalmia. There have been five retrospective reports of findings consistent with neural tube defects, including meningomyelocele. All mothers were exposed to efavirenz-containing regimens in the first trimester. A causal relationship of these events to the use of efavirenz cannot be established.

Malformations have been observed in 3 of 20 fetuses/infants from efavirenz-treated cynomolgus monkeys (versus 0 of 20 concomitant controls) in a developmental toxicity

study. The pregnant monkeys were dosed throughout pregnancy (postcoital days 20–150) with efavirenz 60 mg/kg daily, a dose which resulted in plasma drug concentrations similar to those in humans given 600 mg/day of efavirenz. Anencephaly and unilateral anophthalmia were observed in one fetus, microphthalmia was observed in another fetus, and cleft palate was observed in a third fetus. Efavirenz crosses the placenta in cynomolgus monkeys and produces fetal blood concentrations similar to maternal blood concentrations. Efavirenz has been shown to cross the placenta in rats and rabbits and produces fetal blood concentrations of efavirenz similar to maternal concentrations. An increase in fetal resorptions was observed in rats at efavirenz doses that produced peak plasma concentrations and AUC values in female rats equivalent to or lower than those achieved in humans given 600 mg once daily of efavirenz. Efavirenz produced no reproductive toxicities when given to pregnant rabbits at dose that produced peak plasma concentrations similar to and AUC values approximately half of those achieved in humans given 600 mg once daily of efavirenz.

Emtricitabine: The incidence of fetal variations and malformations was not increased in embryo-fetal toxicity studies performed with emtricitabine in mice at exposures (AUC) approximately 60-fold higher and in rabbits at approximately 120-fold higher than human exposures at the recommended daily dose.

Tenofovir DF: Reproduction studies have been performed in rats and rabbits at doses up to 14 and 19 times the human dose based on body surface area comparisons and revealed no evidence of impaired fertility or harm to the fetus due to tenofovir. Reduced pup body weights, survival and delay in sexual maturation was observed in a peri- and postnatal toxicity study in rats at the maternally toxic doses of 450 and 600 mg/kg (approximately 14 and 19 times the human dose based on body surface area comparisons).

Antiretroviral Pregnancy Registry: To monitor fetal outcomes of pregnant women exposed to ART (antiretroviral therapy) including ATRIPLA, an Antiretroviral Pregnancy Registry has been established. Healthcare providers are encouraged to register patients,
<http://www.apregistry.com>
Telephone: (800) 258-4263
Fax: (800) 800-1052.

Nursing Women:

HIV-1 infected mothers should not breast-feed their infants to avoid risking postnatal transmission of HIV. Studies in rats and rhesus monkeys have demonstrated that tenofovir is secreted in milk. Studies in rats have demonstrated that efavirenz is excreted in milk. It is not known whether efavirenz, emtricitabine or tenofovir is excreted in human milk. Because of both the potential for HIV transmission and the potential for serious adverse reactions in nursing infants, **mothers should be instructed not to breast-feed if they are receiving ATRIPLA.**

Pediatrics (<18 years of age): Safety and effectiveness in pediatric patients have not been established. ATRIPLA is not recommended for pediatric administration.

Geriatrics (>65 years of age): Clinical studies of efavirenz, emtricitabine or tenofovir DF did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently than younger subjects. In general, dose selection for the elderly patient should be cautious, keeping in mind the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

Monitoring and Laboratory Tests

Liver Enzymes: In patients with known or suspected history of Hepatitis B or C infection and in patients treated with other medications associated with liver toxicity, monitoring of liver enzymes is recommended. In patients with persistent elevations of serum transaminases to greater than five times the upper limit of the normal range, the benefit of continued therapy with ATRIPLA needs to be weighed against the unknown risks of significant liver toxicity. (See **ADVERSE REACTIONS: Laboratory Abnormalities** and **WARNINGS AND PRECAUTIONS: Renal**).

ADVERSE REACTIONS

Clinical Trial Adverse Drug Reactions

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

Adverse Drug Reaction Overview

The combination of efavirenz + emtricitabine + tenofovir DF has been studied in 460 patients either as the fixed-dose dose combination tablet ATRIPLA (Study 073) or as the component products (Study 934). Adverse reactions observed in Study 934 (Phase III) and Study 073 (Phase IV) were generally consistent with those seen in previous studies of the individual components.

The most common adverse events reported in Study 934 associated with study regimen of efavirenz + emtricitabine + tenofovir DF through Week 144 were dizziness (25%), nausea (18%), and abnormal dreams (17%). Study regimen was discontinued due to an adverse event for 5% of subjects (13/257) in the efavirenz +emtricitabine + tenofovir DF group and 11% of subjects (29/254) in the lamivudine/zidovudine + efavirenz group.

In study 073, the most frequently reported adverse events associated with ATRIPLA among patients treated up to 48 weeks were psychiatric disorders (16%); nervous system disorders (13%); and gastrointestinal disorders (7%). The majority of adverse events associated with ATRIPLA were mild in severity with dizziness (11%) and abnormal dreams (7%) being the most common specific adverse events reported. Adverse events that led to study drug discontinuation were experienced by 5% of subjects in the ATRIPLA group and 1% of subjects who stayed on their baseline regimen (SBR group); the majority of subjects who discontinued study drug due to adverse events had switched to ATRIPLA from a prior protease inhibitor-based regimen and the events that led to discontinuation were expected events consistent with the known safety profile of efavirenz e.g. nervous system symptoms.

Study 934

Study 934 – Treatment Emergent Adverse Events: Assessment of adverse reactions is based on data from Study 934 in which 511 antiretroviral-naïve patients received either emtricitabine + tenofovir DF administered in combination with efavirenz (N=257) or lamivudine/zidovudine administered in combination with efavirenz (N=254). Adverse events observed in this study were generally consistent with those seen in other studies in treatment experienced or treatment-naïve patients (Table 2). The most common adverse events that occurred in patients receiving efavirenz + emtricitabine + tenofovir DF were mild to moderate dizziness, nausea or diarrhea.

Table 2. Selected Treatment-Emergent Adverse Events (Grades 2–4) Reported in ≥3% in Any Treatment Group in Study 934 (0–48 Weeks)

	FTC+TDF+EFV	AZT/3TC+EFV
	N=257	N=254
Blood and Lymphatic System Disorders		
Anemia	<1%	5%
Gastrointestinal Disorder		
Diarrhea	7%	4%
Nausea	8%	6%
Vomiting	1%	4%
General Disorders and Administration Site Condition		
Fatigue	7%	6%
Infections and Infestations		
Sinusitis	4%	2%
Upper Respiratory Tract Infections	3%	3%
Nasopharyngitis	3%	1%
Nervous System Disorders		
Somnolence	3%	2%
Headache	5%	4%
Dizziness	8%	7%
Psychiatric Disorders		
Depression	4%	7%
Insomnia	4%	5%
Abnormal Dreams	4%	3%
Skin and Subcutaneous Tissue Disorders		
Rash	5%	4%

Patients who received efavirenz + emtricitabine + tenofovir DF up to 144 weeks in Study 934 reported adverse events similar in nature and severity to those reported in the first 48 weeks of treatment.

Laboratory Abnormalities: Laboratory abnormalities observed in this study were generally consistent with those seen in other studies (Table 3).

Table 3. Significant Laboratory Abnormalities Reported in $\geq 1\%$ in Any Treatment Group in Study 934 (0–48 Weeks)

	FTC+TDF+EFV	AZT/3TC+EFV
	N=257	N=254
Any \geq Grade 3 Laboratory Abnormality	25%	22%
Fasting Cholesterol (>240 mg/dL)	15%	17%
Creatine Kinase (M: >990 U/L) (F: >845 U/L)	7%	6%
Serum Amylase (>175 U/L)	7%	3%
Alkaline Phosphatase (>550 U/L)	1%	0%
AST (M: >180 U/L) (F: >170 U/L)	3%	2%
ALT (M: >215 U/L) (F: >170 U/L)	2%	2%
Hemoglobin (<8.0 mg/dL)	0%	3%
Hyperglycemia (>250 mg/dL)	1%	1%
Hematuria (>75 RBC/HPF)	2%	2%
Neutrophil (<750 /mm ³)	3%	4%
Fasting Triglycerides (>750 mg/dL)	4%	2%

Laboratory abnormalities in patients who received treatment up to 144 weeks in Study 934 were consistent with those observed in the first 48 weeks of treatment.

Through 48 weeks, 7 patients in the efavirenz + emtricitabine + tenofovir DF group and 5 patients in the efavirenz + lamivudine/zidovudine group experienced a new CDC Class C event (10 and 6 patients, respectively, through 144 weeks). Renal safety assessed by laboratory abnormalities was similar in the two groups and no patient discontinued study drug due to renal events. At Weeks 48 and 144, total limb fat (as measured by dual-energy x-ray absorptiometry) was significantly less in a subgroup of patients in the efavirenz + lamivudine/zidovudine group compared to the efavirenz + emtricitabine + tenofovir DF subgroup (see Table 4).

Table 4. Study 934 Total Limb Fat at Week 144* (Dual-Energy X-Ray Absorptiometry)

	FTC + TDF + EFV	AZT/3TC +EFV
Week 48¹	N=51	N=49
Total Limb Fat (kg) Mean ± S.D (Median)	8.9 ± 5.4 (7.4)	6.9 ± 3.9 (6.0)
Week 144²	N=145	N=124
Total Limb Fat (kg) Mean ± S.D (Median)	9.2 ± 5.4 (7.9)	6.5 ± 4.4 (5.4)
Change from Week 48 to Week 144³	N=48	N=38
Total Limb Fat (kg) Mean ± S.D (Median)	1.1 ± 1.9 (0.9)	-1.1 ± 1.7 (-0.8)

* Baseline data are not available.

¹P=0.03 for the comparison between arms

²P<0.001 for the comparison between arms

³P<0.001 for the comparison between arms; P<0.001 for the comparison within arms

Lipids:

In Study 934 at Week 144, the mean increase from baseline fasting triglyceride concentrations was 4 mg/dL for the tenofovir DF, emtricitabine and efavirenz group and 36 mg/dL for the zidovudine/lamivudine and efavirenz group. For fasting total, LDL, and HDL cholesterol concentrations, the mean increases from baseline were 24 mg/dL, 13 mg/dL, and 10 mg/dL, respectively, for the tenofovir DF group and 36 mg/dL, 16 mg/dL, and 12 mg/dL, respectively, for the zidovudine/lamivudine group. The differences between treatment groups reached statistical significance for fasting triglycerides (p=0.047) and fasting total serum cholesterol (p=0.005).

Hepatic Events:

In Study 934, 10 patients treated with efavirenz, emtricitabine, and tenofovir DF and 16 patients treated with efavirenz and fixed-dose zidovudine/lamivudine were Hepatitis C antibody positive. Among these HCV coinfecting patients, one patient (1/10) in the efavirenz, emtricitabine and tenofovir DF arm had elevations in ALT and AST to greater than five times ULN through 144 weeks. One patient (1/16) in the fixed-dose zidovudine/lamivudine arm had elevations in ALT to greater than five times ULN through 144 weeks, and one patient (1/16) in the fixed-dose zidovudine/lamivudine arm had elevations in AST to greater than five times ULN through 144 weeks. Nine patients treated with efavirenz, emtricitabine and tenofovir DF and 4 patients treated with efavirenz and fixed-dose zidovudine/lamivudine were Hepatitis B surface antigen positive. None of these patients had treatment-emergent elevations in ALT and AST to greater than five times ULN through 144 weeks. No HBV and/or HCV coinfecting patient discontinued from the study due to hepatobiliary disorders (see **WARNINGS AND PRECAUTIONS, Hepatic/Biliary/Pancreatic**).

Study 073

In study 073, patients with stable virologic suppression on antiretroviral therapy and no history of virologic failure were randomized to receive ATRIPLA or to stay on their baseline regimen. The adverse reactions observed in Study 073 were generally consistent with those seen in Study 934 and those seen with individual components of ATRIPLA when each was administered in combination with other antiretroviral agents.

In such an open label switch study, subjects may experience an adverse event or laboratory abnormality when they switch to a new treatment compared to subjects who remain on their baseline regimen.

Treatment-emergent AEs such as nervous system symptoms were more frequent in subjects who switched to ATRIPLA than those who stayed on their baseline regimen (SBR group). Such events can be expected in subjects receiving an efavirenz based regimen for the first time, and are consistent with the known safety profile of efavirenz.

Table 5. Selected Treatment-Emergent Adverse Events Reported for $\geq 3\%$ of Subjects in Any Treatment Group in Study 073 (0–48 Weeks)

	FTC+TDF+EFV	SBR
	N=203	N=97
Gastrointestinal Disorder		
Diarrhea	3%	0%
Nausea	3%	2%
Nervous System Disorders		
Somnolence	3%	0%
Dizziness	11%	1%
Psychiatric Disorders		
Insomnia	4%	0%
Abnormal Dreams	7%	0%

Laboratory abnormalities observed in the ATRIPLA group in Study 073 (Table 6) were generally consistent with those in Study 934.

Table 6. Laboratory Abnormalities (≥ Grade 3) Reported in ≥1% of Subjects in Any Treatment Group in Study 073 (0–48 Weeks)

	FTC+TDF+EFV	SBR
	N=203	N=97
Any ≥Grade 3 Laboratory Abnormality	9%	15%
Fasting Cholesterol (>300 mg/dL)	<1%	2%
Serum Amylase (>200 u/L)	3%	2%
AST (M: >180 U/L) (F: >170 U/L)	1%	1%
ALT (M: >215 U/L) (F: >170 U/L)	2%	0%
Hemoglobin (<7.5 g/dL)	0%	1%
Hyperglycemia (>250 mg/dL)	1%	1%
Neutrophil (<750/mm ³)	0%	1%
Fasting Triglycerides (>750 mg/dL)	2%	2%
Total Bilirubin (>3 mg/dL)	0	7%

In addition to the adverse events in Study 934 (Table 2) and Study 073 (Table 5), the following adverse events were reported in clinical studies of efavirenz, emtricitabine, or tenofovir DF in combination with other antiretroviral agents. This list is not all inclusive. For additional safety information about SUSTIVA (efavirenz), EMTRIVA (emtricitabine) or VIREAD (tenofovir DF), consult the Product Monographs for these products.

Efavirenz: The most significant adverse events observed in patients treated with efavirenz are nervous system symptoms, psychiatric symptoms, and rash (see **WARNINGS AND PRECAUTIONS**). Selected clinical adverse events of moderate or severe intensity reported in ≥2% of efavirenz-treated patients in three controlled clinical trials included abdominal pain, dyspepsia, anorexia, nausea, vomiting, diarrhea, anxiety, insomnia, dizziness, impaired concentration, nervousness, fatigue, pain, abnormal dreams, headache, depression, rash, somnolence, lipodystrophy, and pruritus.

Other adverse events of moderate or severe intensity reported in <2% of patients in all Phase II/III studies as well as postmarketing spontaneous reports included:

Body as a Whole: alcohol intolerance, allergic reaction, asthenia, fever, hot flushes, influenza-like symptoms, malaise, pain, peripheral edema, syncope, dysregulated body temperature, flank pain, hypersensitivity reactions. Redistribution/accumulation of body fat (see **Warnings and PRECAUTIONS, Endocrine and Metabolism, Fat Redistribution**).

Cardiovascular: arrhythmia, flushing, palpitations, tachycardia, thrombophlebitis, hypertension, congestive heart failure, chest pain

Central and Peripheral Nervous System: ataxia, confusion, convulsions, impaired coordination, migraine headaches, neuralgia, parasthesia, hypoesthesia, peripheral neuropathy, speech disorder, stupor, tremor, neuromuscular paresis, paranoid reaction

Gastrointestinal: dry mouth, pancreatitis, constipation, malabsorption

Liver and Biliary System: hepatic enzymes increased (including ALT, AST and GGT), hepatitis, jaundice, hepatomegaly

Metabolic and Nutritional: hypercholesterolemia, hypertriglyceridemia

Miscellaneous: thrombocytopenia, proteinuria, anemia, pancytopenia, increased sweating

Psychiatric: aggressive reactions, abnormal thinking, aggravated depression, agitation, delusions, amnesia, anxiety, apathy, delirium, depersonalization, emotional lability, euphoria, hallucination, manic reaction, psychosis, neurosis, paranoia, suicide, suicidal ideation, non-fatal suicide attempts

Respiratory: asthma, apnea, dyspnea

Skin and Appendages: acne, alopecia, eczema, folliculitis, skin defoliation, urticaria, erythema nodosum, erythema multiforme, Stevens-Johnson syndrome, verruca, nail disorders, skin disorders, photosensitivity reaction

Special Senses: abnormal vision, diplopia, glaucoma, iritis, parosmia, taste perversion, tinnitus

Urinary System: polyuria

For more information, please consult the SUSTIVA Product Monograph.

Emtricitabine and Tenofovir: Adverse events that occurred in at least 3-5% of patients receiving emtricitabine or tenofovir DF with other antiretroviral agents in clinical trials include: anorexia, anxiety, arthralgia, asthenia, increased cough, depressive disorders, dyspepsia, fever, flatulence, myalgia, pain, abdominal pain, back pain, chest pain, paresthesia, peripheral neuropathy (including peripheral neuritis and neuropathy), pneumonia, rhinitis, rash event (including rash, pruritus, maculopapular rash, urticaria, vesiculobullous rash, pustular rash and allergic reaction), sweating, and weight loss. Skin discoloration has been reported with higher frequency among emtricitabine treated patients.

Skin discoloration, manifested by hyperpigmentation on the palms and/or soles was generally mild and asymptomatic and of little clinical significance. The mechanism is unknown. In addition to the laboratory abnormalities described for Study 934 (Table 3), Grade 3/4 elevations of bilirubin ($>2.5 \times \text{ULN}$), pancreatic amylase ($>2.0 \times \text{ULN}$), serum glucose (<40 or >250 mg/dL), serum lipase ($>2.0 \times \text{ULN}$), and urine glucose ($\geq 3+$) occurred in up to 3% of patients treated with emtricitabine or tenofovir DF with other antiretroviral agents in clinical trials. For more information, please consult the VIREAD and EMTRIVA Product Monographs.

Post Market Adverse Drug Reactions

In addition to the adverse reaction reports from clinical trials, the following possible adverse reactions have been identified during post-approval use of efavirenz, emtricitabine, or tenofovir DF. Because these events have been reported voluntarily from a population of unknown size, estimates of frequency cannot be made. These events have been considered possible adverse reactions due to a combination of their seriousness, frequency of reporting or potential causal relationship with treatment.

Efavirenz:

Additional adverse experiences* reported in postmarketing surveillance include:

<i>Endocrine</i>	Gynecomastia
<i>Eye disorders</i>	Blurred vision
<i>Hepatobiliary disorders</i>	Hepatic failure
<i>Immune System disorders</i>	Immune reconstitution syndrome
<i>Musculoskeletal</i>	Rhabdomyolysis
<i>Nervous system disorders</i>	Cerebellar coordination and balance disturbances
<i>Psychiatric</i>	Neurosis
<i>Renal and urinary disorders</i>	Increased CPK
<i>Skin and subcutaneous tissue disorders</i>	Photoallergic dermatitis

*Those adverse events not already included under **Clinical Trial Adverse Drug Reactions, Efavirenz**

Additional cases of pancreatitis have been reported in postmarketing surveillance. Please see **WARNINGS AND PRECAUTIONS: Hepatic/Biliary/Pancreatitis**.

Emtricitabine:

The following adverse experiences have been reported in post-marketing experience without regard to causality; some events represent a single report.

Blood and lymphatic system disorders: Thrombocytopenia

Gastrointestinal disorders: Pancreatitis

General disorders and administrative site conditions: Pyrexia

Metabolism and nutrition disorders: Lactic acidosis

Tenofovir DF:

Immune system disorders: Allergic reaction (including angioedema)

Metabolism and nutrition disorders: Lactic acidosis, hypokalemia, hypophosphatemia

Respiratory, thoracic and mediastinal disorders: Dyspnea

Gastrointestinal disorders: Pancreatitis, increased amylase, abdominal pain

Hemic and lymphatic system disorders: Thrombocytopenia

Hepatobiliary disorders: Hepatic steatosis, hepatitis, increased liver enzymes (most commonly AST, ALT, GGT)

Skin and Subcutaneous Tissue Disorders: Rash

Musculoskeletal and Connective Tissue Disorders: Rhabdomyolysis, osteomalacia (manifested as bone pain and infrequently contributing to fractures), muscular weakness, myopathy

Renal and urinary disorders: Acute renal failure, renal failure, acute tubular necrosis, Fanconi syndrome, proximal renal tubulopathy, interstitial nephritis (including acute cases), nephrogenic diabetes insipidus, renal insufficiency, increased creatinine, proteinuria, polyuria

General disorders and administrative site conditions: Asthenia

The following adverse reactions, listed under the body system headings above, sometimes appeared to be concurrent with proximal renal tubulopathy: rhabdomyolysis, osteomalacia (manifested as bone pain and infrequently contributing to fractures), hypokalemia, muscular weakness, myopathy, hypophosphatemia.

There have been three post marketing reports of acute renal failure in patients on concomitant NSAIDS therapy where a relationship to tenofovir DF could not be excluded. These events mostly occurred in medically complex patients, where underlying disease processes confound interpretation.

Immune Reconstitution Syndrome

In HIV-infected patients with severe immune deficiency at the time of initiation of antiretroviral therapy an inflammatory reaction to infectious pathogens (active or inactive) may arise (see **WARNINGS and PRECAUTIONS**).

Exacerbations of Hepatitis after Discontinuation of Treatment

Tenofovir DF and emtricitabine-containing products: In HIV infected patients coinfecting with HBV, clinical and laboratory evidence of exacerbations of hepatitis have occurred after discontinuation of treatment (see **WARNINGS AND PRECAUTIONS: Special Populations - Patients with HIV and Hepatitis B Virus Coinfection**).

DRUG INTERACTIONS

Drug-Drug Interactions

No drug interaction studies have been conducted with ATRIPLA tablets. The drug interactions described in Tables 7 and 8 are based on studies conducted with efavirenz, emtricitabine or tenofovir DF as individual agents or are potential drug interactions. The tables include potentially significant interactions, but may not be inclusive of all potential interactions.

Table 7. Drugs That Are Not Recommended for Use With ATRIPLA

Drug Class: Drug Name	Clinical Comment
Antiretrovirals (Related Drugs): EMTRIVA (emtricitabine), VIREAD (tenofovir DF), TRUVADA (emtricitabine/tenofovir DF), SUSTIVA (efavirenz), HEPSERA (adefovir dipivoxil)	Not for use with ATRIPLA because the active ingredients of EMTRIVA (emtricitabine), VIREAD (tenofovir DF), TRUVADA (emtricitabine/tenofovir DF) and SUSTIVA (efavirenz) are components of ATRIPLA. Co-administration of ATRIPLA with HEPSERA is not recommended due to the possible risk of renal toxicity.
Antiretrovirals (Similar Drugs): 3TC (lamivudine), Combivir (lamivudine/zidovudine), Heptovir (lamivudine-HBV), Kivexa (abacavir sulfate/lamivudine), Trizivir (abacavir sulfate/lamivudine/zidovudine)	Lamivudine (3TC), which is similar to emtricitabine, is a component of Combivir, Heptovir, Kivexa, and Trizivir.
Protease inhibitor: atazanavir	NOT RECOMMENDED: Sufficient data are not available to make a dosing recommendation for atazanavir or atazanavir/ritonavir with ATRIPLA. Plasma concentrations of atazanavir were decreased by both efavirenz and tenofovir DF. Atazanavir has been shown to increase tenofovir concentrations.
St. John's Wort (<i>Hypericum perforatum</i>)	NOT RECOMMENDED: Expected to substantially decrease plasma levels of efavirenz; has not been studied in combination with efavirenz.

*Cisapride is not marketed in Canada.

Table 8. Established¹ and Other Potentially Significant² Drug Interactions: Alteration in Dose or Regimen May Be Recommended Based on Drug Interaction Studies or Predicted Interaction

Concomitant Drug Class: Drug Name	Effect³	Clinical Comment
<i>Antiretroviral Agents</i>		
Protease Inhibitor: Amprenavir	↓ amprenavir concentration	Efavirenz has the potential to decrease serum concentrations of amprenavir.
Protease Inhibitor: Fosamprenavir calcium	↓ amprenavir concentration	Fosamprenavir (unboosted): Appropriate doses of fosamprenavir and ATRIPLA with respect to safety and efficacy have not been established. Fosamprenavir/ritonavir: An additional 100 mg/day (300 mg total) of ritonavir is recommended when ATRIPLA is administered with fosamprenavir/ritonavir once daily. No change in the ritonavir dose is required when ATRIPLA is administered with fosamprenavir plus ritonavir twice daily.

Concomitant Drug Class: Drug Name	Effect ³	Clinical Comment
Protease Inhibitor: Indinavir	↓ indinavir concentration	The optimal dose of indinavir, when given in combination with efavirenz, is not known. Increasing the indinavir dose to 1000 mg every 8 hours does not compensate for the increased indinavir metabolism due to efavirenz.
Protease Inhibitor: Lopinavir/ritonavir	↓ lopinavir concentration ↑ tenofovir concentration	For lopinavir/ritonavir capsules or oral solution, a dose increase to 533/133 mg (4 capsules or 6.5 mL) twice daily taken with food is recommended when used in combination with efavirenz in patients for whom reduced susceptibility to lopinavir is clinically suspected (by treatment history or laboratory evidence). A dose increase of lopinavir/ritonavir tablets to 600/150 mg (3 tablets) twice daily may be considered when used in combination with efavirenz in treatment-experienced patients where decreased susceptibility to lopinavir is clinically suspected (by treatment history or laboratory evidence). Patients should be monitored for tenofovir-associated adverse events. ATRIPLA should be discontinued in patients who develop tenofovir-associated adverse events.
Protease Inhibitor: Ritonavir	↑ ritonavir concentration ↑ efavirenz concentration	When ritonavir 500 mg every 12 hours was coadministered with efavirenz 600 mg once daily, the combination was associated with a higher frequency of adverse clinical experiences (eg, dizziness, nausea, paresthesia) and laboratory abnormalities (elevated liver enzymes). Monitoring of liver enzymes is recommended when ATRIPLA is used in combination with ritonavir.
Protease Inhibitor: Saquinavir	↓ saquinavir concentration	Should not be used as sole protease inhibitor in combination with ATRIPLA.
NRTI: Didanosine	↑ didanosine concentration	Higher didanosine concentrations could potentiate didanosine-associated adverse events, including pancreatitis, and neuropathy. In adults weighing ≥60 kg, the didanosine EC dose should be reduced to 250 mg if coadministered with ATRIPLA. For patients with body weight <60 kg, and creatinine clearance ≥60 mL/min, the recommended dose of ddI-EC is 200 mg. Data are not available to recommend a dose adjustment of didanosine for patients with creatinine clearance <60mL/min. Coadministration of ATRIPLA and didanosine EC should be undertaken with caution and patients receiving this combination should be monitored closely for didanosine-associated adverse events. For additional information, please consult the Videx[®]/ Videx EC[®] (didanosine) prescribing information.

Concomitant Drug Class: Drug Name	Effect³	Clinical Comment
<i>Other Agents</i>		
Anticoagulant: Warfarin	↑ or ↓ warfarin concentration	Plasma concentrations and effects potentially increased or decreased by efavirenz.
Anticonvulsants: Carbamazepine	↓ carbamazepine concentration ↓ efavirenz concentration	Periodic monitoring of carbamazepine plasma levels should be conducted. There are insufficient data to make a dose recommendation for ATRIPLA. Alternative anticonvulsant treatment should be considered.
Anticonvulsants: Phenytoin Phenobarbital	↓ anticonvulsant concentration ↓ efavirenz concentration	Potential for reduction in anticonvulsant and/or efavirenz plasma levels; periodic monitoring of anticonvulsant plasma levels should be conducted.
Antidepressant: Sertraline	↓ sertraline concentration	It may be necessary to retitrate the sertraline dose to achieve the desired clinical effect. In a drug interaction study in healthy subjects, an increased incidence of impaired concentration was seen in subjects receiving sertraline concomitantly with efavirenz.
Antifungals: Itraconazole Ketoconazole	 ↓ itraconazole concentration ↓ hydroxy-itraconazole concentration ↓ ketoconazole concentration	See CONTRAINDICATIONS for other antifungals. Since no dose recommendation for itraconazole can be made, alternative antifungal treatment should be considered. Drug interaction studies with ATRIPLA and ketoconazole have not been conducted. Efavirenz has the potential to decrease plasma concentrations of ketoconazole.
Anti-infective: Clarithromycin	↓ clarithromycin concentration ↑ 14-OH metabolite concentration	Clinical significance unknown. In uninfected volunteers, 46% developed rash while receiving efavirenz and clarithromycin. No dose adjustment of ATRIPLA is recommended when given with clarithromycin. Alternatives to clarithromycin, such as azithromycin, should be considered. Not all macrolide antibiotics have been studied in combination with efavirenz.
Antimycobacterial: Rifabutin	↓ rifabutin concentration	Increase daily dose of rifabutin by 50%. Consider doubling the rifabutin dose in regimens where rifabutin is given 2 or 3 times a week.

Concomitant Drug Class: Drug Name	Effect³	Clinical Comment
Antimycobacterial: Rifampin	↓ efavirenz concentration	An additional 200 mg/day (800 mg total) of efavirenz is recommended when ATRIPLA is administered with rifampin
Calcium Channel Blockers: Diltiazem	↓ diltiazem ↓ desacetyl diltiazem ↓ N-mono desmethyl diltiazem	Diltiazem levels are markedly decreased when coadministered with efavirenz. Efavirenz levels decreased to a lesser extent (see Tables 9 and 10). Patients should be closely monitored for possible decreased diltiazem effects and increased adverse events and laboratory abnormalities associated with efavirenz. Refer to the prescribing information for diltiazem for guidance on dose adjustment.
Others (e.g. felodipine, nifedipine, verapamil)	↓ calcium channel blocker	No data are available on the potential interactions of efavirenz with other calcium channel blockers that are substrates of the CYP3A4 enzyme. The potential exists for reduction in plasma concentrations of the calcium channel blocker. Dose adjustments should be guided by clinical response (refer to the prescribing information for the calcium channel blocker).
Immunosuppressants: cyclosporine, tacrolimus, sirolimus	↓ cyclosporine, tacrolimus, sirolimus concentration	When an immunosuppressant metabolized by CYP3A4 is administered with efavirenz, decreased exposure of the immunosuppressant may be expected due to CYP3A4 induction. Immunosuppressants metabolized by CYP3A4 are not anticipated to affect exposure of efavirenz. There were no clinically significant pharmacokinetic interactions when emtricitabine/tenofovir DF was coadministered with tacrolimus. Dose adjustments of the immunosuppressant may be required when used with ATRIPLA. Close monitoring of immunosuppressant concentrations for at least 2 weeks (until stable concentrations are reached) is recommended when starting or stopping treatment with ATRIPLA.
HMG-CoA Reductase inhibitors: Atorvastatin Pravastatin Simvastatin	↓ atorvastatin concentration ↓ pravastatin concentration ↓ simvastatin concentration	Plasma concentrations of atorvastatin, pravastatin, and simvastatin decreased with efavirenz. Consult the complete prescribing information for the HMG-CoA reductase inhibitor for guidance on individualizing the dose.

Concomitant Drug Class: Drug Name	Effect³	Clinical Comment
Hormonal contraceptive: Oral: Ethinyl estradiol/ Norgestimate Implant: Etonogestrel	↓ active metabolites of norgestimate ↓ etonogestrel	A reliable method of barrier contraception must be used in addition to hormonal contraceptives. Efavirenz had no effect on ethinyl estradiol concentrations, but progestin levels (norelgestromin and levonorgestrel) were markedly decreased. The clinical significance of these effects is not known. No effect of ethinyl estradiol/norgestimate on efavirenz plasma concentrations was observed. The interaction between etonogestrel and efavirenz has not been studied. Decreased exposure of etonogestrel may be expected (CYP3A4 induction), and there have been occasional postmarketing reports of contraceptive failure with etonogestrel in efavirenz-exposed patients.
Narcotic Analgesic: Methadone	↓ methadone concentration	Coadministration of efavirenz in HIV-infected individuals with a history of injection drug use resulted in decreased plasma levels of methadone and signs of opiate withdrawal. Patients should be monitored for signs of withdrawal and their methadone dose increased as required to alleviate withdrawal symptoms.

1. See Tables 9-12.
2. This table is not all inclusive.
3. Increase = ↑; Decrease = ↓

Efavirenz:

Efavirenz has been shown in vivo to cause hepatic enzyme induction, thus increasing the biotransformation of some drugs metabolized by CYP3A4. In vitro studies have demonstrated that efavirenz inhibits 2C9, 2C19 and 3A4 isozymes in the range of observed efavirenz plasma concentrations. Coadministration of efavirenz with drugs primarily metabolized by these isozymes may result in altered plasma concentrations of the coadministered drug. Therefore, appropriate dose adjustments may be necessary for these drugs.

Drugs which induce CYP3A4 activity (e.g., phenobarbital, rifampin, rifabutin) would be expected to increase the clearance of efavirenz resulting in lowered plasma concentrations.

Drug interaction studies were performed with efavirenz and other drugs likely to be coadministered or drugs commonly used as probes for pharmacokinetic interaction. There was no clinically significant interaction observed between efavirenz and zidovudine, lamivudine, azithromycin, fluconazole, lorazepam, cetirizine, or paroxetine. Single doses of famotidine or an aluminum and magnesium antacid with simethicone had no effects on efavirenz exposures. The effects of coadministration of efavirenz on C_{max} and AUC are summarized in Table 9 (effect of other drugs on efavirenz) and Table 10 (effect of efavirenz on other drugs). (See **CONTRAINDICATIONS**.)

Specific drug interaction studies have not been performed with efavirenz and NRTIs other than lamivudine, zidovudine and tenofovir DF. The steady-state pharmacokinetics of efavirenz and tenofovir were unaffected when efavirenz and tenofovir DF were administered together versus each agent dosed alone. Clinically significant interactions would not be expected since the NRTIs are metabolized via a different route than efavirenz and would be unlikely to compete for the same metabolic enzymes and elimination pathways.

Table 9. Drug Interactions: Changes in Pharmacokinetic Parameters for Efavirenz in the Presence of the Coadministered Drug

Coadministered Drug	Dose of Coadministered Drug (mg)	Efavirenz Dose (mg)	N	Mean % Change of Efavirenz Pharmacokinetic Parameters ¹	
				C _{max}	AUC
Atazanavir ⁴	400 mg qd x 20 days	600 mg qd d 7-20	--	↔	↔
Indinavir	800 mg q8h x 14 days	200 mg x 14 days	11	↔	↔
Lopinavir/ Ritonavir	400/100 mg q12h x 9 days	600 mg x 9 days	11, 12 ²	↔	↓ 16
Nelfinavir	750 mg q8h x 7 days	600 mg x 7 days	10	↔	↔
Ritonavir	500 mg q12h x 8 days	600 mg x 10 days	9	↑ 14	↑ 21
Saquinavir SGC ³	1200 mg q8h x 10 days	600 mg x 10 days	13	↓ 13	↓ 12
Tenofovir disoproxil fumarate	300 mg qd	600 mg qd x 14 days	30	↔	↔
Clarithromycin	500 mg q12h x 7 days	400 mg x 7 days	12	↑ 11	↔
Azithromycin	600 mg single dose	400 mg x 7 days	14	↔	↔
Rifabutin	300 mg qd x 14 days	600 mg x 14 days	11	↔	↔
Rifampin	600 mg x 7 days	600 mg x 7 days	12	↓ 20	↓ 26

Coadministered Drug	Dose of Coadministered Drug (mg)	Efavirenz Dose (mg)	N	Mean % Change of Efavirenz Pharmacokinetic Parameters ¹	
				C _{max}	AUC
Antacid: Aluminium hydroxide 400 mg Magnesium hydroxide 400 mg + simethicone 30 mg	30 mL single dose	400 mg single dose	17	↔	↔
Carbamazepine	200 mg qd × 3 days, 200 mg BID × 3 days, then 400 mg qd × 15 days	600 mg × 35 days	14	↓ 21	↓ 36
Paroxetine	20 mg qd × 14 days	600 mg × 14 days	12	↔	↔
Sertraline	50 mg qd × 14 days	600 mg × 14 days	13	↑ 11	↔
Fluconazole	200 mg x 7 days	400 mg x 7 days	10	↔	↑ 16
Itraconazole	200 mg q 12h x 14 days	600 mg x 28 days	16	↔	↔
Voriconazole	400 mg po q12h × 1 day then 200 mg po q12h × 8 days	400 mg × 9 days	—	↑ 38	↑ 44
	300 mg po q12h days 2-7	300 mg × 7 days	—	↓ 14 ⁵	↔ ⁵
	400 mg po q12h days 2-7	300 mg × 7 days	—	↔ ⁵	↑ 17 ⁵
Diltiazem	240 mg x 14 days	600 mg x 28 days	12	↑ 16	↑ 11
Cetirizine	10 mg single dose	600 mg x 10 days	11	↔	↔
Famotidine	40 mg single dose	400 mg single dose	17	↔	↔
Atorvastatin	10 mg qd x 4 days	600 mg x 15 days	14	↔	↔
Pravastatin	40 mg qd x 4 days	600 mg x 15 days	11	↔	↔
Simvastatin	40 mg qd x 4 days	600 mg x 15 days	14	↓ 12	↔

1. Increase = ↑; Decrease = ↓; No Effect = ↔
2. Parallel-group design; N for efavirenz + lopinavir/ritonavir, N for efavirenz alone.
3. Soft Gelatin Capsule
4. Comparison with historical data suggests that Atazanavir does not affect Efavirenz PK
5. Relative to steady-state administration of efavirenz (600 mg once daily for 9 days).

Table 10. Drug Interactions: Changes in Pharmacokinetic Parameters for Coadministered Drug in the Presence of Efavirenz

Coadministered Drug	Dose of Coadministered Drug (mg)	Efavirenz Dose (mg)	N	Mean % Change of Coadministered Drug Pharmacokinetic Parameters ¹	
				C _{max}	AUC
Amprenavir	1200 mg q 12h × 7 days	600 mg × 7 days		↓ 33 ²	↓ 24 ²
Atazanavir	400 mg qd with a light meal d 1–20	600 mg qd with a light meal d 7–20	27	↓ 59	↓ 74
	400 mg qd d 1–6, then 300 mg qd d 7–20 with ritonavir 100 mg qd and a light meal	600 mg qd 2 h after atazanavir and ritonavir d 7–20	13	↑ 14 ³	↑ 39 ³
	300 mg qd/ritonavir 100 mg qd d 1-10 (pm), then 400 mg qd/ritonavir 100 mg qd d 11-24 (pm) (simultaneous with efavirenz)	600 mg qd d 11-24 (pm)	14	↑ 17%	↔
Indinavir	1000 mg q8h × 10 days	600 mg × 10 days	20		
	After morning dose			↔ ⁴	↓ 33 ⁴
	After afternoon dose			↔ ⁴	↓ 37 ⁴
	After evening dose			↓ 29 ⁴	↓ 46 ⁴
Indinavir/Ritonavir	Indinavir 800 mg + ritonavir 100 mg q12 hr d 1–29	600 mg d 15-29		↓ 17 ⁵	↓ 25 ⁵
Lopinavir/ Ritonavir	400/100 mg (capsule) q12h × 9 days	600 mg × 9 days	11, 7 ⁶	↔ ⁷	↓ 19 ⁷
	600/150 mg (tablet) q12h × 10 days with efavirenz compared to 400/100 mg q12h alone	600 mg × 9 days	23	↑ 36	↑ 36
Nelfinavir Metabolite AG-1402	750 mg q8h × 7 days	600 mg × 7 days	10	↑ 21 ↓ 40	↑ 20 ↓ 37
Ritonavir	500 mg q12h × 8 days	600 mg × 10 days	11		
	After AM dose			↑ 24	↑ 18
	After PM dose			↔	↔

Coadministered Drug	Dose of Coadministered Drug (mg)	Efavirenz Dose (mg)	N	Mean % Change of Coadministered Drug Pharmacokinetic Parameters ¹	
				C _{max}	AUC
Saquinavir SGC ⁹	1200 mg q8h × 10 days	600 mg × 10 days	12	↓ 50	↓ 62
Lamivudine	150 mg q 12 h x 14 days	600 mg × 14 days	9	↔	↔
Zidovudine	300 mg q 12 h x 14 days	600 mg × 14 days	9	↔	↔
Tenofovir disoproxil fumarate	300 mg qd	600 mg × 14 days	29	↔	↔
Clarithromycin 14-OH Metabolite	500 mg q12h × 7 days	400 mg × 7 days	11	↓ 26 ↑ 49	↓ 39 ↑ 34
Azithromycin	600 mg single dose	400 mg x 7 days	14	↑ 22	↔
Rifabutin 25-0- desacetyl rifabutin	300 mg qd × 14 days	600 mg × 14 days		↓ 32 ↓ 49 ²	↓ 38 ↓ 74 ²
Carbamazepine	200 mg qd × 3 days, 200 mg BID × 3 days, then 400 mg qd × 29 days	600 mg × 14 days	12	↓ 20	↓ 27
Epoxide Metabolite				↔	↔
Paroxetine	20 mg daily x 14 days	600 mg x 14 days	16	↔	↔
Oral contraceptive: Ethinyl estradiol/ Norgestimate Ethinyl estradiol Norelgestromin Levonorgestrel	0.035 mg/ 0.25 mg x 14 days	600 mg x 14 days	21 21 6	↔ ↓ 46% ↓ 80%	↔ ↓ 64% ↓ 83%
Methadone	Stable maintenance 35–100 mg daily	600 mg 14-21 days	11	↓ 45	↓ 52
Sertraline N desmethyl sertraline	50 mg qd × 14 days	600 mg × 14 days	13	↓ 29 ↓ 17	↓ 39 ↓ 20
Fluconazole	200 mg x 7 days	400 mg x 7 days	10	↔	↔
Itraconazole			18	↓ 37	↓ 39
Hydro-intraconazole	200 mg q12h x 28 days	600 mg x 14 days		↓ 35	↓ 37
Voriconazole	400 mg po q12h × 1 day then 200 mg po q12h x 8 days	400 mg × 9 days	—	↓ 61	↓ 77
	300 mg po q12h days 2-7	300 mg × 7 days	—	↓ 36 ⁸	↓ 55 ⁸

Coadministered Drug	Dose of Coadministered Drug (mg)	Efavirenz Dose (mg)	N	Mean % Change of Coadministered Drug Pharmacokinetic Parameters ¹	
				C _{max}	AUC
	400 mg po q12h days 2-7	300 mg × 7 days	—	↑23 ⁸	↔ ⁸
Lorazepam	2 mg single dose	600 mg × 10 days	12	↑ 16	↔
Diltiazem	240 mg x 21 days	600 mg x 14 days	13	↓60	↓69
Desacetyl diltiazem				↓64	↓75
N-monodesmethyl diltiazem				↓28	↓37
Cetirzine	10 mg single dose	600 mg × 10 days	11	↓24	↔
Atorvastatin	10 mg qd × 4 days	600 mg × 15 days	14	↓ 14	↓ 43
Total active (including metabolites)				↓ 15	↓ 32
Pravastatin	40 mg qd × 4 days	600 mg × 15 days	13	↓ 32	↓ 44
Simvastatin	40 mg qd × 4 days	600 mg × 15 days	14	↓ 72	↓ 68
Total active (including metabolites)				↓ 68	↓ 60

1. Increase = ↑; Decrease = ↓; No Effect = ↔.
2. Based on arithmetic means.
3. Compared with atazanavir 400 mg qd alone.
4. Comparator dose of indinavir was 800 mg q8h × 10 days.
5. Compared to indinavir 800 twice daily given with ritonavir 100 mg twice daily without efavirenz. The geometric C_{min} for indinavir (0.33 mg/L) when given with ritonavir and efavirenz was higher than the mean historical C_{min} (0.15 mg/L) when indinavir was given alone at 800 mg every 8 hours. When efavirenz 600 mg once daily was given with indinavir/ritonavir 800/100 mg twice daily in HIV-1-infected patients (n=6), the pharmacokinetics of indinavir and efavirenz were generally comparable to these data from uninfected volunteers.
6. Parallel-group design; N for efavirenz + lopinavir/ritonavir, N for lopinavir/ritonavir alone.
7. Values are for lopinavir. The pharmacokinetics of ritonavir 100 mg q12h are unaffected by concurrent efavirenz.
8. Relative to steady-state administration of voriconazole (400 mg for 1 day, then 200 mg po q12h for 2 days).
9. Soft Gelatin Capsule.

Emtricitabine and tenofovir DF:

The steady state pharmacokinetics of emtricitabine and tenofovir were unaffected when emtricitabine and tenofovir DF were administered together versus each agent dosed alone.

In vitro and clinical pharmacokinetic drug-drug interaction studies have shown that the potential for CYP450 mediated interactions involving emtricitabine and tenofovir with other medicinal products is low.

Emtricitabine and tenofovir are primarily excreted by the kidneys by a combination of glomerular filtration and active tubular secretion. No drug-drug interactions due to competition for renal excretion have been observed. Since emtricitabine and tenofovir are primarily eliminated by the kidneys, coadministration of ATRIPLA with drugs that reduce renal function (see **WARNINGS AND PRECAUTIONS: Renal**) or compete for active tubular secretion may increase serum concentrations of emtricitabine, tenofovir, and/or other renally eliminated drugs. Some examples of drugs that compete for active tubular secretion include, but are not limited to, cidofovir, acyclovir, valganciclovir, ganciclovir, and valganciclovir.

ATRIPLA should not be administered with HEPSERA (adefovir dipivoxil).

No clinically significant drug interactions have been observed between emtricitabine and famciclovir, indinavir, zidovudine, stavudine, and tenofovir DF. Similarly, no clinically significant drug interactions have been observed between tenofovir DF and abacavir, ribavirin, efavirenz, emtricitabine, entecavir, indinavir, lamivudine, lopinavir/ritonavir, methadone, oral contraceptives, nelfinavir or saquinavir/ritonavir in studies conducted in healthy volunteers.

Atazanavir and lopinavir/ritonavir have been shown to increase tenofovir concentrations (see Table 7 and Table 8). The mechanism of this interaction is unknown. The effects of coadministration of tenofovir on C_{max} , C_{min} and AUC are summarized in Table 11 (effect of other drugs on tenofovir) and Table 12 (effect of tenofovir on other drugs).

Table 11. Drug Interactions: Changes in Pharmacokinetic Parameters for Tenofovir in the Presence of the Coadministered Drug^{1,2}

Coadministered Drug	Dose of Coadministered Drug (mg)	N	Mean % Change of Tenofovir Pharmacokinetic Parameters ³ (90% CI)		
			C_{max}	AUC	C_{min}
Atazanavir ⁴	400 once daily × 14 days	33	↑ 14 (↑ 8 to ↑ 20)	↑ 24 (↑ 21 to ↑ 28)	↑ 22 (↑ 15 to ↑ 30)
Didanosine (enteric-coated)	400 once	25	↔	↔	↔
Didanosine (buffered)	250 or 400 once daily × 7 days	14	↔	↔	↔
Entecavir	1 mg once daily × 10 days	28	↔	↔	↔
Lopinavir/Ritonavir	400/100 twice daily × 14 days	24	↔	↑ 32 (↑ 25 to ↑ 38)	↑ 51 (↑ 37 to ↑ 66)

1. All interaction studies conducted in healthy volunteers.

2. Patients received tenofovir DF 300 mg once daily.

3. Increase = ↑; Decrease = ↓; No Effect = ↔

4. Reyataz Prescribing Information

Table 12. Drug Interactions: Changes in Pharmacokinetic Parameters for Coadministered Drug in the Presence of Tenofovir^{1,2}

Coadministered Drug	Dose of Coadministered Drug (mg)	N	Mean % Change of Coadministered Drug Pharmacokinetic Parameters ³ (90% CI)		
			C _{max}	AUC	C _{min}
Atazanavir ⁴	400 once daily × 14 days	34	↓ 21 (↓ 27 to ↓ 14)	↓ 25 (↓ 30 to ↓ 19)	↓ 40 (↓ 48 to ↓ 32)
	Atazanavir/ritonavir 300/100 once daily × 42 days	10	↓ 28 (↓ 50 to ↑ 5)	↓ 25 ⁵ (↓ 42 to ↓ 3)	↓ 23 ⁵ (↓ 46 to ↑ 10)
Entecavir	1 mg once daily × 10 days	28	↔	↑ 13 (↑ 11 to ↑ 15)	↔
Lopinavir	Lopinavir/ritonavir 400/100 twice daily × 14 days	24	↔	↔	↔
Ritonavir	Lopinavir/ritonavir 400/100 twice daily × 14 days	24	↔	↔	↔

1. All interaction studies conducted in healthy volunteers.
2. Patients received tenofovir DF 300 mg once daily.
3. Increase = ↑; Decrease = ↓; No Effect = ↔
4. Reyataz Prescribing Information
5. In HIV-infected patients, addition of tenofovir DF to atazanavir 300 mg plus ritonavir 100 mg, resulted in AUC and C_{min} values of atazanavir that were 2.3- and 4-fold higher than the respective values observed for atazanavir 400 mg when given alone.

Coadministration of tenofovir DF with didanosine results in changes in the pharmacokinetics of didanosine that may be of clinical significance. Table 13 summarizes the effects of tenofovir DF on the pharmacokinetics of didanosine. Concomitant dosing of tenofovir DF with didanosine buffered tablets or enteric-coated capsules significantly increased the C_{max} and AUC of didanosine. When didanosine 250 mg enteric-coated capsules were administered with tenofovir DF, systemic exposures of didanosine were similar to those seen with the 400 mg enteric-coated capsules alone under fasted conditions. The mechanism of this interaction is unknown. **Increases in didanosine concentrations of this magnitude could potentiate didanosine-associated adverse events, including pancreatitis, lactic acidosis, and neuropathy.** In addition, suppression of CD4 counts has been observed in patients receiving tenofovir DF with didanosine at a dose of 400 mg daily.

Therefore, the recommended dose of didanosine EC is 250 mg for HIV infected adults with body weight ≥60 kg and creatinine clearance ≥60 mL/min when coadministered with ATRIPLA. For patients with body weight <60 kg, and creatinine clearance ≥60 mL/min, the recommended dose of ddi-EC is 200 mg. Data are not available to recommend a dose adjustment of didanosine for patients with creatinine clearance <60mL/min (for didanosine EC dosing adjustment recommendations, see Table 8). **Coadministration of ATRIPLA and**

didanosine should be undertaken with caution and patients receiving this combination should be monitored closely for didanosine-associated adverse events.

Caution should be used when coadministering reduced-dose didanosine, tenofovir, and an NNRTI in treatment-naïve patients with high viral loads at baseline since such use has been associated with reports of a high rate of virologic failure and emergence of resistance at an early stage. All patients receiving tenofovir disoproxil fumarate and didanosine concomitantly should be closely monitored for didanosine-related adverse events and clinical response.

Table 13. Drug Interactions: Changes in Pharmacokinetic Parameters for Didanosine in the Presence of Tenofovir Disoproxil Fumarate^{1,2}

Didanosine Dose (mg)/Method of Administration ⁴	Tenofovir DF Method of Administration ^{2,4}	N	Mean % Change (90% CI) vs. Didanosine 400 mg Alone, Fasted ³	
			C _{max}	AUC
Buffered Tablets				
400 once daily ⁵ × 7 days	Fasted 1 hour after didanosine	14	↑ 28 (↑ 11 to ↑ 48)	↑ 44 (↑ 31 to ↑ 59)
Enteric coated Capsules				
400 Once, Fasted	With food, 2 hr after didanosine	26	↑ 48 (↑ 25 to ↑ 76)	↑ 48 (↑ 31 to ↑ 67)
400 Once, with Food	Simultaneously with didanosine	26	↑ 64 (↑ 41 to ↑ 89)	↑ 60 (↑ 44 to ↑ 79)
250 Once, Fasted	With food, 2 hr after didanosine	28	↓ 10 (↓ 22 to ↑ 3)	↔
250 Once, Fasted	Simultaneously with didanosine	28	↔	↑ 14 (0 to ↑ 31)
250 Once, with Food	Simultaneously with didanosine	28	↓ 29 (↓ 39 to ↓ 18)	↓ 11 (↓ 23 to ↑ 2)

1. All interaction studies conducted in healthy volunteers.
2. Patients received tenofovir DF 300 mg once daily.
3. Increase = ↑; Decrease = ↓; No Effect = ↔
4. Administration with food was with a light meal (~373 kcal, 20% fat).
5. Includes 4 subjects weighing <60 kg receiving ddi 250 mg.

Drug-Food Interactions

Interactions of ATRIPLA with food have not been established. In studies of the individual components of ATRIPLA, increased efavirenz and tenofovir concentrations were observed following administration with food (**see ACTION AND CLINICAL PHARMACOLOGY, Effect of Food on Oral Absorption**).

Drug-Herb Interactions

Interactions of ATRIPLA with herbs have not been established. St. John's Wort is expected to substantially decrease plasma levels of efavirenz but has not been studied in combination with efavirenz. ATRIPLA should not be co-administered with St. John's Wort.

Drug-Laboratory Interactions

Interactions of ATRIPLA with laboratory tests have not been established.

Efavirenz Assay Interference

Cannabinoid Test Interaction: Efavirenz does not bind to cannabinoid receptors. False-positive urine cannabinoid test results have been observed in non-HIV-infected volunteers receiving efavirenz when the Microgenics Cedia DAU Multi-Level THC assay was used for screening. Negative results were obtained when more specific confirmatory testing was performed with gas chromatography/mass spectrometry. For more information, please consult the SUSTIVA Product Monograph.

DOSAGE AND ADMINISTRATION

The dose of ATRIPLA is one tablet (containing 600 mg of efavirenz, 200 mg of emtricitabine and 300 mg of tenofovir disoproxil fumarate) once daily taken orally on an empty stomach. Dosing at bedtime may improve the tolerability of nervous system symptoms (see **WARNINGS AND PRECAUTIONS, Neurologic and INFORMATION FOR THE CONSUMER**).

Dose Adjustment for Renal Impairment

Because ATRIPLA is a fixed-dose combination, it should not be prescribed for patients requiring dosage adjustment such as those with moderate to severe renal impairment (creatinine clearance <50 mL/min).

Missed Dose

If a patient misses a dose at the regularly scheduled time, but then remembers it that same day, the patient should take the missed dose immediately. The patient should not take more than 1 dose of ATRIPLA in a day.

OVERDOSAGE

For management of a suspected drug overdose, please contact your regional Poison Control Centre.
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If overdose occurs the patient must be monitored for evidence of toxicity, and standard supportive treatment applied as necessary including monitoring of vital signs and observation

of the patient's clinical status. Administration of activated charcoal may be used to aid removal of efavirenz. Hemodialysis can remove both emtricitabine and tenofovir DF (refer to information below), but, since efavirenz is highly protein bound, dialysis is unlikely to significantly remove the drug from blood. There is no specific antidote for overdose with efavirenz.

Efavirenz:

Some patients accidentally taking 600 mg twice daily have reported increased nervous system symptoms. One patient experienced involuntary muscle contractions and a second patient experienced vomiting after taking twice the recommended dose.

Emtricitabine:

Limited clinical experience is available at doses higher than the therapeutic dose of EMTRIVA. In one clinical pharmacology study single doses of emtricitabine 1200 mg were administered to 11 patients. No severe adverse reactions were reported. The effects of higher doses are not known.

Hemodialysis treatment removes approximately 30% of the emtricitabine dose over a 3-hour dialysis period starting within 1.5 hours of emtricitabine dosing (blood flow rate of 400 mL/min and a dialysate flow rate of 600 mL/min), however, a single treatment does not significantly affect emtricitabine C_{max} or AUC. It is not known whether emtricitabine can be removed by peritoneal dialysis.

Tenofovir DF:

Limited clinical experience is available at doses higher than the therapeutic dose of tenofovir DF 300 mg. In one study, 600 mg tenofovir DF was administered to 8 patients orally for 28 days. No severe adverse reactions were reported. The effects of higher doses are not known.

Tenofovir is efficiently removed by hemodialysis with an extraction coefficient of approximately 54%. Following a single 300 mg dose of tenofovir DF, a four-hour hemodialysis session removed approximately 10% of the administered tenofovir dose.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

Efavirenz:

Efavirenz is a non-nucleoside reverse transcriptase (RT) inhibitor of human immunodeficiency virus type 1 (HIV-1). Efavirenz is predominantly a non-competitive inhibitor of HIV-1 RT. HIV-2 RT and human cellular DNA polymerases α , β , γ , and δ are not inhibited by efavirenz.

Emtricitabine:

Emtricitabine, a synthetic nucleoside analog of cytidine, is phosphorylated by cellular enzymes to form emtricitabine 5'-triphosphate. Emtricitabine 5'-triphosphate inhibits the activity of the HIV-1 RT by competing with the natural substrate deoxycytidine 5'-triphosphate and by being incorporated into nascent viral DNA which results in chain termination. Emtricitabine 5'-triphosphate is a weak inhibitor of mammalian DNA polymerases α , β , ϵ , and mitochondrial DNA polymerase γ .

Tenofovir DF:

Tenofovir DF is an acyclic nucleoside phosphonate diester analog of adenosine monophosphate. Tenofovir DF requires initial diester hydrolysis for conversion to tenofovir and subsequent phosphorylations by cellular enzymes to form tenofovir diphosphate. Tenofovir diphosphate inhibits the activity of HIV-1 RT by competing with the natural substrate deoxyadenosine 5'-triphosphate and, after incorporation into DNA, by DNA chain termination. Tenofovir diphosphate is a weak inhibitor of mammalian DNA polymerases α , β , and mitochondrial DNA polymerase γ .

Pharmacodynamics

Antiviral Activity In Vitro:

Efavirenz, emtricitabine and tenofovir DF: In combination studies evaluating the in vitro antiviral activity of efavirenz and emtricitabine together, efavirenz and tenofovir together and emtricitabine and tenofovir together, additive to synergistic antiviral effects were observed.

Efavirenz: The concentration of efavirenz inhibiting replication of wild-type laboratory adapted strains and clinical isolates in cell culture by 90–95% (EC_{90-95}) ranged from 1.7–25 nM in lymphoblastoid cell lines, peripheral blood mononuclear cells, and macrophage/monocyte cultures. Efavirenz demonstrated antiviral activity against most non-clade B isolates (subtypes A, AE, AG, C, D, F, G, J, and N), but had reduced antiviral activity against group O viruses.

Emtricitabine: The in vitro antiviral activity of emtricitabine against laboratory and clinical isolates of HIV was assessed in lymphoblastoid cell lines, the MAGI-CCR5 cell line, and peripheral blood mononuclear cells. The EC_{50} (50% effective concentration) values for emtricitabine were in the range of 0.0013–0.64 μ M (0.0003–0.158 μ g/mL). Emtricitabine displayed antiviral activity in vitro against HIV-1 clades A, B, C, D, E, F, and G (EC_{50} values

ranged from 0.007–0.075 μM) and showed strain specific activity against HIV-2 (EC_{50} values ranged from 0.007–1.5 μM).

Tenofovir DF: The antiviral activity in cell culture of tenofovir against laboratory and clinical isolates of HIV-1 was assessed in lymphoblastoid cell lines, primary monocyte/macrophage cells and peripheral blood lymphocytes. The EC_{50} values for tenofovir were in the range of 0.04–8.5 μM . Tenofovir displayed antiviral activity in vitro against HIV-1 clades A, B, C, D, E, F, G, and O (EC_{50} values ranged from 0.5–2.2 μM) and showed strain specific activity against HIV-2 (EC_{50} values ranged from 1.6 μM to 4.9 μM).

Pharmacokinetics

ATRIPLA:

One ATRIPLA Tablet was bioequivalent to one SUSTIVA tablet (600 mg) plus one EMTRIVA Capsule (200 mg) plus one VIREAD Tablet (300 mg) following single-dose administration to fasting healthy subjects (N=45).

Efavirenz:

In HIV-infected patients time-to-peak plasma concentrations were approximately 3–5 hours and steady-state plasma concentrations were reached in 6–10 days. In 35 patients receiving efavirenz 600 mg once daily, steady-state C_{max} was 12.9 μM , C_{min} was 5.6 μM , and AUC was 184 $\mu\text{M}/\text{h}$. Efavirenz is highly bound (approximately 99.5–99.75%) to human plasma proteins, predominantly albumin. Following administration of ^{14}C -labeled efavirenz, 14–34% of the dose was recovered in the urine (mostly as metabolites) and 16–61% was recovered in feces (mostly as parent drug).

In vitro studies suggest CYP3A4 and CYP2B6 are the major isozymes responsible for efavirenz metabolism. Efavirenz has been shown to induce P450 enzymes, resulting in induction of its own metabolism.

Efavirenz has been shown in vivo to cause hepatic enzyme induction, thus increasing the biotransformation of some drugs metabolized by CYP3A4. In vitro studies have demonstrated that efavirenz inhibits 2C9, 2C19 and 3A4 isozymes with K_i values (8.5–17 μM) in the range of observed efavirenz plasma concentrations. In in vitro studies, efavirenz did not inhibit CYP2E1 and inhibited CYP2D6 and CYP1A2 (K_i values 82–160 μM) only at concentrations well above those achieved clinically. Coadministration of efavirenz with drugs primarily metabolized by 2C9, 2C19, and 3A4 isozymes may result in altered plasma concentrations of the coadministered drug. Drugs which induce CYP3A4 activity would be expected to increase the clearance of efavirenz resulting in lowered plasma concentrations.

Efavirenz plasma exposure may be increased in patients with the homozygous G156T genetic variant of the CYP2B6 isozyme. The clinical implications of such an association are unknown; however, the potential for an increased frequency and severity of efavirenz-associated adverse events cannot be excluded.

Efavirenz has a terminal half-life of 52–76 hours after single doses and 40–55 hours after multiple doses.

Emtricitabine:

Following oral administration, emtricitabine is rapidly absorbed with peak plasma concentrations occurring at 1–2 hours post-dose. In vitro binding of emtricitabine to human plasma proteins is <4% and is independent of concentration over the range of 0.02–200 µg/mL. Following administration of radiolabelled emtricitabine, approximately 86% is recovered in the urine and 13% is recovered as metabolites. The metabolites of emtricitabine include 3'-sulfoxide diastereomers and their glucuronic acid conjugate. Emtricitabine is eliminated by a combination of glomerular filtration and active tubular secretion. Following a single oral dose, the plasma emtricitabine half-life is approximately 10 hours.

Tenofovir DF:

Following oral administration of tenofovir DF, maximum tenofovir serum concentrations are achieved in 1.0 ± 0.4 hour. In vitro binding of tenofovir to human plasma proteins is <0.7% and is independent of concentration over the range of 0.01–25 µg/mL. Approximately 70–80% of the intravenous dose of tenofovir is recovered as unchanged drug in the urine. Tenofovir is eliminated by a combination of glomerular filtration and active tubular secretion. Following a single oral dose of tenofovir DF, the terminal elimination half-life of tenofovir is approximately 17 hours.

Effect of Food on Oral Absorption

ATRIPLA has not been evaluated in the presence of food. Administration of efavirenz tablets with a high fat meal increased the mean AUC and C_{max} of efavirenz by 28% and 79%, respectively, compared to administration in the fasted state. Compared to fasted administration, dosing of tenofovir DF/ emtricitabine in combination with either a high fat meal or a light meal increased the mean AUC and C_{max} of tenofovir by 35% and 15%, respectively, without affecting emtricitabine exposures.

Special Populations and Conditions

Pediatrics and Geriatrics: Pharmacokinetics of tenofovir have not been evaluated in children (<18 years). Efavirenz has not been studied in children below 3 years of age or who weigh less than 13 kg. Emtricitabine has been studied in pediatric patients from 3 months to 17 years of age. ATRIPLA is not recommended for pediatric administration. Pharmacokinetics of efavirenz, emtricitabine, and tenofovir have not been fully evaluated in the elderly (>65 years).

Race:

Efavirenz: The pharmacokinetics of efavirenz in patients appear to be similar among the ethnic groups studied.

Emtricitabine: No pharmacokinetic differences due to race have been identified following the administration of emtricitabine.

Tenofovir DF: There were insufficient numbers from racial and ethnic groups other than Caucasian to adequately determine potential pharmacokinetic differences among these populations following administration of tenofovir DF.

Gender: Efavirenz, emtricitabine and tenofovir pharmacokinetics are similar in male and female patients.

Hepatic Insufficiency: The pharmacokinetics of tenofovir following a 300 mg single dose of tenofovir DF have been studied in non-HIV infected subjects with moderate to severe hepatic impairment. There were no substantial alterations in tenofovir pharmacokinetics in subjects with hepatic impairment compared with unimpaired subjects. The pharmacokinetics of efavirenz or emtricitabine have not been studied in patients with hepatic impairment; however, emtricitabine has not been shown to be significantly metabolized by liver enzymes, so the impact of liver impairment should be limited (**see WARNINGS AND PRECAUTIONS, Hepatic/Biliary/Pancreatic**).

Renal Insufficiency: The pharmacokinetics of efavirenz have not been studied in patients with renal insufficiency; however, less than 1% of efavirenz is excreted unchanged in the urine, so the impact of renal impairment on efavirenz elimination should be minimal. The pharmacokinetics of emtricitabine and tenofovir are altered in patients with renal insufficiency (**see WARNINGS AND PRECAUTIONS, Renal, Nephrotoxicity**). In patients with creatinine clearance <50 mL/min, C_{max} and $AUC_{0-\infty}$ of emtricitabine and tenofovir were increased. Because ATRIPLA is a fixed dose combination, it should not be prescribed for patients requiring dosage adjustment such as those with moderate to severe renal impairment (creatinine clearance <50 mL/min).

STORAGE AND STABILITY

Store at 15–30° C (59–86° F).

- Keep container tightly closed
- Dispense only in original container
- Do not use if seal over bottle opening is broken or missing.

SPECIAL HANDLING INSTRUCTIONS

There are no special handling instructions.

DOSAGE FORMS, COMPOSITION AND PACKAGING

ATRIPLA is available as tablets. Each tablet contains 600 mg of efavirenz, 200 mg of emtricitabine and 300 mg of tenofovir disoproxil fumarate (which is equivalent to 245 mg of

tenofovir disoproxil), as active ingredients. The tablets also include the following inactive ingredients: croscarmellose sodium, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, and sodium lauryl sulfate. The tablets are coated with Opadry II Pink 85F94172 which contains iron oxide black, iron oxide red, polyethylene glycol, polyvinyl alcohol, talc, and titanium dioxide. The tablets are pink, capsule-shaped and film-coated, debossed with “123” on one side and plain-faced on the other side. Each bottle contains 30 tablets and a silica gel desiccant and is closed with a child-resistant closure.

PART II. SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

ATRIPLA Tablets are fixed dose combination tablets containing efavirenz, emtricitabine and tenofovir disoproxil fumarate. SUSTIVA is the brand name for efavirenz, a non-nucleoside reverse transcriptase inhibitor. EMTRIVA is the brand name for emtricitabine, a synthetic nucleoside analog of cytidine. VIREAD is the brand name for tenofovir disoproxil fumarate (also known as tenofovir DF), which is converted in vivo to tenofovir, an acyclic nucleoside phosphonate (nucleotide) analog of adenosine 5'-monophosphate. All three compounds exhibit inhibitory activity against HIV-1 reverse transcriptase. VIREAD and EMTRIVA are the components of TRUVADA.

ATRIPLA Tablets are for oral administration. Each tablet contains 600 mg of efavirenz, 200 mg of emtricitabine and 300 mg of tenofovir disoproxil fumarate (which is equivalent to 245 mg of tenofovir disoproxil), as active ingredients. The tablets also include the following inactive ingredients: croscarmellose sodium, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, and sodium lauryl sulfate. The tablets are coated with Opadry II Pink 85F94172 which contains iron oxide black, iron oxide red, polyethylene glycol, polyvinyl alcohol, talc, and titanium dioxide.

Efavirenz:

Drug Substance

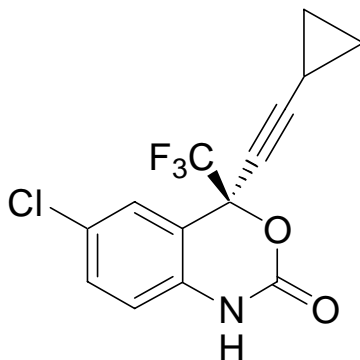
Common Name: efavirenz (USAN)

Chemical Name: (S)-6-chloro-4-(cyclopropylethynyl)-1,4-dihydro-4-(trifluoromethyl)-2*H*-3,1-benzoxazin-2-one.

Empirical Formula: C₁₄H₉ClF₃NO₂

Molecular Weight: 315.68

Structural Formula:



Physicochemical Properties:

Description: Efavirenz is a white to slightly pink crystalline powder.

Solubility: It is practically insoluble in water (<10 µg/mL). The pKa is 10.2. The partition coefficient (log P) is = 5.4.

Emtricitabine:

Drug Substance

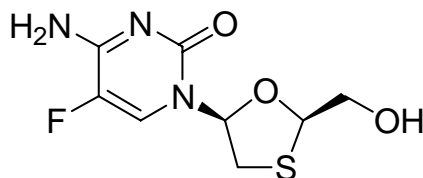
Common Name: emtricitabine (USAN)

Chemical Name: 5-fluoro-1-(2*R*,5*S*)-[2-(hydroxymethyl)-1,3-oxathiolan-5-yl]cytosine

Empirical Formula: C₈H₁₀FN₃O₃S

Molecular Weight: 247.24

Structural Formula:



Physicochemical Properties:

Description: Emtricitabine is a white to off-white crystalline powder.

Solubility: The solubility is approximately 112 mg/mL in water at 25 °C. The partition coefficient (log P) is -0.43 and the pKa is 2.65.

Tenofovir DF:

Drug Substance

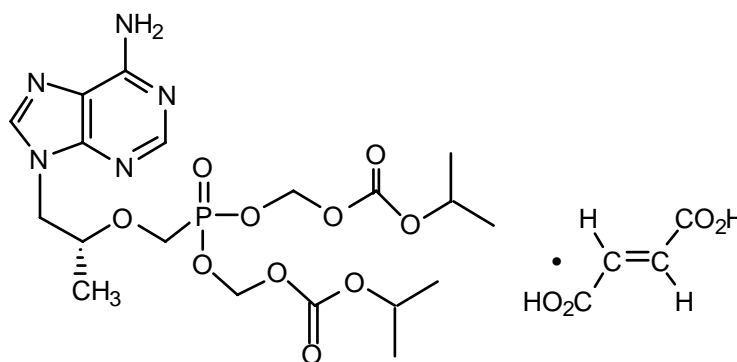
Common Name: tenofovir disoproxil fumarate (USAN)

Chemical Name: 9-[(R)-2-[[bis[[isopropoxycarbonyl]oxy]methoxy]phosphinyl]-methoxy]propyl]adenine fumarate (1:1)

Empirical Formula: C₁₉H₃₀N₅O₁₀P • C₄H₄O₄

Molecular Weight: 635.52

Structural Formula:



Physicochemical Properties:

Description: Tenofovir disoproxil fumarate is a white to off-white crystalline powder.

Solubility: The solubility is 13.4 mg/mL in water at 25 °C. The partition coefficient (log P) is 1.25 and the pKa is 3.75.

CLINICAL TRIALS

Study Demographics and Trial Design

Description of Clinical Studies

For safety and efficacy studies using SUSTIVA, EMTRIVA or VIREAD in combination with other antiretroviral agents, also consult the Product Monographs for these products.

Study 934

Study 934 was a phase III 144-week open-label randomized non-inferiority clinical study comparing the safety and efficacy of a once-daily regimen containing the individual agents efavirenz, emtricitabine and tenofovir disoproxil fumarate versus the fixed-dose combination of lamivudine and zidovudine (Combivir[®]) administered twice-daily and efavirenz once daily

in HIV-1 infected antiretroviral treatment-naive patients. Patients in both arms of Study 934 who completed 144 weeks were given the option to roll-over into an open-label extension phase of the study and switch their ARV regimen to the once-daily fixed-dose combination single tablet regimen, efavirenz/emtricitabine/tenofovir DF (ATRIPLA) for an additional 96 weeks.

In this study, 59% of patients were Caucasian and 23% were Black. The mean baseline CD4 cell count was 245 cells/mm³ (range 2–1191) and median baseline plasma HIV-1 RNA was 5.01 log₁₀ copies/mL (range 3.56–6.54). Patients were stratified by baseline CD4 count (< or ≥200 cells/mm³); 41% had CD4 cell counts <200 cells/mm³ and 51% of patients had baseline viral loads >100,000 copies/mL.

Table 14. Study 934 Efavirenz + Emtricitabine + Tenofovir DF Compared with Efavirenz + Lamivudine/Zidovudine

Study Number	Trial Design	Dosage, Route of Administration and Duration	Study Subjects	Mean Age	Gender
GS-01-934	Randomized, open-label, parallel, multicenter, active controlled study. Arm 1: emtricitabine+ tenofovir DF+ efavirenz Arm 2: lamivudine/ zidovudine + efavirenz	Arm 1¹: efavirenz 600 mg once daily, emtricitabine 200 mg once and tenofovir DF 300 mg once daily; Oral administration. Arm 2: efavirenz 600 mg once daily and Combivir (lamivudine/ zidovudine) 150/300 mg twice daily. Oral administration. 144 weeks	Antiretroviral naive patients (HIV-1 RNA >10,000 copies/mL) (N=517 randomized)	Mean 38 years (18–80)	Male : 86% Female: 14%

¹From weeks 96 to 144 of the study, patients randomized to emtricitabine + tenofovir DF received emtricitabine/tenofovir DF fixed-dose combination (Truvada) with efavirenz in place of emtricitabine + tenofovir DF with efavirenz.

Study 073

Study 073 was a 48-week open-label, randomized clinical study in patients with stable virologic suppression on combination antiretroviral therapy. The study compared the efficacy of ATRIPLA to antiretroviral therapy consisting of at least two nucleoside reverse transcriptase inhibitors (NRTIs) administered in combination with a protease inhibitor (with or without ritonavir) or a non-nucleoside reverse transcriptase inhibitor. At baseline, patients had been virologically suppressed (HIV-1 RNA <200 copies/mL) on their current antiretroviral therapy for at least 12 weeks prior to study entry, and had no known HIV-1

substitutions conferring resistance to the components of ATRIPLA or history of virologic failure. At baseline, 140 subjects (47%) were receiving an NNRTI-based regimen and 160 subjects (53%) were receiving a PI-based regimen.

Table 15. Study 073 ATRIPLA Compared with 2 NRTI + NNRTI or 2 NRTI + PI (+/- Ritonavir)

Study Number	Trial Design	Dosage, Route of Administration and Duration	Study Subjects	Mean Age	Gender
AI266073	Open-label, randomized, parallel multicenter, active-controlled study.	Arm 1: ATRIPLA once daily. Oral administration. 48 weeks Arm 2: Standard Baseline Regimen (SBR)* either 2 NRTI + NNRTI or PI (+/- ritonavir). Oral administration. 48 weeks	Virologically suppressed (≥ 3 months) HIV-1 infected patients (HIV-1 RNA <200 copies/mL) (N=306 randomized)	Mean 43 years (22–73)	Male : 88% Female: 12%

*Dosage and dosing frequency administered according to manufacturers' dosing instructions.

Patients were randomized in a 2:1 ratio to switch to ATRIPLA (N=203) or stay on their baseline regimen (SBR) (N=97). Patients had a mean age of 43 years (range 22 to 73 years), 88% were male, 68% were white, 29% were black or African-American, and 3% were of other races. At baseline, median CD4⁺ cell count was 516 cells/mm³ and all but 11 patients (3.7%) had HIV-1 RNA <50 copies/mL. The median time since onset of antiretroviral therapy was 3 years.

In antiretroviral treatment-experienced patients, the use of ATRIPLA Tablets may be considered for patients with HIV strains that are expected to be susceptible to the components of ATRIPLA as assessed by treatment history or by genotypic or phenotypic testing (see **VIROLOGY, Resistance and Cross Resistance**).

Study Result

Efavirenz, Emtricitabine, and Tenofovir DF

Study 934: Efavirenz + Emtricitabine + Tenofovir DF Compared with Efavirenz + Lamivudine/Zidovudine

Treatment outcomes through 48 and 144 weeks for those patients who did not have efavirenz resistance at baseline are presented in Table 16.

Table 16. Outcomes of Randomized Treatment at Weeks 48 and 144 (Study 934)

Outcomes	At Week 48		At Week 144 ¹	
	FTC+TDF + EFV (N=244)	3TC/AZT +EFV (N=243)	FTC+TDF + EFV (N=227)	3TC/AZT +EFV (N=229)
Responder ²	84%	73%	71%	58%
Virologic Failure ³	2%	4%	3%	6%
Rebound	1%	3%	2%	5%
Never Suppressed	0%	0%	0%	0%
Change in Antiretroviral Regimen	1%	1%	1%	1%
Death	<1%	1%	1%	1%
Discontinued Due to Adverse Event	4%	9%	5%	12%
Discontinued for Other Reasons ⁴	10%	14%	20%	22%

1. Patients who did not consent to continue study beyond Week 48 were excluded from analysis.
2. Patients achieved and maintained confirmed HIV-1 RNA <400 copies/mL through Week 48 and 144.
3. Includes confirmed viral rebound and failure to achieve confirmed <400 copies/mL through Week 48 and 144.
4. Includes lost to follow-up, patient withdrawal, non-compliance, protocol violation and other reasons.

In this study, efavirenz +emtricitabine + tenofovir DF demonstrated statistically significant superiority to lamivudine/zidovudine in combination with efavirenz in achieving and maintaining HIV-1 RNA <400 copies/mL through 48 and 144 weeks (Table 16). The difference in the percentages of responders, stratified by baseline CD4 cell count (< or ≥200 cells/mm³), between the efavirenz +emtricitabine + tenofovir DF group and the efavirenz + lamivudine/zidovudine group was 11.4%, and the 95% CI was 4.3% to 18.6% (p=0.002) at Week 48 and was 13% at Week 144, 95% CI= 4% to 22% (p=0.004). Through 48 weeks of therapy, 80% and 70% of patients in the efavirenz +emtricitabine + tenofovir DF and the efavirenz + lamivudine/zidovudine arms, respectively, achieved and maintained HIV-1 RNA <50 copies/mL (64 and 56%, respectively, through Week 144). The difference in the percentages of responders stratified by baseline CD4 cell count (< or ≥ 200 cells/mm³) between the efavirenz + emtricitabine + tenofovir DF group and the efavirenz + lamivudine/zidovudine group was 9.1%, and the 95% CI was 1.6% to 16.6% (p=0.02) at Week 48 and was 8% at Week 144, 95% CI = -1% to 17% (p=0.082). The mean increase from baseline in CD4 cell count was 190 cells/mm³ for the efavirenz +emtricitabine + tenofovir DF arm, and 158 cells/mm³ for the efavirenz + lamivudine/zidovudine arm (p=0.002) at Week 48 (312 and 271 cells/mm³, respectively, at Week 144, p=0.089).

The difference in the proportion of patients who achieved and maintained HIV-1 RNA <400 copies/mL through 48 weeks largely results from the higher number of discontinuations due to adverse events and other reasons in the zidovudine/lamivudine group in this open label study.

Study 073: ATRIPLA Compared with 2 NRTI + NNRTI or 2 NRTI + PI (+/- Ritonavir)

Table 17 summarizes treatment outcomes through week 48.

Table 17 Outcomes of Randomized Treatment at Week 48 (Study 073)

Outcomes	ATRIPLA (N=203)	Stayed on Baseline Regimen (SBR) (N=97)
Responder ^a	89% (87%)	88% (85%)
Virologic failure ^b	1% (2%)	1% (4%)
Death	0%	0%
Discontinued due to adverse event	5%	1%
Discontinued for other reasons ^c	5%	10%

- a. Patients who maintained confirmed HIV-1 RNA <200 copies/mL (<50 copies/mL) through Week 48. Primary endpoint of the study was HIV-1 RNA <200 copies/mL
- b. Virologic failure defined as rebound [two consecutive HIV-1 RNA \geq 200 copies/mL (\geq 50 copies/mL) or last observed HIV RNA \geq 200 copies/mL (\geq 50 copies/mL)].
- c. Includes lost to follow-up, patient consent withdrawal, investigator's discretion, and protocol violation.

The responder difference (HIV-1 RNA <200 copies/mL), ATRIPLA minus SBR, was 1% (95% CI: -7%, 9%, p=0.82) at week 48. ATRIPLA was non-inferior to SBR in this study. At week 48, the median change from baseline in CD4+ cell count was 3 cells/mm³ in the ATRIPLA group and 9 cells/mm³ in the SBR group.

When treatment responses were evaluated at 48 weeks by baseline treatment regimen strata (NNRTI or PI), 92% vs. 84% of patients on prior NNRTI-based antiretroviral therapy maintained HIV-1 RNA <200 copies/mL when randomized to switch to ATRIPLA vs. continue SBR, respectively. For patients on prior PI-based antiretroviral therapy, 87% vs. 90% of patients maintained HIV-1 RNA <200 copies/mL when randomized to receive ATRIPLA vs. SBR, respectively.

Comparative Bioavailability Studies

Study GS-US-177-0105 was a single-dose, randomized, open-label, two-way crossover study conducted in 45 healthy male and female adults to establish comparative bioavailability between the combination tablet (containing 600 mg efavirenz/200 mg emtricitabine/300 mg tenofovir disoproxil fumarate) and concurrent administration of the individual dosage forms, the SUSTIVA (efavirenz) 600 mg tablet, the EMTRIVA (emtricitabine) 200 mg capsule and the VIREAD (tenofovir disoproxil fumarate) 300 mg tablet, administered under fasting conditions, by evaluation of C_{max} and AUC of efavirenz, emtricitabine and tenofovir. The bioavailability of one tablet of the fixed dose combination containing 600 mg efavirenz/200 mg emtricitabine/300 mg tenofovir disoproxil fumarate was found to be comparable to the bioavailability of one dose of the three individual dosage forms taken in combination. The results of Study GS-US-177-0105 are summarized in Tables 18, 19 and 20 below.

Table 18. Summary of Efavirenz Pharmacokinetic Parameters (GS-US-177-0105) – Fasted

Efavirenz 600mg From Measured Data				
Geometric Mean				
Arithmetic Mean (CV%)				
Parameter	Test Treatment B²	Reference Treatment A¹	% Ratio of Geometric Means⁵	90% Confidence Interval⁵
AUC ₀₋₇₂ (ng•h/mL)	52098.79 53366.2 (22.2)	54570.04 56352.5 (22.9)	95.47	(90.69, 100.51)
AUC ₁ (ng•h/mL)	139109.1 146074.9 (33.1)	145097.8 155518.6 (34.6)	95.87	(89.63, 102.55)
C _{max} (ng/mL)	2179.16 2264.3 (26.8)	2205.86 2308.6 (30.3)	98.79	(92.28, 105.76)
T _{max} ³ (h)	3.50 (2.00–8.00)	3.50 (1.50–5.50)		
T _{1/2} ⁴ (h)	180.6 (45.3)	182.5 (38.3)		

1. Treatment A = concurrent administration of SUSTIVA (efavirenz) 600 mg tablet , EMTRIVA (emtricitabine) 200 mg capsule and VIREAD (tenofovir DF) 300 mg tablet to fasted subjects.
2. Treatment B = 600 mg efavirenz/ 200 mg emtricitabine/300 mg tenofovir DF combination tablet administered to fasted subjects.
3. Expressed as the median (range) only.
4. Expressed as the arithmetic mean (CV%) only.
5. Based on geometric least squares mean.

Table 19. Summary of Emtricitabine Pharmacokinetic Parameters (GS-US-177-0105) – Fasted

Emtricitabine 200mg From Measured Data				
Geometric Mean				
Arithmetic Mean (CV%)				
Parameter	Test Treatment B²	Reference Treatment A¹	% Ratio of Geometric Means⁵	90% Confidence Interval⁵
AUC _T (ng•h/mL)	10523.83 10682.6 (18.1)	10740.78 10874.4 (14.9)	97.98	94.90–101.16
AUC ₁ (ng•h/mL)	10694.43 10854.9 (17.9)	10916.98 11054.3 (14.9)	97.96	94.86–101.16
C _{max} (ng/mL)	2066.48 2130.6 (25.3)	2325.96 2384.4 (20.4)	88.84	84.02–93.94
T _{max} ³ (h)	1.50 (1.00–5.00)	1.50 (0.77–2.50)		
T _{1/2} ⁴ (h)	14.5 (53.8)	14.6 (47.8)		

1. Treatment A = concurrent administration of SUSTIVA (efavirenz) 600 mg tablet , EMTRIVA (emtricitabine) 200 mg capsule and VIREAD (tenofovir DF) 300 mg tablet to fasted subjects.
2. Treatment B = 600 mg efavirenz/ 200 mg emtricitabine/300 mg tenofovir DF combination tablet administered to fasted subjects.
3. Expressed as the median (range) only.
4. Expressed as the arithmetic mean (CV%) only.
5. Based on geometric least squares mean.

Table 20. Summary of Tenofovir Pharmacokinetic Parameters (GS-US-177-0105) – Fasted

Tenofovir 300mg From Measured Data				
Geometric Mean Arithmetic Mean (CV%)				
Parameter	Test Treatment B ²	Reference Treatment A ¹	% Ratio of Geometric Means ⁵	90% Confidence Interval ⁵
AUC _T (ng•h/mL)	1845.03 1948.8 (32.9)	1858.15 1969.0 (32.8)	99.29	91.02–108.32
AUC _I (ng•h/mL)	2218.24 2314.0 (29.2)	2208.41 2319.4 (30.3)	100.45	93.22–108.23
C _{max} (ng/mL)	307.25 325.1 (34.2)	335.93 352.9 (29.6)	91.46	84.64–98.83
T _{max} ³ (h)	1.00 (0.50–3.50)	0.75 (0.50–2.00)		
T _½ ⁴ (h)	18.9 (20.8)	17.8 (22.6)		

1. Treatment A=concurrent administration of SUSTIVA (efavirenz) 600 mg tablet , EMTRIVA (emtricitabine) 200 mg capsule and VIREAD (tenofovir DF) 300 mg tablet to fasted subjects
2. Treatment B=600 mg efavirenz/ 200 mg emtricitabine/300 mg tenofovir DF combination tablet administered to fasted subjects.
3. Expressed as the median (range) only.
4. Expressed as the arithmetic mean (CV%) only.
5. Based on geometric least squares mean.

DETAILED PHARMACOLOGY

VIROLOGY (MICROBIOLOGY)

Efavirenz:

Efavirenz is a non-nucleoside reverse transcriptase (RT) inhibitor of human immunodeficiency virus type 1 (HIV-1). Efavirenz is predominantly a non-competitive inhibitor of HIV-1 RT. HIV-2 RT and human cellular DNA polymerases α , β , γ , and δ are not inhibited by efavirenz.

Emtricitabine:

Emtricitabine, a synthetic nucleoside analog of cytidine, is phosphorylated by cellular enzymes to form emtricitabine 5'-triphosphate. Emtricitabine 5'-triphosphate inhibits the activity of the HIV-1 RT by competing with the natural substrate deoxycytidine 5'-triphosphate and by being incorporated into nascent viral DNA which results in chain termination. Emtricitabine 5'-triphosphate is a weak inhibitor of mammalian DNA polymerases α , β , ϵ , and mitochondrial DNA polymerase γ .

Tenofovir DF:

Tenofovir DF is an acyclic nucleoside phosphonate diester analog of adenosine monophosphate. Tenofovir DF requires initial diester hydrolysis for conversion to tenofovir and subsequent phosphorylations by cellular enzymes to form tenofovir diphosphate. Tenofovir diphosphate inhibits the activity of HIV-1 RT by competing with the natural substrate deoxyadenosine 5'-triphosphate and, after incorporation into DNA, by DNA chain termination. Tenofovir diphosphate is a weak inhibitor of mammalian DNA polymerases α , β , and mitochondrial DNA polymerase γ .

Antiviral Activity

Efavirenz, emtricitabine and tenofovir DF:

In combination studies evaluating the in vitro antiviral activity of emtricitabine and efavirenz together, efavirenz and tenofovir together and emtricitabine and tenofovir together, additive to synergistic antiviral effects were observed.

Efavirenz:

The clinical significance of in vitro susceptibility of HIV-1 to efavirenz has not been established. The in vitro antiviral activity of efavirenz was assessed in lymphoblastoid cell lines, peripheral blood mononuclear cells (PBMCs) and macrophage/monocyte cultures enriched from PBMCs. The concentration of efavirenz inhibiting replication of wild-type laboratory adapted strains and clinical isolates in cell culture by 90–95% (EC_{90-95}) ranged from 1.7 to ≤ 25 nM. Efavirenz demonstrated additive to synergistic antiviral activity against HIV-1 in cell culture when combined with non-nucleoside reverse transcriptase inhibitors (NNRTIs) (delavirdine and nevirapine), nucleoside reverse transcriptase inhibitors (NRTIs) (abacavir, didanosine, lamivudine, stavudine, zalcitabine, and zidovudine), protease inhibitors (PIs) (amprenavir, indinavir, lopinavir, nelfinavir, ritonavir, and saquinavir), and the fusion inhibitor enfuvirtide. Efavirenz demonstrated antiviral activity against most non-clade B isolates (subtypes A, AE, AG, C, D, F, G, J, and N), but had reduced antiviral activity against group O viruses.

Emtricitabine:

The in vitro antiviral activity of emtricitabine against laboratory and clinical isolates of HIV was assessed in lymphoblastoid cell lines, the MAGI-CCR5 cell line, and peripheral blood mononuclear cells. The EC_{50} values for emtricitabine were in the range of 0.0013–0.64 μ M (0.0003–0.158 μ g/mL). In drug combination studies of emtricitabine with nucleoside reverse transcriptase inhibitors (abacavir, lamivudine, stavudine, zalcitabine, and zidovudine), non-nucleoside reverse transcriptase inhibitors (delavirdine, efavirenz, and nevirapine), and protease inhibitors (amprenavir, nelfinavir, ritonavir, and saquinavir), additive to synergistic effects were observed. Most of these drug combinations have not been studied in humans. Emtricitabine displayed antiviral activity in vitro against HIV-1 clades A, B, C, D, E, F, and G (EC_{50} values ranged from 0.007–0.075 μ M) and showed strain specific activity against HIV-2 (EC_{50} values ranged from 0.007–1.5 μ M).

Tenofovir DF:

The in vitro antiviral activity of tenofovir against laboratory and clinical isolates of HIV-1 was assessed in lymphoblastoid cell lines, primary monocyte/macrophage cells and peripheral blood lymphocytes. The EC₅₀ values for tenofovir were in the range of 0.04–8.5 µM. In drug combination studies of tenofovir with nucleoside reverse transcriptase inhibitors (abacavir, didanosine, lamivudine, stavudine, zalcitabine, and zidovudine), non-nucleoside reverse transcriptase inhibitors (delavirdine, efavirenz, and nevirapine), and protease inhibitors (amprenavir, indinavir, nelfinavir, ritonavir, and saquinavir), additive to synergistic effects were observed. Most of these drug combinations have not been studied in humans. Tenofovir displayed antiviral activity in vitro against HIV-1 clades A, B, C, D, E, F, G, and O (IC₅₀ values ranged from 0.5–2.2 µM).

Resistance

Efavirenz, emtricitabine and tenofovir DF:

HIV-1 isolates with reduced susceptibility to the combination of emtricitabine and tenofovir have been selected in cell culture and in clinical studies. Genotypic analysis of these isolates identified the M184V/I and/or K65R amino acid substitutions in the viral RT.

In a clinical study of treatment-naïve patients (Study 934, see Clinical Studies) resistance analysis was performed on HIV isolates from all virologic failure patients with >400 copies/mL of HIV-1 RNA at Week 144 or early discontinuations. The resistance analysis population consisted of 19/244 (8%) patients in the emtricitabine + tenofovir DF group and 29/243 (12%) patients in the zidovudine/lamivudine fixed-dose combination group with available genotypic data. Genotypic resistance to efavirenz, predominantly the K103N mutation, was the most common form of resistance that developed. Resistance to efavirenz occurred in 13/19 (68%) analyzed patients (13/244, 5% of total patients) in the emtricitabine + tenofovir DF group and in 21/29 (72%) analyzed patients (21/243, 9% of total patients) in the zidovudine/lamivudine fixed-dose combination group. The M184V amino acid substitution, associated with resistance to emtricitabine and lamivudine, was observed in 2/19 (11%) analyzed patient isolates (2/244, 0.8% of total patients) in the emtricitabine + tenofovir DF group and in 10/29 (34%) analyzed patient isolates (10/243, 4.1% of total patients) in the zidovudine/lamivudine group; this difference was statistically significant (p=0.021). Through 144 weeks of Study 934, no patients developed a detectable K65R mutation in their HIV as analyzed through standard genotypic analysis.

Only limited resistance data are available from a clinical study of treatment-experienced patients with stable virologic suppression and no history of virologic failure (Study 073, see CLINICAL TRIALS section) since virologic failure was observed in only 3 subjects in the ATRIPLA treatment group. One of the 3 subjects who experienced virologic failure on ATRIPLA had confirmed development of resistance, specifically the K103N efavirenz resistance mutation only.

In a clinical study of treatment-naïve patients, isolates from 8/47 (17%) analyzed patients receiving tenofovir DF developed the K65R substitution through 144 weeks of therapy; 7 of these occurred in the first 48 weeks of treatment and one at Week 96. In treatment

experienced patients, 14/304 (5%) of tenofovir DF treated patients with virologic failure through Week 96 showed >1.4 fold (median 2.7) reduced susceptibility to tenofovir. Genotypic analysis of the resistant isolates showed a mutation in the HIV-1 RT gene resulting in the K65R amino acid substitution.

Efavirenz:

HIV-1 isolates with reduced susceptibility to efavirenz (>380-fold increase in EC₉₀) compared to baseline emerged rapidly under selection in cell culture in the presence of drug.

Genotypic characterization of these viruses identified mutations resulting in single amino acid substitutions L100I or V179D, double substitutions L100I/V108I and triple substitutions L100I/V179D/Y181C in RT. Other resistance mutations observed to emerge commonly included L100I (7%), K101E/Q/R(14%), V108I (11%), G190S/T/A(7%), P225H(18%), and M230I/L(11%).

Phenotypic (N=26) changes in evaluable HIV-1 isolates and genotypic (N=104) changes in plasma virus from selected patients treated with efavirenz in combination with indinavir, or with zidovudine plus lamivudine, were monitored. Clinical isolates with reduced susceptibility in vitro to efavirenz have been obtained. One or more RT mutations at amino acid positions 98, 100, 101, 103, 106, 108, 188, 190 and 225, and 227 were observed in all 102 of 104 patients with a frequency of at least 9% compared to baseline. The mutation at RT amino acid position 103 (lysine to asparagine) was the most frequently observed (≥90%). A mean loss in susceptibility (EC₉₀) to efavirenz of 47-fold was observed in 26 clinical isolates. Five clinical isolates were evaluated for both genotypic and phenotypic changes from baseline. Decreases in efavirenz susceptibility (range from 9 to >312-fold increase in EC₉₀) were observed for these isolates in vitro compared to baseline. All 5 isolates possessed at least one of the efavirenz-associated RT mutations. The clinical relevance of phenotypic and genotypic changes associated with efavirenz therapy has not been established.

Emtricitabine:

Emtricitabine-resistant isolates of HIV have been selected in vitro. Genotypic analysis of these isolates showed that the reduced susceptibility to emtricitabine was associated with a mutation in the HIV RT gene at codon 184 which resulted in an amino acid substitution of methionine by valine or isoleucine (M184V/I).

Tenofovir DF:

HIV-1 isolates with reduced susceptibility to tenofovir have been selected in vitro. These viruses expressed a K65R mutation in RT and showed a 2–4 fold reduction in susceptibility to tenofovir.

Cross-resistance

Efavirenz, emtricitabine and tenofovir DF:

Cross-resistance has been recognized among NNRTIs. Cross-resistance has also been recognized among certain NRTIs. The M184V/I and/or K65R substitutions selected in vitro by the combination of emtricitabine and tenofovir are also observed in some HIV-1 isolates

from subjects failing treatment with tenofovir in combination with either lamivudine or emtricitabine, and either abacavir or didanosine. Therefore, cross-resistance among these drugs may occur in patients whose virus harbors either or both of these amino acid substitutions. Delaviradine and/or nevirapine resistant clinical viral isolates with NNRTI resistance-associated substitutions (A98G, L100I, K101E/P, K103N/S, V106A, Y181X, Y188X, G190X, P225H, F227L, or M230L) showed reduced susceptibility to efavirenz in cell culture.

Efavirenz:

Rapid emergence of HIV-1 strains that are cross-resistant to non-nucleoside RT inhibitors has been observed in vitro. Thirteen clinical isolates previously characterized as efavirenz-resistant were also phenotypically resistant to nevirapine and delavirdine in vitro compared to baseline. Clinically derived zidovudine-resistant HIV-1 isolates tested in vitro retained susceptibility to efavirenz. Cross-resistance between efavirenz and HIV protease inhibitors is unlikely because of the different enzyme targets involved.

Emtricitabine:

Emtricitabine-resistant isolates (M184V/I) were cross-resistant to lamivudine and zalcitabine but retained susceptibility in vitro to didanosine, stavudine, tenofovir, zidovudine, and NNRTIs (delavirdine, efavirenz, and nevirapine). Isolates from heavily treatment-experienced patients containing the M184V/I amino acid substitution in the context of other NRTI resistance-associated substitutions may retain susceptibility to tenofovir. HIV-1 isolates containing the K65R substitution, selected in vivo by abacavir, didanosine, tenofovir, and zalcitabine, demonstrated reduced susceptibility to inhibition by emtricitabine. Viruses harboring mutations conferring reduced susceptibility to stavudine and zidovudine (M41L, D67N, K70R, L210W, T215Y/F, K219Q/E) or didanosine (L74V) remained sensitive to emtricitabine. HIV-1 containing the K103N substitution associated with resistance to NNRTIs was susceptible to emtricitabine.

Tenofovir DF:

The K65R mutation selected by tenofovir is also selected in some HIV-1 infected patients treated with abacavir, didanosine, or zalcitabine. HIV-1 isolates with the K65R mutation also showed reduced susceptibility to emtricitabine and lamivudine. Therefore, cross-resistance among these drugs may occur in patients whose virus harbors the K65R mutation. HIV-1 isolates from patients (N=20) whose HIV-1 expressed a mean of 3 zidovudine-associated RT amino acid substitutions (M41L, D67N, K70R, L210W, T215Y/F, or K219Q/E/N) showed a 3.1-fold decrease in the susceptibility to tenofovir. Multinucleoside resistant HIV-1 with a T69S double insertion mutation in the RT showed reduced susceptibility to tenofovir.

TOXICOLOGY

For additional information on toxicology, reproductive toxicology, mutagenicity and carcinogenicity, please consult the Product Monographs for SUSTIVA, EMTRIVA, and VIREAD.

Carcinogenesis

Efavirenz:

Oral carcinogenicity studies in mice and rats were carried out with efavirenz. Mice were dosed with 0, 25, 75, 150, or 300 mg/kg/day for 2 years. Incidences of hepatocellular adenomas and carcinomas and pulmonary alveolar/bronchiolar adenomas at all doses were increased above background in females. No increases in tumor incidence above background were seen in males. The systemic exposure (based on AUCs) in mice was approximately 1.7-fold that in humans receiving the 600-mg/day dose. In studies in which rats were administered efavirenz at doses of 0, 25, 50, or 100 mg/kg/day for 2 years, no increases in tumor incidence above background were observed. The exposure to efavirenz in rats at all doses was lower than that in humans.

The findings from the efavirenz mouse carcinogenicity study may not represent a significant risk to patients based upon the following: An increase in the incidence of hepatic tumors in efavirenz-treated mice was not unexpected as efavirenz is known to induce hepatic drug-metabolizing enzyme activity and enzyme inducers are known to increase the incidence of hepatic tumors in rodents, but not in humans. While the cause of the increased incidence of pulmonary tumors is not known, this finding also may not constitute a significant risk for patients given efavirenz because: (1) efavirenz is not genotoxic, (2) the strain of mice used in these studies is documented to have a high spontaneous background incidence of this tumor type, and (3) a decrease in the incidence of pulmonary tumors was observed in efavirenz-treated male mice. In male mice, plasma efavirenz concentrations were equal to or greater than in female mice.

Emtricitabine:

In long-term oral carcinogenicity studies of emtricitabine, no drug-related increase in tumor incidence was found in mice at doses up to 750 mg/kg/day (26 times the human systemic exposure at the therapeutic dose of 200 mg/day) or in rats at doses up to 600 mg/kg/day (31 times the human systemic exposure at the therapeutic dose).

Tenofovir DF:

Long-term oral carcinogenicity studies were conducted in mice and rats receiving tenofovir DF. In the mouse study, one male and two female mice in the 600 mg/kg/day group (15 times the human systemic exposure at the recommended human dose of 300 mg/day) had duodenal tumors. The mechanism underlying this effect is uncertain but may relate to high local drug concentrations in the gastrointestinal tract. No treatment-related tumors were seen in mice in the 100 or 300 mg/kg/day groups. In the rat study at doses of 30, 100, and 300 mg/kg/day (approximately 5 times human exposure), no treatment-related increase in tumor incidence was observed.

Mutagenesis

Efavirenz:

Efavirenz was not genotoxic in a battery of in vitro and in vivo genotoxicity assays. This included assays in four in vitro assay systems: (1) bacterial mutation assays in *Salmonella*

typhimurium and *Escherichia coli*, (2) a Chinese hamster ovary (CHO) cell/hypoxanthine-guanine phosphoribosyl-transferase (HGPRT) forward mutation assay, (3) a chromosome aberration assay in human peripheral lymphocytes, and (4) a chromosome aberration assay in CHO cells, and in one in vivo system (mouse micronucleus assay). All assays were conducted employing maximally soluble or minimally toxic doses/concentrations of efavirenz.

Emtricitabine:

Emtricitabine was not genotoxic in the reverse mutation bacterial test (Ames test), mouse lymphoma or mouse micronucleus assays.

Tenofovir DF:

Tenofovir DF was not mutagenic in the in vitro bacterial mutation (Ames) assay (*Salmonella-Escherichia coli*/Mammalian-Microsome Reverse Mutation Assay) but was mutagenic in the in vitro mouse lymphoma assay (L5178Y TK +/- Forward Mutation Assay), with and without metabolic activation. Tenofovir disoproxil fumarate was not clastogenic in the in vivo mouse micronucleus assay at plasma exposure levels of more than 10x the human exposure.

Reproductive Toxicology

Efavirenz:

Malformations were observed in 3 of 20 fetuses/infants from efavirenz-treated cynomolgus monkeys versus 0 of 20 concurrent control animals. The pregnant monkeys were given 60 mg/kg/day of efavirenz throughout pregnancy (post-coital days 20–150) and plasma drug concentrations were similar to those in humans given 600 mg/day. Anencephaly and unilateral anophthalmia were observed in one fetus, microphthalmia was observed in another fetus, and cleft palate was observed in a third fetus.

No malformations were observed in fetuses from efavirenz-treated rats; however an increase in fetal resorptions and a slight increase in pup mortality was observed at doses that produced peak plasma concentrations and AUC values in pregnant female rats similar to or lower than those achieved in humans at the recommended clinical dose. Efavirenz was not teratogenic or embryotoxic when given to pregnant rabbits.

A 5–8% decrease in mean rat pup weights versus control, and a slight increase in pup mortality was observed at ≥ 50 mg/kg BID when efavirenz was given to pregnant rats during gestation and through lactation until weaning. Peak plasma concentrations and AUC values in rats were similar to or lower than those in humans at the recommended clinical dose. No efavirenz-related effects were observed on the fertility, mating behavior, sexual maturation, learning, or behavior of the F1 generation derived from female rats given 100 mg/kg BID.

In a five-week oral infant rhesus toxicity study, infant monkeys given 30 mg/kg BID exhibited slight transitory decreases in body weight gain and food intake. Doses of 45 mg/kg BID produced adverse clinical signs including; vomiting, lethargy, dehydration, poor appetite, and/or weakness and slight decreases in body weight gain.

No efavirenz-related effects were observed on the fertility or reproductive performance of female rats given 100 mg/kg BID, or on the reproductive performance or sperm motility and morphology of male rats given 200 mg/kg BID.

Fetal Exposure to efavirenz was documented in pregnant rats, rabbits and cynomolgous monkeys. Maternal and fetal blood concentrations were equivalent in pregnant rabbits and cynomolgous monkeys and fetal blood concentrations were approximately 25% to 49% lower than the corresponding maternal concentrations in pregnant rats. Results of these studies indicated that efavirenz crossed the placenta of all species tested.

The excretion of efavirenz into rat milk was demonstrated. Efavirenz milk concentrations in rats were approximately 8-fold higher than corresponding maternal efavirenz plasma concentrations.

Emtricitabine:

The incidence of fetal variations and malformations was not increased in embryofetal toxicity studies performed with emtricitabine in mice at exposures (AUC) approximately 60-fold higher and in rabbits at approximately 120-fold higher than human exposures at the recommended daily dose.

Emtricitabine did not affect fertility in male rats at approximately 140-fold or in male and female mice at approximately 60-fold higher exposures (AUC) than in humans given the recommended 200 mg daily dose. Fertility was normal in the offspring of mice exposed daily from before birth (in utero) through sexual maturity at daily exposures (AUC) of approximately 60-fold higher than human exposures at the recommended 200 mg daily dose.

Tenofovir DF:

Tenofovir DF had no adverse effects on embryo-fetal development in rats at doses 450 mg/kg/day and in rabbits at doses up to 300 mg/kg/day (14 and 19 times the human dose based on body surface area comparisons).

Tenofovir DF had no adverse effects on fertility or general reproductive performance in rats at doses up to 600 mg/kg/day. In a study of effects on peri- and postnatal development in rats, reduced pup body weights, survival and delay in sexual maturation was in rats at the maternally toxic doses of 450 and 600 mg/kg (approximately 14 and 19 times the human dose based on body surface area comparisons). There were no adverse effects on growth, development, behavior, or reproductive parameters at non-maternally toxic doses (150 mg/kg/day).

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PART III: CONSUMER INFORMATION

Pr ATRIPLA[®]
(Efavirenz / Emtricitabine /
Tenofovir Disoproxil Fumarate Tablets)

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

ABOUT THIS MEDICATION

What the medication is used for:

ATRIPLA contains 3 medicines, SUSTIVA[®] (efavirenz), EMTRIVA[®] (emtricitabine) and VIREAD[®] (tenofovir DF), combined in one pill. EMTRIVA and VIREAD are HIV (human immunodeficiency virus) nucleoside analog reverse transcriptase inhibitor (NRTI) and SUSTIVA is an HIV non-nucleoside analog reverse transcriptase inhibitor (NNRTI). VIREAD and EMTRIVA are components of TRUVADA[®]. ATRIPLA can be used alone as a complete regimen, or in combination with other medications to treat people with HIV infection. ATRIPLA is for adults age 18 and older. ATRIPLA has not been studied in children under age 18 or adults over age 65.

What it does:

ATRIPLA helps block HIV reverse transcriptase, a chemical in your body (enzyme) that is needed for HIV to multiply. ATRIPLA lowers the amount of HIV in the blood (viral load). Lowering the amount of HIV in the blood lowers the chance of infections that happen when your immune system is weak (opportunistic infections).

HIV infection destroys CD4 (T) cells, which are important to the immune system. The immune system helps fight infection. After a large number of T cells are destroyed, acquired immune deficiency syndrome (AIDS) develops. ATRIPLA may also help to increase the number of T cells (CD4 cells).

ATRIPLA does not cure HIV infection or AIDS. The long-term effects of ATRIPLA are not known at this time. People taking ATRIPLA may still get opportunistic infections or other conditions that happen with HIV infection. Opportunistic infections

are infections that develop because the immune system is weak. Some of these conditions are pneumonia, herpes virus infections, and *Mycobacterium avium* complex (MAC) infections. **It is very important that you see your doctor regularly while taking ATRIPLA.**

ATRIPLA has not been shown to reduce the risk of passing HIV to others through sexual contact or blood contamination. Continue to practice safe sex and do not use or share dirty needles.

When it should not be used:

Together with your doctor, you need to decide whether ATRIPLA is right for you.

Do not take ATRIPLA if:

- you are taking any medication that is listed in this pamphlet under “**Drugs that should not be taken with ATRIPLA**” (see **INTERACTIONS WITH THIS MEDICATION**).
- you have or are at known risk for any type of bone disease or bone related problems and have not discussed this with your doctor.
- you are allergic to ATRIPLA or any of its ingredients. The medicinal ingredients are efavirenz, emtricitabine and tenofovir DF (See: **What the important nonmedicinal ingredients in ATRIPLA are**).

What the medicinal ingredients are:

efavirenz
 emtricitabine
 tenofovir disoproxil fumarate (tenofovir DF)

What the important nonmedicinal ingredients are:

croscarmellose sodium, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, sodium lauryl sulfate, black iron oxide, polyethylene glycol, polyvinyl alcohol, red iron oxide, talc and titanium dioxide.

What dosage forms it comes in:

ATRIPLA is available as tablets. Each tablet contains 600 mg of efavirenz, 200 mg of emtricitabine and 300 mg of tenofovir disoproxil fumarate (which is equivalent to 245 mg of tenofovir disoproxil), as active ingredients. The tablets are pink, modified capsule-shaped, film-coated, debossed with “123” on one side and plain-faced on the other side. Each bottle contains 30 tablets and a silica gel desiccant and is closed with a child-resistant closure.

WARNINGS AND PRECAUTIONS

Serious Warnings and Precautions

- The class of medicines to which emtricitabine and tenofovir DF, two of the components of ATRIPLA, belong (NRTIs) can cause a condition called lactic acidosis, together with an enlarged liver. Non-specific symptoms such as nausea, vomiting and stomach pain might indicate the development of lactic acidosis. This rare but serious side effect has occasionally been fatal. Lactic acidosis occurs more often in women, particularly if they are very overweight. You should consult your doctor immediately if such symptoms occur while you are receiving ATRIPLA. The symptoms that may indicate a liver problem include: feeling very weak, tired or uncomfortable; unusual or unexpected stomach discomfort; feeling cold; feeling dizzy or lightheaded; suddenly developing a slow or irregular heartbeat. If you notice these symptoms, stop taking ATRIPLA and consult a doctor immediately.
 - **“Flare-ups” of Hepatitis B Virus infection**, in which the disease suddenly returns in a worse way than before, can occur if you stop taking ATRIPLA. Do not stop taking ATRIPLA without your doctor’s advice. If you stop taking ATRIPLA, tell your doctor immediately about any new, unusual or worsening symptoms that you notice after stopping treatment. After you stop taking ATRIPLA, your doctor will still need to check your health and take blood tests to check your liver. ATRIPLA is not approved for the treatment of hepatitis B virus infection.
 - The most serious possible side effect is harm to the kidneys, including damage to kidney cells, kidney tissue inflammation and kidney failure. Your doctor may monitor your kidney function before beginning and while receiving ATRIPLA. Some patients treated with tenofovir DF (a component of ATRIPLA) have had kidney problems. Your doctor may need to perform additional blood tests if you have had kidney problems in the past or need to take another drug that can cause kidney problems.
 - Tenofovir DF, a component of ATRIPLA, caused harm to the bones of animals. If you notice bone pain, suffer a bone fracture, or other bone problem, consult your doctor. **If you have bone problems, you may wish to discuss calcium and/or vitamin D supplementation with your doctor. The effect of supplementation with calcium and/or vitamin D is unknown.**
- Patients who experience dizziness, trouble concentrating or drowsiness should avoid driving or operating machinery.
 - **Women should not become pregnant while taking ATRIPLA and for 12 weeks after stopping it.** Serious birth defects have been seen in animals and women treated with efavirenz (a component of ATRIPLA) during pregnancy. It is not known whether efavirenz caused these defects. A reliable form of barrier contraception must always be used even if you or your partner are using other methods of contraception such as the pill or other hormonal therapy (e.g. implants, injections). ATRIPLA may remain in your blood for a time after therapy is stopped. Therefore, you should continue use of a reliable form of contraception for 12 weeks after stopping treatment with ATRIPLA.
 - A small number of patients taking efavirenz, one of the components of ATRIPLA, have had severe depression, strange thoughts, or angry behavior. Some patients have had thoughts of suicide and a few patients have actually committed suicide. These problems tend to occur more often in patients with a history of mental illness. You should contact your doctor immediately if you think you are having these symptoms, so your doctor can decide whether you should continue to take ATRIPLA.
 - Consult your doctor if you have a rash since some rashes may be serious.
 - If you have advanced HIV infection (AIDS) and have an infection, you may develop symptoms of infection and inflammation or worsening of the symptoms of an existing infection once treatment with ATRIPLA is started. These symptoms may indicate that your body’s improved immune system is fighting infection. If you notice signs of inflammation or infection, tell your doctor at once.
- BEFORE you use ATRIPLA (efavirenz/ emtricitabine/tenofovir DF) talk to your doctor or pharmacist:**
- If you are breast-feeding:*** Do not breast-feed if you are taking ATRIPLA. Do not breast-feed if you have HIV. If you are a woman who has or will have a baby, talk with your doctor about the best way to feed your baby. If your baby does not already have HIV, there is a chance that the baby can get HIV through breast-feeding. Also, ATRIPLA may pass through milk and cause serious harm to the baby.
- If you have other medical conditions:*** Let your doctor know if you have other medical conditions, especially liver, kidney problems, pancreatitis (inflammation of the pancreas), seizures or mental illness, or have or are at risk for bone disease or bone related problems.

If you are taking other medicines: Some medicines can interact when taken together, including prescription and non-prescription medicines, herbal products and dietary supplements (see **INTERACTIONS WITH THIS MEDICATION**).

INTERACTIONS WITH THIS MEDICATION

Drugs that should not be taken with ATRIPLA:

- **Propulsid (cisapride)*, Versed (midazolam), Halcion (triazolam), ergot medications (for example Wigraine and Cafergot) or Orap (pimozide).** Taking these medications with ATRIPLA could create the potential for serious or life-threatening side effects.
- Vfend (voriconazole) since it may lose its effect or may increase the chance of having side effects from ATRIPLA.
- Do not take ATRIPLA if you are on other medications that may affect your kidneys and have not discussed this with your doctor.
- Combivir, EMTRIVA, 3TC, Heptovir, HEPSERA, Kivexa, SUSTIVA, Trizivir, TRUVADA or VIREAD. **ATRIPLA should not be used with those medicines.**
- **St. John's Wort (*Hypericum perforatum*), or products containing St. John's Wort.** St. John's Wort is a herbal product sold as a dietary supplement. Taking St. John's Wort may stop ATRIPLA from working properly.

*Cisapride is not marketed in Canada.

It is also important to tell your doctor if you are taking any of the following:

- Reyataz (atazanavir sulfate), Fortovase, Invirase (saquinavir) or Biaxin (clarithromycin); **these medicines need to be replaced with another medicine when taken with ATRIPLA. If your doctor does prescribe Reyataz and ATRIPLA together, you may need to be monitored more carefully for side effects.**
- Crixivan (indinavir), Methadone, Mycobutin (rifabutin) or Zoloft (sertraline); **these medicines may need to have their dose changed when taken with ATRIPLA.**
- Videx or Videx EC (didanosine); Tenofovir DF (a component of ATRIPLA) may increase the amount of didanosine in your blood, which could result in more side effects. **You may need to be monitored more carefully** if you are taking ATRIPLA and didanosine together. Also, the dose of didanosine may need to be changed.
- Kaletra (lopinavir/ritonavir); these medicines may increase the amount of Tenofovir DF (a component of ATRIPLA) in your blood, which

could result in more side effects. **You may need to be monitored more carefully** if you are taking ATRIPLA and Kaletra together. Also, the dose of Kaletra may need to be changed.

- Medicines for seizures [for example, Dilantin (phenytoin), Tegretol (carbamazepine), or phenobarbital]; your doctor may want to switch you to another medicine or check drug levels in your blood from time to time.
- Sporanox (itraconazole) may need to be replaced by another medicine when taken with ATRIPLA.
- The cholesterol-lowering medicines Lipitor (atorvastatin), Pravachol (pravastatin sodium), and Zocor (simvastatin).
- Rifadin (rifampin) or the rifampin-containing medicines Rofact and Rifater.
- Calcium channel blockers such as Cardizem, or Tiazac (diltiazem), Covera HS, Isoptin SR or Tarka (verapamil), and others.
- Immunosuppressants such as Neoral (cyclosporin), Advagraf or Prograf (tacrolimus), Rapamune or Torisel (sirolimus).
- The effect of combining alcohol or recreational (street, illicit) drugs with efavirenz has not been studied. Because they may interact with each other, speak with your doctor before you combine ATRIPLA with these drugs.

These are not all the medicines that may cause problems if you take ATRIPLA. Be sure to tell your doctor about all the medicines you take.

Keep a complete list of all the prescription and nonprescription medicines as well as any herbal medicines that you are taking, how much you take and how often you take them. Make a new list when medicines or herbal medicines are added or stopped, or if the dose changes. Give copies of this list to all your doctors and pharmacists every time you visit them or fill a prescription. This will give your doctor a complete picture of the medicines you use. Then he or she can decide the best approach for the situation.

PROPER USE OF THIS MEDICATION

Stay under a doctor's care when taking ATRIPLA. Do not change your treatment or stop treatment without first talking with your doctor.

Take ATRIPLA every day exactly as your doctor prescribed it. Follow the directions from your doctor, exactly as written on the label. Set up a dosing schedule and follow it carefully.

When your ATRIPLA supply starts to run low, get more from your doctor or pharmacy. This is very important because the amount of virus in your blood may increase if the medicine is stopped for even a short time. The virus may develop resistance to ATRIPLA and become harder to treat.

Only take medicine that has been prescribed specifically for you. Do not give ATRIPLA to others or take medicine prescribed for someone else. Do not use if seal over bottle opening is broken or missing.

Usual Adult Dose:

- The usual dose of ATRIPLA is one tablet orally (by mouth) once a day.
- ATRIPLA should be taken on an empty stomach. Taking ATRIPLA at bedtime may make some side effects less bothersome.

Overdosage:

If you suspect that you took more than the prescribed dose of ATRIPLA, contact your local poison control center or emergency room right away. As with all medicines, ATRIPLA should be kept out of reach of children.

Missed Dose:

It is important that you do not miss any doses. If you miss a dose of ATRIPLA, take it as soon as you remember that day. **Do not** take more than 1 dose of ATRIPLA in a day. **Do not** take 2 doses at the same time. Call your doctor or pharmacist if you are not sure what to do.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

The most common side effects of SUSTIVA, EMTRIVA and VIREAD are:

- Nervous system symptoms such as dizziness, trouble sleeping, drowsiness, trouble concentrating, unusual dreams
- Headache
- Diarrhea
- Nausea
- Vomiting
- Rash
- Flatulence (intestinal gas)
- Tiredness
- Itching
- Allergic reaction (including swelling of the face, lips, tongue or throat)
- Abdominal pain

Other side effects may include pancreatitis (inflammation of the pancreas) and shortness of breath.

Skin discoloration (small spots or freckles) may also happen with ATRIPLA.

A small number of patients taking efavirenz, one of the components of ATRIPLA, have had severe depression, strange thoughts, or angry behavior. Some patients have had thoughts of suicide and a few patients have actually committed suicide. These problems tend to occur more often in patients with a history of mental illness. Contact your doctor immediately if you think you are having these symptoms so your doctor can decide whether you should continue to take ATRIPLA.

Changes in body fat have been seen in some patients taking antiretroviral therapy. These changes may include increased amounts of fat in the upper back and neck (“buffalo hump”), breast, and around the trunk. Loss of fat from the legs, arms and face may also happen. The cause and long term health effects of these conditions are not known at this time.

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM				
Symptoms / Effect		Talk with your doctor or pharmacist		Stop taking drug and call your doctor or pharmacist
		Only if severe	In all cases	
Rare	Effect: Lactic acidosis Symptoms <ul style="list-style-type: none"> • Feeling very weak or tired • Unusual muscle pain • Stomach pain with nausea and vomiting • Feeling cold especially in arms and legs • Feeling dizzy or lightheaded • Fast or irregular heartbeat 		<ul style="list-style-type: none"> ✓ ✓ ✓ ✓ ✓ 	
Very Rare	Effect: Hepatotoxicity (severe liver problems) with hepatomegaly (liver enlargement) and steatosis (fat in the liver) Symptoms <ul style="list-style-type: none"> • Jaundice (skin or the white part of eyes turn yellow) • Urine turns dark • Bowel movements (stools) turn light in color • Loss of appetite for several days or longer • Feeling sick to your 		<ul style="list-style-type: none"> ✓ ✓ ✓ ✓ ✓ 	

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM				
Symptoms / Effect		Talk with your doctor or pharmacist		Stop taking drug and call your doctor or pharmacist
		Only if severe	In all cases	
	stomach (nausea) • Lower stomach pain		✓	
Very Rare	Effect: Flare-ups of hepatitis B virus infection following drug discontinuation Symptoms • Jaundice (skin or the white part of eyes turn yellow) • Urine turns dark • Bowel movements (stools) turn light in color • Loss of appetite for several days or longer • Feeling sick to your stomach (nausea) • Lower stomach pain		✓ ✓ ✓ ✓ ✓	
Rare	Effect: Kidney problems Symptoms • You may have increased or decreased urination as well as increased thirst • You may have swelling of your legs and feet • You may feel listless and tired		✓ ✓ ✓	

Lactic acidosis is a medical emergency and must be treated in the hospital. You may be more likely to get lactic acidosis or serious liver problems if you are very overweight (obese) or have been taking nucleoside analog medicines, like ATRIPLA, for a long time.

Muscle pain, muscle weakness, bone pain and softening of the bone (infrequently contributing to fractures) have also been reported due to tenofovir DF (a component of ATRIPLA).

There have been other side effects in patients taking SUSTIVA, EMTRIVA or VIREAD. This is **not** a complete list of side effects. If you have questions about side effects, ask your doctor, nurse, or pharmacist. You should report any new or continuing symptoms to your doctor right away. Your doctor may be able to help you manage these side effects.

HOW TO STORE IT

- Keep ATRIPLA and all other medications out of reach of children.
- ATRIPLA should be stored at 15–30 °C (59–86 °F). It should remain stable until the expiration date printed on the label.
- Do not keep your medicine in places that are too hot or cold.
- Do not keep medicine that is out of date or that you no longer need. If you throw any medicines away make sure that children will not find them.
- Keep ATRIPLA in its original container and keep the container tightly closed.

REPORTING SUSPECTED SIDE EFFECTS

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

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

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

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

MORE INFORMATION

This document plus the full Product Monograph, prepared for health professionals can be found at: www.gilead.ca or by contacting the sponsor, Gilead Sciences Canada, Inc., at: 1-866-207-4267

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Bristol-Myers Squibb & Gilead Sciences, LLC
Foster City, CA 94404
USA

Gilead Sciences Canada, Inc.
Mississauga, ON L5N 2W3

Bristol-Myers Squibb Canada
Montreal, Quebec H4S 0A4

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