

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

EMTRIVA® (emtricitabine) capsule, for oral use

EMTRIVA® (emtricitabine) oral solution

Initial U.S. Approval: 2003

WARNING: LACTIC ACIDOSIS/SEVERE HEPATOMEGALY WITH STEATOSIS and POST TREATMENT EXACERBATION OF HEPATITIS B

See full prescribing information for complete boxed warning.

- Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of nucleoside analogs. (5.1)
- Emtriva is not approved for the treatment of chronic Hepatitis B virus (HBV) infection. Severe acute exacerbations of Hepatitis B have been reported in patients who have discontinued EMTRIVA. Hepatic function should be monitored closely in patients coinfecting with HIV-1 and HBV. If appropriate, initiation of anti-Hepatitis B therapy may be warranted. (5.2)

RECENT MAJOR CHANGES

Indications and Usage (1)	11/2012
Warnings and Precautions	
Coadministration with Other Products (5.3)	11/2012
Immune Reconstitution Syndrome (5.6)	11/2011

INDICATIONS AND USAGE

EMTRIVA, a nucleoside analog HIV-1 reverse transcriptase inhibitor, is indicated in combination with other antiretroviral agents for the treatment of HIV-1 infection. (1)

DOSAGE AND ADMINISTRATION

- EMTRIVA may be taken without regard to food. (2.1)
- Adult Patients (18 years of age and older) (2.2):
 - EMTRIVA capsules: one 200 mg capsule administered once daily orally.
 - EMTRIVA oral solution: 240 mg (24 mL) administered once daily orally.
- Pediatric Patients (0–3 months of age) (2.3):
 - EMTRIVA oral solution: 3 mg/kg administered once daily orally.
- Pediatric Patients (3 months through 17 years) (2.4):
 - EMTRIVA oral solution: 6 mg/kg up to a maximum of 240 mg (24 mL) administered once daily orally.
 - EMTRIVA capsules: for children weighing more than 33 kg who can swallow an intact capsule, one 200 mg capsule administered once daily orally.
- Dose interval adjustment in adult patients with renal impairment (2.5):

Formulation	Creatinine Clearance (mL/min)			
	≥50 mL/min	30–49 mL/min	15–29 mL/min	<15 mL/min or on hemodialysis ^a
Capsule (200 mg)	200 mg every 24 hours	200 mg every 48 hours	200 mg every 72 hours	200 mg every 96 hours
Oral Solution (10 mg/mL)	240 mg every 24 hours (24 mL)	120 mg every 24 hours (12 mL)	80 mg every 24 hours (8 mL)	60 mg every 24 hours (6 mL)

- a. Hemodialysis Patients: If dosing on day of dialysis, give dose after dialysis.

DOSAGE FORMS AND STRENGTHS

- Capsules: 200 mg (3)
- Oral solution: 10 mg per mL (3)

CONTRAINDICATIONS

EMTRIVA is contraindicated in patients with previously demonstrated hypersensitivity to any of the components of the products. (4)

WARNINGS AND PRECAUTIONS

- Products with same active ingredient: Do not use with other emtricitabine-containing products (e.g., ATRIPLA, COMPLERA, STRIBILD, and TRUVADA). (5.3)
- Redistribution/accumulation of body fat: Observed in patients receiving antiretroviral therapy. (5.5)
- Immune reconstitution syndrome: May necessitate further evaluation and treatment. (5.6)

ADVERSE REACTIONS

Most common adverse reactions (incidence ≥10%) are headache, diarrhea, nausea, fatigue, dizziness, depression, insomnia, abnormal dreams, rash, abdominal pain, asthenia, increased cough, and rhinitis. Skin hyperpigmentation was very common (≥10%) in pediatric patients. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Gilead Sciences, Inc. at 1-800-GILEAD-5 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

USE IN SPECIFIC POPULATIONS

- Nursing mothers: Women infected with HIV should be instructed not to breast feed. (8.3)
- Pediatrics: Dose adjustment based on age and weight. (2.3, 2.4, 12.3)

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

Revised: November 2012

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FULL PRESCRIBING INFORMATION

WARNING: LACTIC ACIDOSIS/SEVERE HEPATOMEGALY WITH STEATOSIS and POST TREATMENT EXACERBATION OF HEPATITIS B

Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of nucleoside analogs alone or in combination with other antiretrovirals [See *Warnings and Precautions (5.1)*].

EMTRIVA is not approved for the treatment of chronic hepatitis B virus (HBV) infection and the safety and efficacy of EMTRIVA have not been established in patients coinfecting with HBV and HIV-1. Severe acute exacerbations of hepatitis B have been reported in patients who have discontinued EMTRIVA. Hepatic function should be monitored closely with both clinical and laboratory follow-up for at least several months in patients who are coinfecting with HIV-1 and HBV and discontinue EMTRIVA. If appropriate, initiation of anti-hepatitis B therapy may be warranted [See *Warnings and Precautions (5.2)*].

1 INDICATIONS AND USAGE

EMTRIVA[®] is indicated in combination with other antiretroviral agents for the treatment of HIV-1 infection.

Additional important information regarding the use of EMTRIVA for the treatment of HIV-1 Infection:

- EMTRIVA should not be coadministered with ATRIPLA[®], COMPLERA[®], STRIBILD[™], TRUVADA[®], or lamivudine-containing products [See *Warnings and Precautions (5.3)*].
- In treatment-experienced patients, the use of EMTRIVA should be guided by laboratory testing and treatment history [See *Clinical Pharmacology (12.4)*].

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dose

EMTRIVA may be taken without regard to food.

2.2 Adult Patients (18 years of age and older)

- **EMTRIVA capsules:** one 200 mg capsule administered once daily orally.
- **EMTRIVA oral solution:** 240 mg (24 mL) administered once daily orally.

2.3 Pediatric Patients (0–3 months of age)

- **EMTRIVA oral solution:** 3 mg per kg administered once daily orally.

2.4 Pediatric Patients (3 months through 17 years)

- **EMTRIVA oral solution:** 6 mg per kg up to a maximum of 240 mg (24 mL) administered once daily orally.

- **EMTRIVA capsules:** for children weighing more than 33 kg who can swallow an intact capsule, one 200 mg capsule administered once daily orally.

2.5 Dose Adjustment in Adult Patients with Renal Impairment

Significantly increased drug exposures were seen when EMTRIVA was administered to subjects with renal impairment [See *Clinical Pharmacology (12.3)*]. Therefore, the dosing interval or dose of EMTRIVA should be adjusted in patients with baseline creatinine clearance less than 50 mL per min using the following guidelines (see Table 1). The safety and effectiveness of these dose adjustment guidelines have not been clinically evaluated. Therefore, clinical response to treatment and renal function should be closely monitored in these patients.

Table 1 Dose Adjustment in Adult Patients with Renal Impairment

Formulation	Creatinine Clearance (mL/min)			
	≥50 mL/min	30–49 mL/min	15–29 mL/min	<15 mL/min or on hemodialysis ^a
Capsule (200 mg)	200 mg every 24 hours	200 mg every 48 hours	200 mg every 72 hours	200 mg every 96 hours
Oral Solution (10 mg/mL)	240 mg every 24 hours (24 mL)	120 mg every 24 hours (12 mL)	80 mg every 24 hours (8 mL)	60 mg every 24 hours (6 mL)

a. Hemodialysis Patients: If dosing on day of dialysis, give dose after dialysis.

Although there are insufficient data to recommend a specific dose adjustment of EMTRIVA in pediatric patients with renal impairment, a reduction in the dose and/or an increase in the dosing interval similar to adjustments for adults should be considered.

3 DOSAGE FORMS AND STRENGTHS

EMTRIVA is available as capsules and oral solution.

EMTRIVA capsules, containing 200 mg of emtricitabine, are size 1 hard gelatin capsules with a blue cap and white body, printed with “200 mg” in black on the cap and “GILEAD” and the corporate logo in black on the body.

EMTRIVA oral solution is a clear, orange to dark orange liquid containing 10 mg of emtricitabine per mL.

4 CONTRAINDICATIONS

EMTRIVA is contraindicated in patients with previously demonstrated hypersensitivity to any of the components of the products.

5 WARNINGS AND PRECAUTIONS

5.1 Lactic Acidosis/Severe Hepatomegaly with Steatosis

Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of nucleoside analogs alone or in combination, including emtricitabine and other antiretrovirals. 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 diseases; however, cases have also been reported in patients with no known risk factors. Treatment with EMTRIVA 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).

5.2 Patients Coinfected with HIV-1 and HBV

It is recommended that all patients with HIV-1 be tested for the presence of chronic Hepatitis B virus (HBV) before initiating antiretroviral therapy. EMTRIVA is not approved for the treatment of chronic HBV infection and the safety and efficacy of EMTRIVA have not been established in patients coinfecting with HBV and HIV-1. Severe acute exacerbations of Hepatitis B have been reported in patients after the discontinuation of EMTRIVA. 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 monitored closely with both clinical and laboratory follow-up for at least several months in patients who are coinfecting with HIV-1 and HBV and discontinue EMTRIVA. If appropriate, initiation of anti-Hepatitis B therapy may be warranted.

5.3 Coadministration with Related Products

EMTRIVA is a component of ATRIPLA (a fixed-dose combination of efavirenz, emtricitabine, and tenofovir disoproxil fumarate), COMPLERA (a fixed-dose combination of emtricitabine, rilpivirine, and tenofovir disoproxil fumarate), STRIBILD (a fixed-dose combination of elvitegravir, cobicistat, emtricitabine, and tenofovir disoproxil fumarate), and TRUVADA (a fixed-dose combination of emtricitabine and tenofovir disoproxil fumarate). EMTRIVA should not be coadministered with ATRIPLA, COMPLERA, STRIBILD, or TRUVADA. Due to similarities between emtricitabine and lamivudine, EMTRIVA should not be coadministered with other drugs containing lamivudine, including Combivir (lamivudine/zidovudine), Epivir or Epivir-HBV (lamivudine), Epzicom (abacavir sulfate/lamivudine), or Trizivir (abacavir sulfate/lamivudine/zidovudine).

5.4 New Onset or Worsening Renal Impairment

Emtricitabine is principally eliminated by the kidney. Reduction of the dosage of EMTRIVA is recommended for patients with impaired renal function [*See Dosage and Administration (2.5) and Clinical Pharmacology (12.3)*].

5.5 Fat Redistribution

Redistribution/accumulation of body fat including central obesity, dorsocervical fat enlargement (buffalo hump), peripheral wasting, facial wasting, breast enlargement, and “cushingoid appearance” have been observed in patients receiving antiretroviral therapy. The mechanism and long-term consequences of these events are currently unknown. A causal relationship has not been established.

5.6 Immune Reconstitution Syndrome

Immune reconstitution syndrome has been reported in patients treated with combination antiretroviral therapy, including EMTRIVA. During the initial phase of combination antiretroviral treatment, patients whose immune system responds may develop an inflammatory response to indolent or residual opportunistic infections [such as *Mycobacterium avium* infection, cytomegalovirus, *Pneumocystis jirovecii* pneumonia (PCP), or tuberculosis], which may necessitate further evaluation and treatment.

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

6 ADVERSE REACTIONS

The following adverse reactions are discussed in other sections of the labeling:

- Lactic acidosis/severe hepatomegaly with steatosis [See *Boxed Warning, Warnings and Precautions (5.1)*].
- Severe acute exacerbations of Hepatitis B [See *Boxed Warning, Warnings and Precautions (5.2)*].
- Immune reconstitution syndrome [See *Warnings and Precautions (5.6)*].

6.1 Adverse Reactions from Clinical Trials Experience

Clinical Trials in Adult Subjects

More than 2,000 adult subjects with HIV-1 infection have been treated with EMTRIVA alone or in combination with other antiretroviral agents for periods of 10 days to 200 weeks in clinical trials.

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The most common adverse reactions (incidence greater than or equal to 10%, any severity) identified from any of the 3 large controlled clinical trials include headache, diarrhea, nausea, fatigue, dizziness, depression, insomnia, abnormal dreams, rash, abdominal pain, asthenia, increased cough, and rhinitis.

Studies 301A and 303– Treatment Emergent Adverse Reactions: The most common adverse reactions that occurred in subjects receiving EMTRIVA with other antiretroviral agents in clinical trials 301A and 303 were headache, diarrhea, nausea, and rash, which were generally of mild to moderate severity. Approximately 1% of subjects discontinued participation in the clinical trials due to these events. All adverse reactions were reported with similar frequency in EMTRIVA and control treatment groups with the exception of skin discoloration which was reported with higher frequency in the EMTRIVA treated group.

Skin discoloration, manifested by hyperpigmentation on the palms and/or soles was generally mild and asymptomatic. The mechanism and clinical significance are unknown.

A summary of EMTRIVA treatment-emergent clinical adverse reactions in Studies 301A and 303 is provided in Table 2.

Table 2 Selected Treatment-Emergent Adverse Reactions (All Grades, Regardless of Causality) Reported in ≥3% of EMTRIVA-Treated Subjects in Either Study 301A or 303 (0–48 Weeks)

	303		301A	
	EMTRIVA + ZDV/d4T + NNRTI/PI (N=294)	Lamivudine + ZDV/d4T + NNRTI/PI (N=146)	EMTRIVA + didanosine + efavirenz (N=286)	Stavudine + didanosine + efavirenz (N=285)
Body as a Whole				
Abdominal pain	8%	11%	14%	17%
Asthenia	16%	10%	12%	17%
Headache	13%	6%	22%	25%
Digestive System				
Diarrhea	23%	18%	23%	32%
Dyspepsia	4%	5%	8%	12%
Nausea	18%	12%	13%	23%
Vomiting	9%	7%	9%	12%
Musculoskeletal				
Arthralgia	3%	4%	5%	6%
Myalgia	4%	4%	6%	3%
Nervous System				
Abnormal dreams	2%	<1%	11%	19%
Depressive disorders	6%	10%	9%	13%
Dizziness	4%	5%	25%	26%
Insomnia	7%	3%	16%	21%
Neuropathy/peripheral neuritis	4%	3%	4%	13%
Paresthesia	5%	7%	6%	12%
Respiratory				
Increased cough	14%	11%	14%	8%
Rhinitis	18%	12%	12%	10%
Skin				
Rash event ^a	17%	14%	30%	33%

a. Rash event includes rash, pruritus, maculopapular rash, urticaria, vesicubullous rash, pustular rash, and allergic reaction.

Studies 301A and 303– Laboratory Abnormalities: Laboratory abnormalities in these trials occurred with similar frequency in the EMTRIVA and comparator groups. A summary of Grades 3-4 laboratory abnormalities is provided in Table 3 below.

Table 3 Treatment-Emergent Grades 3-4 Laboratory Abnormalities Reported in ≥1% of EMTRIVA-Treated Subjects in Either Study 301A or 303

	303		301A	
	EMTRIVA + ZDV/d4T + NNRTI/PI (N=294)	Lamivudine + ZDV/d4T + NNRTI/PI (N=146)	EMTRIVA + Didanosine + Efavirenz (N=286)	Stavudine + Didanosine + Efavirenz (N=285)
Percentage with grade 3 or grade 4 laboratory abnormality	31%	28%	34%	38%
ALT (>5.0 x ULN ^a)	2%	1%	5%	6%
AST (>5.0 x ULN)	3%	<1%	6%	9%
Bilirubin (>2.5 x ULN)	1%	2%	<1%	<1%
Creatine kinase (>4.0 x ULN)	11%	14%	12%	11%
Neutrophils (<750 mm ³)	5%	3%	5%	7%
Pancreatic amylase (>2.0 x ULN)	2%	2%	<1%	1%
Serum amylase (>2.0 x ULN)	2%	2%	5%	10%
Serum glucose <40 or >250 mg/dL)	3%	3%	2%	3%
Serum lipase (>2.0 x ULN)	<1%	<1%	1%	2%
Triglycerides (>750 mg/dL)	10%	8%	9%	6%

a. ULN = Upper limit of normal

Study 934– Treatment Emergent Adverse Reactions: In Study 934, 511 antiretroviral-naïve subjects received either VIREAD[®] + EMTRIVA administered in combination with efavirenz (N=257) or zidovudine/lamivudine administered in combination with efavirenz (N=254). Adverse reactions observed in this trial were generally consistent with those seen in previous trials in treatment-experienced or treatment-naïve subjects (Table 4).

Table 4 Selected Treatment-Emergent Adverse Reactions^a (Grades 2–4) Reported in ≥5% in Any Treatment Group in Study 934 (0–144 Weeks)

	TDF ^b + EMTRIVA + EFV	AZT/3TC + EFV
	N=257	N=254
Gastrointestinal Disorder		
Diarrhea	9%	5%
Nausea	9%	7%
Vomiting	2%	5%
General Disorders and Administration Site Condition		
Fatigue	9%	8%
Infections and Infestations		
Sinusitis	8%	4%
Upper respiratory tract infections	8%	5%
Nasopharyngitis	5%	3%
Nervous System Disorders		
Headache	6%	5%
Dizziness	8%	7%
Psychiatric Disorders		
Depression	9%	7%
Insomnia	5%	7%
Skin and Subcutaneous Tissue Disorders		
Rash event ^c	7%	9%

- a. Frequencies of adverse reactions are based on all treatment-emergent adverse events, regardless of relationship to study drug.
- b. From Weeks 96 to 144 of the trial, subjects received TRUVADA with efavirenz in place of VIREAD + EMTRIVA with efavirenz.
- c. Rash event includes rash, exfoliative rash, rash generalized, rash macular, rash maculo-papular, rash pruritic, and rash vesicular.

Study 934 – Laboratory Abnormalities: Significant laboratory abnormalities observed in this trial are shown in Table 5.

Table 5 Significant Laboratory Abnormalities Reported in ≥1% of Subjects in Any Treatment Group in Study 934 (0–144 Weeks)

	TDF ^a + EMTRIVA + EFV	AZT/3TC + EFV
	N=257	N=254
Any ≥ Grade 3 Laboratory Abnormality	30%	26%
Fasting Cholesterol (>240 mg/dL)	22%	24%
Creatine Kinase (M: >990 U/L) (F: >845 U/L)	9%	7%
Serum Amylase (>175 U/L)	8%	4%
Alkaline Phosphatase (>550 U/L)	1%	0%
AST (M: >180 U/L) (F: >170 U/L)	3%	3%
ALT (M: >215 U/L) (F: >170 U/L)	2%	3%
Hemoglobin (<8.0 mg/dL)	0%	4%
Hyperglycemia (>250 mg/dL)	2%	1%
Hematuria (>75 RBC/HPF)	3%	2%
Glycosuria (3+)	<1%	1%
Neutrophils (<750/mm ³)	3%	5%
Fasting Triglycerides (>750 mg/dL)	4%	2%

a. From Weeks 96 to 144 of the trial, subjects received TRUVADA with efavirenz in place of VIREAD + EMTRIVA with efavirenz.

Clinical Trials in Pediatric Subjects

Assessment of adverse reactions is based on data from Study 203, an open label, uncontrolled trial of 116 HIV-1-infected pediatric subjects who received emtricitabine through 48 weeks. The adverse reaction profile in pediatric subjects was generally comparable to that observed in clinical trials of EMTRIVA in adult subjects [See *Adverse Reactions (6.1)*]. Hyperpigmentation was more frequent in children. Additional adverse reactions identified from this trial include anemia.

Selected treatment-emergent adverse events, regardless of causality, reported in subjects during 48 weeks of treatment were the following: infection (44%), hyperpigmentation (32%), increased cough (28%), vomiting (23%), otitis media (23%),

rash (21%), rhinitis (20%), diarrhea (20%), fever (18%), pneumonia (15%), gastroenteritis (11%), abdominal pain (10%), and anemia (7%). Treatment-emergent grades 3-4 laboratory abnormalities were experienced by 9% of pediatric subjects, including elevated amylase ($>2.0 \times \text{ULN}$) (n=4), decreased neutrophils ($<750/\text{mm}^3$) (n=3), elevated ALT ($>5 \times \text{ULN}$) (n=2), elevated CPK ($>4 \times \text{ULN}$) (n=2) and one subject each with elevated bilirubin ($>3.0 \times \text{ULN}$), elevated GGT ($>10 \times \text{ULN}$), elevated lipase ($>2.5 \times \text{ULN}$), decreased hemoglobin ($<7 \text{ g/dL}$), and decreased glucose ($<40 \text{ mg/dL}$).

7 DRUG INTERACTIONS

The potential for drug interactions with EMTRIVA has been studied in combination with zidovudine, indinavir, stavudine, famciclovir, and tenofovir disoproxil fumarate. There were no clinically significant drug interactions for any of these drugs. Drug interactions trials are described elsewhere in the labeling [*See Clinical Pharmacology (12.3)*].-

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Pregnancy Category B

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. There are, however, no adequate and well-controlled trials in pregnant women. Because animal reproduction studies are not always predictive of human response, EMTRIVA should be used during pregnancy only if clearly needed.

Antiretroviral Pregnancy Registry: To monitor fetal outcomes of pregnant women exposed to EMTRIVA, an Antiretroviral Pregnancy Registry has been established. Healthcare providers are encouraged to register patients by calling 1-800-258-4263.

8.3 Nursing Mothers

Nursing Mothers: The Centers for Disease Control and Prevention recommend that HIV-1-infected mothers not breast-feed their infants to avoid risking postnatal transmission of HIV-1. Samples of breast milk obtained from five HIV-1 infected mothers show that emtricitabine is secreted in human milk. Breastfeeding infants whose mothers are being treated with emtricitabine may be at risk for developing viral resistance to emtricitabine. Other emtricitabine-associated risks in infants breastfed by mothers being treated with emtricitabine are unknown. Because of both the potential for HIV-1 transmission and the potential for serious adverse reactions in nursing infants, **mothers should be instructed not to breast-feed if they are receiving EMTRIVA.**

8.4 Pediatric Use

The safety and efficacy of emtricitabine in patients between 3 months and 21 years of age is supported by data from three open-label, non-randomized clinical trials in which emtricitabine was administered to 169 HIV-1 infected treatment-naive and experienced (defined as virologically suppressed on a lamivudine containing regimen for which emtricitabine was substituted for lamivudine) subjects [*See Clinical Studies(14.3)*].

The pharmacokinetics of emtricitabine were studied in 20 neonates born to HIV-1-positive mothers [See *Clinical Studies (14.3)*]. All neonates were HIV-1 negative at the end of the trial; the efficacy of emtricitabine in preventing or treating HIV-1 could not be determined.

8.5 Geriatric Use

Clinical trials of EMTRIVA did not include sufficient numbers of subjects aged 65 years and over to determine whether they respond differently from 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.

8.6 Patients with Impaired Renal Function

It is recommended that the dose or dosing interval for EMTRIVA be modified in patients with creatinine clearance less than 50 mL/min or in patients who require dialysis [See *Dosage and Administration (2.5)*].

10 OVERDOSAGE

There is no known antidote for EMTRIVA. Limited clinical experience is available at doses higher than the therapeutic dose of EMTRIVA. In one clinical pharmacology trial single doses of emtricitabine 1200 mg were administered to 11 subjects. No severe adverse reactions were reported.

The effects of higher doses are not known. If overdose occurs the patient should be monitored for signs of toxicity, and standard supportive treatment applied as necessary.

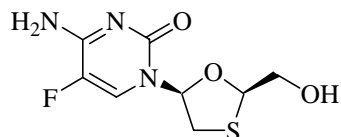
Hemodialysis treatment removes approximately 30% of the emtricitabine dose over a 3-hour dialysis period starting within 1.5 hours of emtricitabine dosing (blood flow rate of 400 mL/min and a dialysate flow rate of 600 mL/min). It is not known whether emtricitabine can be removed by peritoneal dialysis.

11 DESCRIPTION

EMTRIVA is the brand name of emtricitabine, a synthetic nucleoside analog with activity against human immunodeficiency virus type 1 (HIV-1) reverse transcriptase.

The chemical name of emtricitabine is 5-fluoro-1-(2*R*,5*S*)-[2-(hydroxymethyl)-1,3-oxathiolan-5-yl]cytosine. Emtricitabine is the (-) enantiomer of a thio analog of cytidine, which differs from other cytidine analogs in that it has a fluorine in the 5-position.

It has a molecular formula of C₈H₁₀FN₃O₃S and a molecular weight of 247.24. It has the following structural formula:



Emtricitabine is a white to off-white powder with a solubility of approximately 112 mg/mL in water at 25 °C. The log P for emtricitabine is -0.43 and the pKa is 2.65.

EMTRIVA is available as capsules or as an oral solution.

EMTRIVA capsules are for oral administration. Each capsule contains 200 mg of emtricitabine and the inactive ingredients crospovidone, magnesium stearate, microcrystalline cellulose, povidone, titanium dioxide, gelatin, and FD&C blue No. 2.

EMTRIVA oral solution is for oral administration. One milliliter (1 mL) of EMTRIVA oral solution contains 10 mg of emtricitabine in an aqueous solution with the following inactive ingredients: cotton candy flavor, FD&C yellow No. 6, edetate disodium, methylparaben and propylparaben (added as preservatives), sodium phosphate (monobasic), propylene glycol, water, and xylitol (added as a sweetener). Sodium hydroxide and hydrochloric acid may be used to adjust pH.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Emtricitabine is an antiviral drug [See *Clinical Pharmacology* (12.4)].

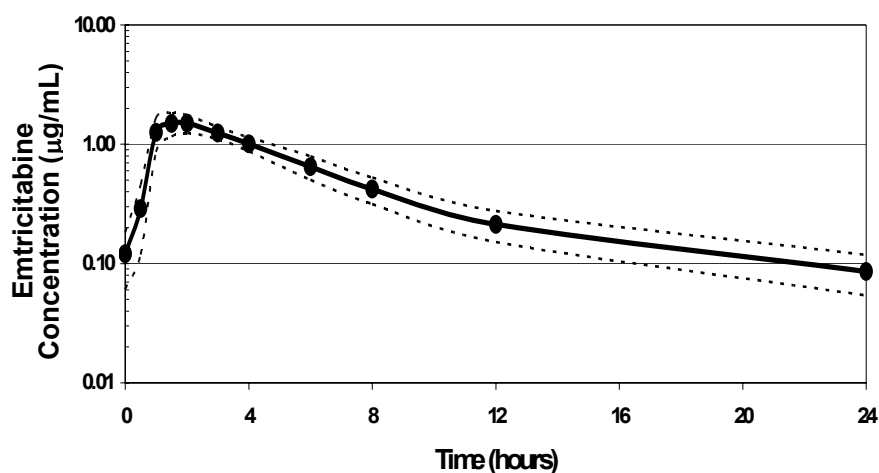
12.3 Pharmacokinetics

Adults

The pharmacokinetics of emtricitabine were evaluated in healthy subjects and HIV-1-infected subjects. Emtricitabine pharmacokinetics are similar between these populations.

Figure 1 shows the mean steady-state plasma emtricitabine concentration-time profile in 20 HIV-1-infected subjects receiving EMTRIVA capsules.

Figure 1 Mean (\pm 95% CI) Steady-State Plasma Emtricitabine Concentrations in HIV-1-Infected Adults (N=20)



Absorption

Emtricitabine is rapidly and extensively absorbed following oral administration with peak plasma concentrations occurring at 1–2 hours post-dose. Following multiple dose oral administration of EMTRIVA capsules to 20 HIV-1-infected subjects, the (mean \pm SD) steady-state plasma emtricitabine peak concentration (C_{max}) was 1.8 ± 0.7 µg/mL and the area-under the plasma concentration-time curve over a 24-hour dosing interval (AUC) was 10.0 ± 3.1 µg·hr/mL. The mean steady state plasma trough concentration at

24 hours post-dose was 0.09 µg/mL. The mean absolute bioavailability of EMTRIVA capsules was 93% while the mean absolute bioavailability of EMTRIVA oral solution was 75%. The relative bioavailability of EMTRIVA oral solution was approximately 80% of EMTRIVA capsules.

The multiple dose pharmacokinetics of emtricitabine are dose proportional over a dose range of 25–200 mg.

Distribution

In vitro binding of emtricitabine to human plasma proteins was less than 4% and independent of concentration over the range of 0.02–200 µg/mL. At peak plasma concentration, the mean plasma to blood drug concentration ratio was ~1.0 and the mean semen to plasma drug concentration ratio was ~4.0.

Metabolism

In vitro studies indicate that emtricitabine is not an inhibitor of human CYP450 enzymes. Following administration of ¹⁴C-emtricitabine, complete recovery of the dose was achieved in urine (~86%) and feces (~14%). Thirteen percent (13%) of the dose was recovered in urine as three putative metabolites. The biotransformation of emtricitabine includes oxidation of the thiol moiety to form the 3'-sulfoxide diastereomers (~9% of dose) and conjugation with glucuronic acid to form 2'-O-glucuronide (~4% of dose). No other metabolites were identifiable.

Elimination

The plasma emtricitabine half-life is approximately 10 hours. The renal clearance of emtricitabine is greater than the estimated creatinine clearance, suggesting elimination by both glomerular filtration and active tubular secretion. There may be competition for elimination with other compounds that are also renally eliminated.

Effects of Food on Oral Absorption

EMTRIVA capsules and oral solution may be taken with or without food. Emtricitabine systemic exposure (AUC) was unaffected while C_{max} decreased by 29% when EMTRIVA capsules were administered with food (an approximately 1000 kcal high-fat meal). Emtricitabine systemic exposure (AUC) and C_{max} were unaffected when 200 mg EMTRIVA oral solution was administered with either a high-fat or low-fat meal.

Special Populations

Race, Gender

The pharmacokinetics of emtricitabine were similar in adult male and female subjects and no pharmacokinetic differences due to race have been identified.

Pediatric Patients

The pharmacokinetics of emtricitabine at steady state were determined in 77 HIV-1-infected pediatric subjects, and the pharmacokinetic profile was characterized in four age groups (Table 6). The emtricitabine exposure achieved in pediatric subjects receiving a daily dose of 6 mg/kg up to a maximum of 240 mg oral solution or a 200 mg

capsule is similar to exposures achieved in adult subjects receiving a once-daily dose of 200 mg.

The pharmacokinetics of emtricitabine were studied in 20 neonates born to HIV-1-positive mothers. Each mother received prenatal and intrapartum combination antiretroviral therapy. Neonates received up to 6 weeks of zidovudine prophylactically after birth. The neonates were administered two short courses of emtricitabine oral solution (each 3 mg/kg once daily x 4 days) during the first 3 months of life. The AUC observed in neonates who received a daily dose of 3 mg/kg of emtricitabine was similar to the AUC observed in pediatric subjects ages 3 months to 17 years who received a daily dose of emtricitabine as a 6 mg/kg oral solution up to 240 mg or as a 200 mg capsule (Table 6).

Table 6 Mean ± SD Pharmacokinetic Parameters by Age Groups for Pediatric Subjects and Neonates Receiving EMTRIVA Capsules or Oral Solution

Age	HIV-1-exposed Neonates	HIV-1-infected Pediatric Subjects			
	0–3 mo (N=20) ^a	3–24 mo (N=14)	25 mo–6 yr (N=19)	7–12yr (N=17)	13–17 yr (N=27)
Formulation					
Capsule (n)	0	0	0	10	26
Oral Solution (n)	20	14	19	7	1
Dose (mg/kg) ^b	3.1 (2.9-3.4)	6.1 (5.5–6.8)	6.1 (5.6–6.7)	5.6 (3.1–6.6)	4.4 (1.8–7.0)
C _{max} (µg/mL)	1.6 ± 0.6	1.9 ± 0.6	1.9 ± 0.7	2.7 ± 0.8	2.7 ± 0.9
AUC (µg·hr/mL)	11.0 ± 4.2	8.7 ± 3.2	9.0 ± 3.0	12.6 ± 3.5	12.6 ± 5.4
T _{1/2} (hr)	12.1 ± 3.1	8.9 ± 3.2	11.3 ± 6.4	8.2 ± 3.2	8.9 ± 3.3

a. Two pharmacokinetic evaluations were conducted in 20 neonates over the first 3 months of life. Median (range) age of infant on day of pharmacokinetic evaluation was 26 (5–81) days.

b. Mean (range)

Geriatric Patients

The pharmacokinetics of emtricitabine have not been fully evaluated in the elderly.

Patients with Impaired Renal Function

The pharmacokinetics of emtricitabine are altered in subjects with renal impairment [See *Warnings and Precautions (5.4)*]. In adult subjects with creatinine clearance less than 50 mL/min or with end-stage renal disease (ESRD) requiring dialysis, C_{max} and AUC of emtricitabine were increased due to a reduction in renal clearance (Table 7). It is recommended that the dosing interval for EMTRIVA be modified in adult patients with creatinine clearance less than 50 mL/min or in adult patients with ESRD who require dialysis [See *Dosage and Administration (2.5)*]. The effects of renal impairment on emtricitabine pharmacokinetics in pediatric patients are not known.

Table 7 Mean ± SD Pharmacokinetic Parameters in Adult Subjects with Varying Degrees of Renal Function

Creatinine Clearance (mL/min)	>80 (N=6)	50–80 (N=6)	30–49 (N=6)	<30 (N=5)	ESRD ^a <30 (N=5)
Baseline creatinine clearance (mL/min)	107 ± 21	59.8 ± 6.5	40.9 ± 5.1	22.9 ± 5.3	8.8 ± 1.4
C _{max} (µg/mL)	2.2 ± 0.6	3.8 ± 0.9	3.2 ± 0.6	2.8 ± 0.7	2.8 ± 0.5
AUC (µg·hr/mL)	11.8 ± 2.9	19.9 ± 1.2	25.1 ± 5.7	33.7 ± 2.1	53.2 ± 9.9
CL/F (mL/min)	302 ± 94	168 ± 10	138 ± 28	99 ± 6	64 ± 12
CLr (mL/min)	213 ± 89	121 ± 39	69 ± 32	30 ± 11	NA ^b

a. ESRD subjects requiring dialysis

b. NA = Not Applicable

Hemodialysis: 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). It is not known whether emtricitabine can be removed by peritoneal dialysis.

Patients with Hepatic Impairment

The pharmacokinetics of emtricitabine have not been studied in subjects with hepatic impairment, however, emtricitabine is not metabolized by liver enzymes, so the impact of liver impairment should be limited.

Assessment of Drug Interactions

At concentrations up to 14-fold higher than those observed *in vivo*, emtricitabine did not inhibit *in vitro* drug metabolism mediated by any of the following human CYP isoforms: CYP1A2, CYP2A6, CYP2B6, CYP2C9, CYP2C19, CYP2D6, and CYP3A4.

Emtricitabine did not inhibit the enzyme responsible for glucuronidation (uridine-5'-disphosphoglucuronyl transferase). Based on the results of these *in vitro* experiments and the known elimination pathways of emtricitabine, the potential for CYP mediated interactions involving emtricitabine with other medicinal products is low.

EMTRIVA has been evaluated in healthy volunteers in combination with tenofovir disoproxil fumarate (DF), zidovudine, indinavir, famciclovir, and stavudine. Tables 8 and 9 summarize the pharmacokinetic effects of coadministered drug on emtricitabine pharmacokinetics and effects of emtricitabine on the pharmacokinetics of coadministered drug.

Table 8 Drug Interactions: Change in Pharmacokinetic Parameters for Emtricitabine in the Presence of the Coadministered Drug^a

Coadministered Drug	Dose of Coadministered Drug (mg)	Emtricitabine Dose (mg)	N	% Change of Emtricitabine Pharmacokinetic Parameters ^b (90% CI)		
				C _{max}	AUC	C _{min}
Tenofovir DF	300 once daily x 7 days	200 once daily x 7 days	17	↔	↔	↑ 20 (↑ 12 to ↑ 29)
Zidovudine	300 twice daily x 7 days	200 once daily x 7 days	27	↔	↔	↔
Indinavir	800 x 1	200 x 1	12	↔	↔	NA
Famciclovir	500 x 1	200 x 1	12	↔	↔	NA
Stavudine	40 x 1	200 x 1	6	↔	↔	NA

a. All interaction trials conducted in healthy volunteers.

b. ↑ = Increase; ↓ = Decrease; ↔ = No Effect; NA = Not Applicable

Table 9 Drug Interactions: Change in Pharmacokinetic Parameters for Coadministered Drug in the Presence of Emtricitabine^a

Coadministered Drug	Dose of Coadministered Drug (mg)	Emtricitabine Dose (mg)	N	% Change of Coadministered Drug Pharmacokinetic Parameters ^b (90% CI)		
				C _{max}	AUC	C _{min}
Tenofovir DF	300 once daily x 7 days	200 once daily x 7 days	17	↔	↔	↔
Zidovudine	300 twice daily x 7 days	200 once daily x 7 days	27	↑ 17 (↑ 0 to ↑ 38)	↑ 13 (↑ 5 to ↑ 20)	↔
Indinavir	800 x 1	200 x 1	12	↔	↔	NA
Famciclovir	500 x 1	200 x 1	12	↔	↔	NA
Stavudine	40 x 1	200 x 1	6	↔	↔	NA

a. All interaction trials conducted in healthy volunteers.

b. ↑ = Increase; ↓ = Decrease; ↔ = No Effect; NA = Not Applicable

12.4 Microbiology

Mechanism of Action

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 reverse transcriptase 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 polymerase α , β , ϵ , and mitochondrial DNA polymerase γ .

Antiviral Activity

The antiviral activity in cell culture of emtricitabine against laboratory and clinical isolates of HIV-1 was assessed in lymphoblastoid cell lines, the MAGI-CCR5 cell line, and peripheral blood mononuclear cells. The 50% effective concentration (EC₅₀) value for emtricitabine was in the range of 0.0013–0.64 μM (0.0003–0.158 μg/mL). In drug combination trials of emtricitabine with nucleoside reverse transcriptase inhibitors (abacavir, lamivudine, stavudine, tenofovir, zalcitabine, zidovudine), non-nucleoside reverse transcriptase inhibitors (delavirdine, efavirenz, nevirapine), and protease inhibitors (amprenavir, nelfinavir, ritonavir, saquinavir), additive to synergistic effects were observed. Emtricitabine displayed antiviral activity in cell culture against HIV-1 clades A, B, C, D, E, F, and G (EC₅₀ values ranged from 0.007–0.075 μM) and showed strain specific activity against HIV-2 (EC₅₀ values ranged from 0.007–1.5 μM).

The *in vivo* activity of emtricitabine was evaluated in two clinical trials in which 101 subjects were administered 25–400 mg a day of EMTRIVA as monotherapy for 10–14 days. A dose-related antiviral effect was observed, with a median decrease from baseline in plasma HIV-1 RNA of 1.3 log₁₀ at a dose of 25 mg once daily and 1.7 log₁₀ to 1.9 log₁₀ at a dose of 200 mg once daily or twice daily.

Resistance

Emtricitabine-resistant isolates of HIV-1 have been selected in cell culture and *in vivo*. Genotypic analysis of these isolates showed that the reduced susceptibility to emtricitabine was associated with a substitution in the HIV-1 reverse transcriptase gene at codon 184 which resulted in an amino acid substitution of methionine by valine or isoleucine (M184V/I).

Emtricitabine-resistant isolates of HIV-1 have been recovered from some subjects treated with emtricitabine alone or in combination with other antiretroviral agents. In a clinical trial of treatment-naïve subjects treated with EMTRIVA, didanosine, and efavirenz [See *Clinical Studies (14.1)*], viral isolates from 37.5% of subjects with virologic failure showed reduced susceptibility to emtricitabine. Genotypic analysis of these isolates showed that the resistance was due to M184V/I substitutions in the HIV-1 reverse transcriptase gene.

In a clinical trial of treatment-naïve subjects treated with either EMTRIVA, VIREAD, and efavirenz or zidovudine/lamivudine and efavirenz [See *Clinical Studies (14.1)*], resistance analysis was performed on HIV-1 isolates from all confirmed virologic failure subjects with greater than 400 copies/mL of HIV-1 RNA at Week 144 or early discontinuation. Development of efavirenz resistance-associated substitutions occurred most frequently and was similar between the treatment arms. The M184V amino acid substitution, associated with resistance to EMTRIVA and lamivudine, was observed in 2/19 analyzed subject isolates in the EMTRIVA + VIREAD group and in 10/29 analyzed subject isolates in the lamivudine/zidovudine group. Through 144 weeks of Study 934, no subjects have developed a detectable K65R substitution in their HIV-1 as analyzed through standard genotypic analysis.

Cross Resistance

Cross-resistance among certain nucleoside analog reverse transcriptase inhibitors has been recognized. Emtricitabine-resistant isolates (M184V/I) were cross-resistant to lamivudine and zalcitabine but retained sensitivity in cell culture to didanosine, stavudine, tenofovir, zidovudine, and NNRTIs (delavirdine, efavirenz, and nevirapine). 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 substitutions 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.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

In long-term oral carcinogenicity studies of emtricitabine, no drug-related increases in tumor incidence were 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 reverse mutation bacterial test (Ames test), 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.

14 CLINICAL STUDIES

14.1 Treatment-Naive Adult Patients

Study 934

Data through 144 weeks are reported for Study 934, a randomized, open-label, active-controlled multicenter clinical trial comparing EMTRIVA + tenofovir disoproxil fumarate (tenofovir DF) administered in combination with efavirenz versus zidovudine/lamivudine fixed-dose combination administered in combination with efavirenz in 511 antiretroviral-naive subjects. From Weeks 96 to 144 of the trial, subjects received EMTRIVA/tenofovir DF fixed-dose combination with efavirenz in place of EMTRIVA + tenofovir DF with efavirenz. Subjects had a mean age of 38 years (range 18–80), 86% were male, 59% 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). Subjects were stratified by baseline CD4⁺ cell count (< or ≥200 cells/mm³); 41% had CD4⁺ cell counts <200 cells/mm³ and 51% of subjects had baseline viral loads >100,000 copies/mL. Treatment outcomes through 48 and 144 weeks for those subjects who did not have efavirenz resistance at baseline are presented in Table 10.

Table 10 Outcomes of Randomized Treatment at Week 48 and 144 (Study 934)

Outcomes	Week 48		Week 144	
	EMTRIVA +TDF +EFV (N=244)	AZT/3TC +EFV (N=243)	EMTRIVA + TDF +EFV (N=227) ^a	AZT/3TC +EFV (N=229) ^a
Responder ^b	84%	73%	71%	58%
Virologic failure ^c	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 ^d	10%	14%	20%	22%

a. Subjects who were responders at Week 48 or Week 96 (HIV-1 RNA <400 copies/mL) but did not consent to continue in the trial after Week 48 or Week 96 were excluded from analysis.

b. Subjects achieved and maintained confirmed HIV-1 RNA <400 copies/mL through Weeks 48 and 144.

c. Includes confirmed viral rebound and failure to achieve confirmed <400 copies/mL through Weeks 48 and 144.

d. Includes lost to follow-up, subject withdrawal, noncompliance, protocol violation and other reasons.

Through Week 48, 84% and 73% of subjects in the EMTRIVA + tenofovir DF group and the zidovudine/lamivudine group, respectively, achieved and maintained HIV-1 RNA <400 copies/mL (71% and 58% through Week 144). The difference in the proportion of subjects 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 trial. In addition, 80% and 70% of subjects in the EMTRIVA + tenofovir DF group and the zidovudine/lamivudine group, respectively, achieved and maintained HIV-1 RNA <50 copies/mL through Week 48 (64% and 56% through Week 144). The mean increase from baseline in CD4⁺ cell count was 190 cells/mm³ in the EMTRIVA + tenofovir DF group and 158 cells/mm³ in the zidovudine/lamivudine group at Week 48 (312 and 271 cells/mm³ at Week 144).

Through 48 weeks, 7 subjects in the EMTRIVA + tenofovir DF group and 5 subjects in the zidovudine/lamivudine group experienced a new CDC Class C event (10 and 6 subjects through 144 weeks).

Study 301A

Study 301A was a 48 week double-blind, active-controlled multicenter clinical trial comparing EMTRIVA (200 mg once daily) administered in combination with didanosine and efavirenz versus stavudine, didanosine and efavirenz in 571 antiretroviral naive adult subjects. Subjects had a mean age of 36 years (range 18–69), 85% were male, 52% Caucasian, 16% African-American and 26% Hispanic. Subjects had a mean baseline CD4⁺ cell count of 318 cells/mm³ (range 5–1317) and a median baseline

plasma HIV-1 RNA of 4.9 log₁₀ copies/mL (range 2.6–7.0). Thirty-eight percent of subjects had baseline viral loads >100,000 copies/mL and 31% had CD4⁺ cell counts <200 cells/mL. Treatment outcomes are presented in Table 11 below.

Table 11 Outcomes of Randomized Treatment at Week 48 (Study 301A)

Outcomes	EMTRIVA + Didanosine + Efavirenz (N=286)	Stavudine + Didanosine + Efavirenz (N=285)
Responder ^a	81% (78%)	68% (59%)
Virologic Failure ^b	3%	11%
Death	0%	<1%
Discontinuation Due to Adverse Event	7%	13%
Discontinuation for Other Reasons ^c	9%	8%

- a. Subjects achieved and maintained confirmed HIV RNA <400 copies/mL (<50 copies/mL) through Week 48.
- b. Includes subjects who failed to achieve virologic suppression or rebounded after achieving virologic suppression.
- c. Includes lost to follow-up, subject withdrawal, non-compliance, protocol violation and other reasons.

The mean increase from baseline in CD4⁺ cell count was 168 cells/mm³ for the EMTRIVA arm and 134 cells/mm³ for the stavudine arm.

Through 48 weeks in the EMTRIVA group, 5 subjects (1.7%) experienced a new CDC Class C event, compared to 7 subjects (2.5%) in the stavudine group.

14.2 Treatment-Experienced Adult Patients

Study 303

Study 303 was a 48 week, open-label, active-controlled multicenter clinical trial comparing EMTRIVA (200 mg once daily) to lamivudine, in combination with stavudine or zidovudine and a protease inhibitor or NNRTI in 440 adult subjects who were on a lamivudine-containing triple-antiretroviral drug regimen for at least 12 weeks prior to trial entry and had HIV-1 RNA ≤400 copies/mL.

Subjects were randomized 1:2 to continue therapy with lamivudine (150 mg twice daily) or to switch to EMTRIVA (200 mg once daily). All subjects were maintained on their stable background regimen. Subjects had a mean age of 42 years (range 22–80), 86% were male, 64% Caucasian, 21% African-American and 13% Hispanic. Subjects had a mean baseline CD4⁺ cell count of 527 cells/mm³ (range 37–1909), and a median baseline plasma HIV-1 RNA of 1.7 log₁₀ copies/mL (range 1.7–4.0).

The median duration of prior antiretroviral therapy was 27.6 months. Treatment outcomes are presented in Table 12 below.

Table 12 Outcomes of Randomized Treatment at Week 48 (Study 303)

Outcomes	EMTRIVA + ZDV/d4T + NNRTI/PI (N=294)	Lamivudine + ZDV/d4T + NNRTI/PI (N=146)
Responder ^a	77% (67%)	82% (72%)
Virologic Failure ^b	7%	8%
Death	0%	<1%
Discontinuation Due to Adverse Event	4%	0%
Discontinuation for Other Reasons ^c	12%	10%

a. Subjects achieved and maintained confirmed HIV RNA <400 copies/mL (<50 copies/mL) through Week 48.

b. Includes subjects who failed to achieve virologic suppression or rebounded after achieving virologic suppression.

c. Includes lost to follow-up, subject withdrawal, non-compliance, protocol violation and other reasons.

The mean increase from baseline in CD4⁺ cell count was 29 cells/mm³ for the EMTRIVA arm and 61 cells/mm³ for the lamivudine arm.

Through 48 weeks, in the EMTRIVA group 2 subjects (0.7%) experienced a new CDC Class C event, compared to 2 subjects (1.4%) in the lamivudine group.

14.3 Pediatric Patients

In three open-label, non-randomized clinical trials, emtricitabine was administered to 169 HIV-1 infected treatment-naive and experienced (defined as virologically suppressed on a lamivudine containing regimen for which emtricitabine was substituted for lamivudine) subjects between 3 months and 21 years of age. Subjects received once-daily EMTRIVA oral solution (6 mg/kg to a maximum of 240 mg/day) or EMTRIVA capsules (a single 200 mg capsule once daily) in combination with at least two other antiretroviral agents.

Subjects had a mean age of 7.9 years (range 0.3–21), 49% were male, 15% Caucasian, 61% Black and 24% Hispanic. Subjects had a median baseline HIV-1 RNA of 4.6 log₁₀ copies/mL (range 1.7–6.4) and a mean baseline CD4⁺ cell count of 745 cells/mm³ (range 2–2650). Through 48 weeks of therapy, the overall proportion of subjects who achieved and sustained an HIV-1 RNA <400 copies/mL was 86%, and <50 copies/mL was 73%. The mean increase from baseline in CD4⁺ cell count was 232 cells/mm³ (-945, +1512). The adverse reaction profile observed during these clinical trials was similar to that of adult subjects, with the exception of the occurrence of anemia and higher frequency of hyperpigmentation in children [See *Adverse Reactions (6)*].

The pharmacokinetics of emtricitabine were studied in 20 neonates born to HIV-1-positive mothers. Each mother received prenatal and intrapartum combination antiretroviral therapy. Neonates received up to 6 weeks of zidovudine prophylactically after birth. The neonates were administered two short courses of emtricitabine oral solution (each 3 mg/kg once daily x 4 days) during the first 3 months of life.

Emtricitabine exposures in neonates were similar to the exposures achieved in subjects ages 3 months to 17 years [See *Clinical Pharmacology (12.3)*]. During the two short dosing periods on emtricitabine there were no safety issues identified in the treated

neonates. All neonates were HIV-1 negative at the end of the trial; the efficacy of emtricitabine in preventing or treating HIV-1 could not be determined.

16 HOW SUPPLIED/STORAGE AND HANDLING

Capsules

The size 1 hard gelatin capsules with a blue cap and white body contain 200 mg of emtricitabine, are printed with “200 mg” in black on the cap and “GILEAD” and the corporate logo in black on the body, and are available in unit of use bottles (closed with induction sealed child-resistant closures) of:

- 30 capsules (NDC 61958–0601–1).

Store at 25 °C (77 °F); excursions permitted to 15 °C–30 °C (59 °F–86 °F).

Oral Solution

The oral solution is a clear, orange to dark orange liquid, contains 10 mg/mL of emtricitabine, and is available in unit of use plastic, amber bottles (closed with child resistant closures and packaged with a marked dosing cup) of:

- 170 mL (NDC 61958–0602–1).

Store refrigerated, 2 °C–8 °C (36 °F–46 °F). EMTRIVA oral solution should be used within 3 months if stored by the patient at 25 °C (77 °F); excursions permitted to 15 °C–30 °C (59 °F–86 °F).

17 PATIENT COUNSELING INFORMATION

See *FDA-approved patient labeling (Patient Information)*.

17.1 Information for Patients

Patients should be advised that:

- EMTRIVA is not a cure for HIV-1 infection and patients may continue to experience illnesses associated with HIV-1 infection, including opportunistic infections. Patients should remain under the care of a physician when using EMTRIVA.
- Patients should avoid doing things that can spread HIV to others.
 - **Do not share needles or other injection equipment.**
 - **Do not share personal items that can have blood or body fluids on them, like toothbrushes and razor blades.**
 - **Do not have any kind of sex without protection.** Always practice safer sex by using a latex or polyurethane condom to lower the chance of sexual contact with semen, vaginal secretions, or blood.
 - **Do not breastfeed.** EMTRIVA is secreted in breast milk. Mothers with HIV-1 should not breastfeed because HIV-1 can be passed to the baby in the breast milk.
- The long term effects of EMTRIVA are unknown.

- EMTRIVA capsules and oral solution are for oral ingestion only.
- It is important to take EMTRIVA with combination therapy on a regular dosing schedule to avoid missing doses.
- Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported. Treatment with EMTRIVA should be suspended in any patient who develops clinical symptoms suggestive of lactic acidosis or pronounced hepatotoxicity (including nausea, vomiting, unusual or unexpected stomach discomfort, and weakness) [See *Warnings and Precautions (5.1)*].
- Patients with HIV-1 should be tested for Hepatitis B virus (HBV) before initiating antiretroviral therapy.
- Severe acute exacerbations of Hepatitis B have been reported in patients who are coinfecting with HBV and HIV-1 and have discontinued EMTRIVA.
- EMTRIVA should not be coadministered with ATRIPLA, COMPLERA, STRIBILD, or TRUVADA; or with other drugs containing lamivudine, including Combivir (lamivudine/zidovudine), Epivir or Epivir-HBV (lamivudine), Epzicom (abacavir sulfate/lamivudine), or Trizivir (abacavir sulfate/lamivudine/zidovudine) [See *Warnings and Precautions (5.3)*].
- Dose or dosing interval of EMTRIVA may need adjustment in patients with renal impairment [See *Dosage and Administration (2.5)*].

PATIENT INFORMATION

EMTRIVA[®] (em-treev'-ah) capsules

EMTRIVA oral solution

Generic name: emtricitabine (em tri SIT uh bean)

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

You should stay under a healthcare provider's care when taking EMTRIVA. **Do not change or stop your medicine without first talking with your healthcare provider.** Talk to your healthcare provider or pharmacist if you have any questions about EMTRIVA.

What is the most important information I should know about EMTRIVA?

- Some people who have taken medicines like EMTRIVA (a nucleoside analog) have developed a serious condition called lactic acidosis (buildup of an acid in the blood). Lactic acidosis can be a medical emergency and may need to be treated in the hospital. Call your healthcare provider right away if you get the following signs of lactic acidosis.
 - You feel very weak or tired.

- You have unusual (not normal) muscle pain.
- You have trouble breathing.
- You have stomach pain with nausea and vomiting.
- You feel cold, especially in your arm and legs.
- You feel dizzy or lightheaded.
- You have a fast or irregular heartbeat.
- **Some people who have taken medicines like EMTRIVA have developed serious liver problems called hepatotoxicity, with liver enlargement (hepatomegaly) and fat in the liver (steatosis). Call your healthcare provider right away if you get the following signs of liver problems.**
 - Your skin or the white part of your eyes turns yellow (jaundice).
 - Your urine turns dark.
 - Your bowel movements (stools) turn light in color.
 - You don't feel like eating food for several days or longer.
 - You feel sick to your stomach (nausea).
 - You have lower stomach area (abdominal) pain.
- **You may be more likely to get lactic acidosis or liver problems** if you are female, very overweight (obese), or have been taking nucleoside analog medicines, like EMTRIVA, for a long time.
- **If you are also infected with the Hepatitis B Virus (HBV),** you need close medical follow-up for several months after stopping treatment with EMTRIVA. Follow-up includes medical exams and blood tests to check for HBV that is getting worse. **Patients with HBV infection, who take EMTRIVA and then stop it, may get “flare-ups” of their hepatitis. A “flare-up” is when the disease suddenly returns in a worse way than before.**

What is EMTRIVA?

EMTRIVA is a type of medicine called an HIV-1 (human immunodeficiency virus) nucleoside reverse transcriptase inhibitor (NRTI). EMTRIVA is always used with other anti-HIV-1 medicines to treat people with HIV-1 infection. **EMTRIVA is for adults and children, but has not been studied fully in adults over age 65.**

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.

EMTRIVA helps to block HIV-1 reverse transcriptase, a chemical in your body (enzyme) that is needed for HIV-1 to multiply. EMTRIVA may lower the amount of HIV-1 in the blood (viral load). EMTRIVA may also help to increase the number of T cells called CD4⁺ cells. Lowering the amount of HIV-1 in the blood lowers the chance of death or infections that happen when your immune system is weak (opportunistic infections).

EMTRIVA does not cure HIV-1 infection or AIDS. The long-term effects of EMTRIVA are not known at this time. People taking EMTRIVA may still get opportunistic infections or other conditions that happen with HIV-1 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 healthcare provider regularly while taking EMTRIVA.**

Who should not take EMTRIVA?

- Do not take EMTRIVA if you are allergic to EMTRIVA or any of its ingredients. The active ingredient is emtricitabine. See the end of this leaflet for a complete list of ingredients.
- Do not take EMTRIVA if you are already taking ATRIPLA[®], COMPLERA[®], STRIBILD[™], TRUVADA[®], Combivir, Efavir, Efavir-HBV, Epzicom, or Trizivir because these medicines contain the same or similar active ingredients.

What should I tell my healthcare provider before taking EMTRIVA?

Tell your healthcare provider

- **If you are pregnant or planning to become pregnant.** We do not know if EMTRIVA can harm your unborn child. You and your healthcare provider will need to decide if EMTRIVA is right for you. If you use EMTRIVA while you are pregnant, talk to your healthcare provider about how you can be on the EMTRIVA Antiviral Pregnancy Registry.
- **If you are breast-feeding.** You should not breast feed if you are HIV-positive because of the chance of passing the HIV virus to your baby. Also, EMTRIVA can pass into your breast milk and it is not known if it can harm your baby. If you are a woman who has or will have a baby, talk with your healthcare provider about the best way to feed your baby.
- **If you have kidney problems.** You may need to take EMTRIVA less often.
- **If you have any liver problems including Hepatitis B Virus infection.**
- **Tell your healthcare provider about all your medical conditions.**
- **Tell your healthcare provider about all the medicines you take** such as prescription and non-prescription medicines and dietary supplements. Keep a complete list of all the medicines that you take. Make a new list when medicines are added or stopped. Give copies of this list to all of your healthcare providers and pharmacist **every** time you visit or fill a prescription.

How should I take EMTRIVA?

- Take EMTRIVA by mouth exactly as your healthcare provider prescribed it. Follow the directions from your healthcare provider, exactly as written on the label.
- Dosing in adults: The usual dose of EMTRIVA is 1 capsule once a day.

- Dosing in children: The child's doctor will calculate the right dose of EMTRIVA (oral solution or capsule) based on the child's weight.
- EMTRIVA is always used with other anti-HIV-1 medicines.
- EMTRIVA may be taken with or without a meal. Food does not affect how EMTRIVA works.
- If you forget to take EMTRIVA, take it as soon as you remember that day. **Do not** take more than 1 dose of EMTRIVA in a day. **Do not** take 2 doses at the same time. Call your healthcare provider or pharmacist if you are not sure what to do. **It is important that you do not miss any doses of EMTRIVA or your other anti-HIV-1 medicines.**
- When your EMTRIVA supply starts to run low, get more from your healthcare provider 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 EMTRIVA and become harder to treat.
- Stay under a healthcare provider's care when taking EMTRIVA. Do not change your treatment or stop treatment without first talking with your healthcare provider.
- If you take too much EMTRIVA, call your local poison control center or emergency room right away.

What should I avoid while taking EMTRIVA?

Avoid doing things that can spread HIV-1 infection.

- **Do not share needles or other injection equipment.**
- **Do not share personal items that can have blood or body fluids on them, like toothbrushes or razor blades.**
- **Do not have any kind of sex without protection.** Always practice safer sex by using a latex or polyurethane condom or other barrier to lower the chance of sexual contact with semen, vaginal secretions, or blood.
- **Do not breastfeed.** EMTRIVA can be passed to your baby in your breast milk. Also, mothers with HIV-1 should not breastfeed because HIV-1 can be passed to the baby in the breast milk.

What are the possible side effects of EMTRIVA?

EMTRIVA may cause the following serious side effects (See "What is the most important information I should know about EMTRIVA?"):

- **lactic acidosis** (buildup of an acid in the blood). Lactic acidosis can be a medical emergency and may need to be treated in the hospital. **Call your doctor right away if you get signs of lactic acidosis** (See "What is the most important information I should know about EMTRIVA?").
- **serious liver problems (hepatotoxicity)**, with liver enlargement (hepatomegaly) and fat in the liver (steatosis). Call your healthcare provider right away if you get any

signs of liver problems (See “What is the most important information I should know about EMTRIVA?”).

- **“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 EMTRIVA. EMTRIVA is not for the treatment of Hepatitis B Virus (HBV) infection.

Other side effects with EMTRIVA when used with other anti-HIV-1 medicines include:

- Changes in body fat have been seen in some patients taking EMTRIVA and other anti-HIV-1 medicines. These changes may include increased amount of fat in the upper back and neck (“buffalo hump”), breast, and around the main part of your body (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.

The most common side effects of EMTRIVA used with other anti-HIV-1 medicines are headache, diarrhea, and nausea. Other side effects include allergic reaction, dizziness, sleeping problems, abnormal dreams, vomiting, indigestion, stomach pain, pain, weakness and rash. Skin discoloration may also happen with EMTRIVA.

There have been other side effects in patients taking EMTRIVA. However, these side effects may have been due to other medicines that patients were taking or to HIV-1 itself. Some of these side effects can be serious.

This list of side effects is **not** complete. If you have questions about side effects, ask your healthcare provider or pharmacist. You should report any new or continuing symptoms to your healthcare provider right away. Your healthcare provider may be able to help you manage these side effects.

How do I store EMTRIVA?

- Keep EMTRIVA and all other medicines out of reach of children.
- Store EMTRIVA capsules between 59 °F and 86 °F (15 °C to 30 °C).
- Store EMTRIVA oral solution in a refrigerator between 36 °F and 46 °F (2 °C–8 °C). Do not freeze. Alternatively, the product may be stored at room temperature for up to 3 months and any remaining solution in the bottle must be discarded after the 3 months.
- 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.

General information about EMTRIVA:

Medicines are sometimes prescribed for conditions that are not mentioned in patient information leaflets. Do not use EMTRIVA for a condition for which it was not prescribed. Do not give EMTRIVA to other people, even if they have the same symptoms you have. It may harm them.

This leaflet summarizes the most important information about EMTRIVA. If you would like more information, talk with your doctor. You can ask your healthcare provider or

pharmacist for information about EMTRIVA that is written for health professionals. For more information, you may also call 1-800-GILEAD5.

What are the ingredients of EMTRIVA?

Active Ingredient: emtricitabine.

Inactive Ingredients for EMTRIVA capsules: crospovidone, magnesium stearate, microcrystalline cellulose, povidone, titanium dioxide, gelatin, and FD&C blue No. 2.

Inactive Ingredients for EMTRIVA oral solution: Cotton candy flavor, FD&C yellow No. 6, edetate disodium, methylparaben and propylparaben, sodium phosphate (monobasic), propylene glycol, water, and xylitol. Sodium hydroxide and hydrochloric acid may be used to adjust pH.

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