

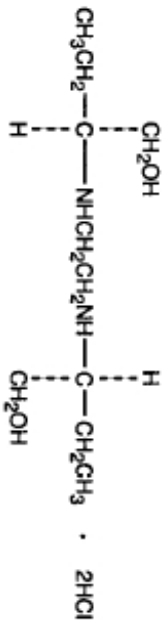
**ETHAMBUTOL HYDROCHLORIDE - ethambutol hydrochloride tablet**  
State of Florida DOH Central Pharmacy

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**ETHAMBUTOL HYDROCHLORIDE TABLETS, USP**

**Rx only**

**Description:**

Ethambutol hydrochloride is an oral chemotherapeutic agent which is specifically effective against actively growing microorganisms of the genus *Mycobacterium*, including *M. tuberculosis*. It is a white, crystalline powder. Freely soluble in water; soluble in alcohol and in methanol; slightly soluble in ether and in chloroform. It has the chemical formula of: (+)-2,2'(Ethylenediimino)-di-1-butanol dihydrochloride. The structural formula is as follows:



**C<sub>10</sub>H<sub>24</sub>N<sub>2</sub>O<sub>2</sub>·2HCl Molecular Weight: 277.23**

Each tablet, for oral administration, contains 400 mg of ethambutol hydrochloride. In addition, each tablet contains the following inactive ingredients: colloidal silicon dioxide, compressible sugar, gelatin, hydroxypropyl methylcellulose, magnesium stearate, methylcellulose, polydextrose, polyethylene glycol, sodium lauryl sulfate, stearic acid, titanium dioxide, and triacetin.

**Clinical Pharmacology:**

Ethambutol hydrochloride following a single oral dose of 25 mg/kg of body weight, attains a peak of 2 to 5 micrograms/mL in serum 2 to 4 hours after administration. When the drug is administered daily for longer periods of time at this dose, serum levels are similar. The serum level of ethambutol hydrochloride falls to undetectable levels by 24 hours after the last dose except in some patients with abnormal renal function. The intracellular concentrations of erythrocytes reach peak values approximately twice those of plasma and maintain this ratio throughout the 24 hours.

During the 24-hour period following oral administration of ethambutol hydrochloride approximately 50 percent of the initial dose is excreted unchanged in the urine, while an additional 8 to 15 percent appears

in the form of metabolites. The main path of metabolism appears to be an initial oxidation of the alcohol to an aldehydic intermediate, followed by conversion to a dicarboxylic acid. From 20 to 22 percent of the initial dose is excreted in the feces as unchanged drug. No drug accumulation has been observed with consecutive single daily doses of 25 mg/kg in patients with normal kidney function, although marked accumulation has been demonstrated in patients with renal insufficiency.

Ethambutol diffuses into actively growing *Mycobacterium* cells such as tubercle bacilli. Ethambutol appears to inhibit the synthesis of one or more metabolites, thus causing impairment of cell metabolism, arrest of multiplication, and cell death. No cross resistance with other available antimycobacterial agents has been demonstrated.

Ethambutol has been shown to be effective against strains of *Mycobacterium tuberculosis* but does not seem to be active against fungi, viruses, or other bacteria. *Mycobacterium tuberculosis* strains previously unexposed to ethambutol have been uniformly sensitive to concentrations of 8 or less micrograms/mL, depending on the nature of the culture media. When ethambutol has been used alone for treatment of tuberculosis, tubercle bacilli from these patients have developed resistance to ethambutol hydrochloride by *in vitro* susceptibility tests; the development of resistance has been unpredictable and appears to occur in a step-like manner. No cross resistance between ethambutol and other antituberculous drugs has been reported. Ethambutol has reduced the incidence of the emergence of mycobacterial resistance to isoniazid when both drugs have been used concurrently.

An agar diffusion microbiologic assay, based upon inhibition of *Mycobacterium smegmatis* (ATCC 607) may be used to determine concentrations of ethambutol in serum and urine.

#### **ANIMAL PHARMACOLOGY:**

Toxicological studies in dogs on high prolonged doses produced evidence of myocardial damage and failure, and depigmentation of the tapetum lucidum of the eyes, the significance of which is not known. Degenerative changes in the central nervous system, apparently not dose-related, have also been noted in dogs receiving ethambutol hydrochloride over a prolonged period.

In the rhesus monkey, neurological signs appeared after treatment with high doses given daily over a period of several months. These were correlated with specific serum levels of ethambutol hydrochloride and with definite neuroanatomical changes in the central nervous system. Focal interstitial carditis was also noted in monkeys which received ethambutol hydrochloride in high doses for a prolonged period.

When pregnant mice or rabbits were treated with high doses of ethambutol hydrochloride, fetal mortality was slightly but not significantly ( $P > 0.05$ ) increased. Female rats treated with ethambutol hydrochloride displayed slight but insignificant ( $P > 0.05$ ) decreases in fertility and litter size.

In fetuses born of mice treated with high doses of ethambutol hydrochloride during pregnancy, a low incidence of cleft palate, exencephaly and abnormality of the vertebral column were observed. Minor abnormalities of the cervical vertebra were seen in the newborn of rats treated with high doses of ethambutol hydrochloride during pregnancy. Rabbits receiving high doses of ethambutol hydrochloride during pregnancy gave birth to two fetuses with monophthalmia, one with a shortened right forearm accompanied by bilateral wrist-joint contracture and one with hare lip and cleft palate.

#### **Indications and Usage:**

Ethambutol Hydrochloride Tablets are indicated for the treatment of pulmonary tuberculosis. It should not be used as the sole antituberculous drug, but should be used in conjunction with at least one other antituberculous drug. Selection of the companion drug should be based on clinical experience, considerations of comparative safety and appropriate *in vitro* susceptibility studies.

In patients who have not received previous antituberculous therapy, i.e., initial treatment, the most frequently used regimens have been the following:

Ethambutol plus isoniazid

Ethambutol plus isoniazid plus streptomycin.

In patients who have received previous antituberculous therapy, mycobacterial resistance to other drugs used in initial therapy is frequent.

Consequently, in such retreatment patients, ethambutol should be combined with at least one of the second line drugs not previously administered to the patient and to which bacterial susceptibility has been indicated by appropriate *in vitro* studies. Antituberculous drugs used with ethambutol have included cycloserine, ethionamide, pyrazinamide, viomycin, and other drugs. Isoniazid, aminosalicylic acid, and streptomycin have also been used in multiple drug regimens. Alternating drug regimens have also been utilized.

### **Contraindications:**

Ethambutol hydrochloride is contraindicated in patients who are known to be hypersensitive to this drug. It is also contraindicated in patients with known optic neuritis unless clinical judgment determines that it may be used.

### **Precautions:**

The effects of combinations of ethambutol hydrochloride with other antituberculous drugs on the fetus is not known. While administration of this drug to pregnant human patients has produced no detectable effect upon the fetus, the possible teratogenic potential in women capable of bearing children should be weighed carefully against the benefits of therapy. There are published reports of five women who received the drug during pregnancy without apparent adverse effect upon the fetus.

Ethambutol is not recommended for use in pediatric patients under thirteen years of age since safe conditions for use have not been established.

Patients with decreased renal function need the dosage reduced as determined by serum levels of ethambutol, since the main path of excretion of this drug is by the kidneys.

Because this drug may have adverse effects on vision, physical examination should include ophthalmoscopy, finger perimetry and testing of color discrimination. In patients with visual defects such as cataracts, recurrent inflammatory conditions of the eye, optic neuritis, and diabetic retinopathy, the evaluation of changes in visual acuity is more difficult, and care should be taken to be sure the variations in vision are not due to the underlying disease conditions. In such patients, consideration should be given to relationship between benefits expected and possible visual deterioration since evaluation of visual changes is difficult. (For recommended procedures, see next paragraphs under **ADVERSE REACTIONS**.)

As with any potent drug, periodic assessment of organ system functions, including renal, hepatic, and hematopoietic, should be made during long-term therapy.

### **Adverse Reactions:**

Ethambutol may produce decreases in visual acuity which appear to be due to optic neuritis. This effect may be related to dose and duration of treatment. This effect is generally reversible when administration of the drug is discontinued promptly. In rare cases recovery may be delayed for up to one year or more. Irreversible blindness has been reported.

Optic neuropathy including optic neuritis or retrobulbar neuritis occurring in association with ethambutol therapy may be characterized by one or more of the following events: decreased visual acuity, scotoma, color blindness, and/or visual defect. These events have also been reported in the absence of a diagnosis of optic or retrobulbar neuritis.

Patients should be advised to report promptly to their physician any change of visual acuity.

The change in visual acuity may be unilateral or bilateral and hence *each eye must be tested separately and both eyes tested together*. Testing of visual acuity should be performed before beginning ethambutol hydrochloride therapy and periodically during drug administration, except that it should be done monthly when a patient is on a dosage of more than 15 mg per kilogram per day. Snellen eye charts are recommended for testing of visual acuity. Studies have shown that there are definite fluctuations of one or two lines of the Snellen chart in the visual acuity of many tuberculous patients *not* receiving ethambutol.

The following table may be useful in interpreting possible changes in visual acuity attributable to ethambutol.

<b>Initial Snellen Reading</b>	<b>Reading Indicating Significant Decrease</b>	<b>Significant Number of Lines</b>	<b>Decrease Number of Points</b>
20/13	20/25	3	12
20/15	20/25	2	10
20/20	20/30	2	10
20/25	20/40	2	15
20/30	20/50	2	20
20/40	20/70	2	30
20/50	20/70	1	20

In general, changes in visual acuity less than those indicated under "Significant Number of Lines" and "Decrease-Number of Points," may be due to chance variation, limitations of the testing method or physiologic variability. Conversely, changes in visual acuity equaling or exceeding those under "Significant Number of Lines" and "Decrease-Number of Points" indicate need for retesting and careful evaluation of the patient's visual status. If careful evaluation confirms the magnitude of visual change and fails to reveal another cause, ethambutol should be discontinued and the patient reevaluated at frequent intervals. Progressive decreases in visual acuity during therapy must be considered to be due to ethambutol.

If corrective glasses are used prior to treatment, these must be worn during visual acuity testing. During 1 to 2 years of therapy, a refractive error may develop which must be corrected in order to obtain accurate test results. Testing the visual acuity through a pinhole eliminates refractive error. Patients developing visual abnormality during ethambutol treatment may show subjective visual symptoms before, or simultaneously with, the demonstration of decreases in visual acuity, and all patients receiving ethambutol should be questioned periodically about blurred vision and other subjective eye symptoms.

Recovery of visual acuity generally occurs over a period of weeks to months after the drug has been discontinued. Some patients have received ethambutol hydrochloride again after such recovery without recurrence of loss of visual acuity.

Other adverse reactions reported include: anaphylactoid reactions, dermatitis, pruritus and joint pain; anorexia, nausea, vomiting, gastrointestinal upset, abdominal pain; fever, malaise, headache, and dizziness; mental confusion, disorientation and possible hallucinations. Numbness and tingling of the extremities due to peripheral neuritis have been reported infrequently.

Elevated serum uric acid levels occur and precipitation of acute gout has been reported. Pulmonary infiltrates and eosinophilia also have been reported during ethambutol hydrochloride therapy. Transient impairment of liver function as indicated by abnormal liver function tests is not an unusual finding. Since ethambutol is recommended for therapy in conjunction with one or more other antituberculous drugs, these changes may be related to the concurrent therapy.

## Dosage and Administration:

Ethambutol hydrochloride should not be used alone, in initial treatment or in retreatment. Ethambutol hydrochloride should be administered on a once every 24-hour basis only. Absorption is not significantly altered by administration with food. Therapy, in general, should be continued until bacteriological conversion has become permanent and maximal clinical improvement has occurred.

Ethambutol hydrochloride is not recommended for use in pediatric patients under thirteen years of age since safe conditions for use have not been established.

## Initial Treatment:

In patients who have not received previous antituberculous therapy, administer ethambutol hydrochloride 15 mg per kilogram (7 mg per pound) of body weight, as a single oral dose once every 24 hours. In the more recent studies, isoniazid has been administered concurrently in a single, daily, oral dose.

## Retreatment:

In patients who have received previous antituberculous therapy, administer ethambutol hydrochloride 25 mg per kilogram (11 mg per pound) of body weight, as a single oral dose once every 24 hours. Concurrently administer at least one other antituberculous drug to which the organisms have been demonstrated to be susceptible by appropriate *in vitro* tests. Suitable drugs usually consist of those not previously used in the treatment of the patient. After 60 days of ethambutol hydrochloride administration, decrease the dose to 15 mg per kilogram (7 mg per pound) of body weight, and administer as a single oral dose once every 24 hours.

During the period when a patient is on a daily dose of 25 mg/kg, monthly eye examinations are advised.

See Table for easy selection of proper weight-dose tablet(s).

### Weight-Dose Table

#### 15 mg/kg (7 mg/lb) Schedule

<u>Weight Range</u>		<u>Daily Dose</u>
<u>Pounds</u>	<u>Kilograms</u>	<u>In mg</u>
Under 85 lbs	Under 37 kg	500
85-94.5	37-43	600
95-109.5	43-50	700
110-124.5	50-57	800
125-139.5	57-64	900
140-154.5	64-71	1000
155-169.5	71-79	1100
170-184.5	79-84	1200
185-199.5	84-90	1300
200-214.5	90-97	1400
215 and Over	Over 97	1500

#### 25 mg/kg (11 mg/lb) Schedule

Under 85 lbs	Under 38 kg	900
85-92.5	38-42	1000
93-101.5	42-45.5	1100
102-109.5	45.5-50	1200
110-118.5	50-54	1300

119-128.5	54-58	1400
129-136.5	58-62	1500
137-146.5	62-67	1600
147-155.5	67-71	1700
156-164.5	71-75	1800
165-173.5	75-79	1900
174-182.5	79-83	2000
183-191.5	83-87	2100
192-199.5	87-91	2200
200-209.5	91-95	2300
210-218.5	95-99	2400
219 and Over	Over 99	2500

**How Supplied:**

Ethambutol Hydrochloride Tablets, USP are supplied by **State of Florida DOH Central Pharmacy** as follows:

<b>NDC</b>	<b>Strength</b>	<b>Quantity/Form</b>	<b>Color</b>	<b>Source Prod. Code</b>
53808-0977-1	400 mg	30 Tablets in a Blister Pack	WHITE	23155-0101

Store at controlled room temperature 15°-30°C (59°-86°F) [see USP].

**MANUFACTURED BY**

**Heritage**

**COLUMBIA, MD**

This Product was Repackaged By:

**State of Florida DOH Central Pharmacy**

104-2 Hamilton Park Drive

Tallahassee, FL 32304

United States

**Label Image for 400mg**

## ETHAMBUTOL HYDROCHLORIDE

ethambutol hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:53808-0977(NDC:23155-0101)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ETHAMBUTOL HYDROCHLORIDE (UNII: QE4VW5FO07) (ETHAMBUTOL - UNII:8G167061QZ)	ETHAMBUTOL HYDROCHLORIDE	400 mg

### Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SUCROSE (UNII: C151H8M554)	
GELATIN (UNII: 2G86QN327L)	
HYDROXYMETHYL CELLULOSE (UNII: 273FM27VK1)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	

### Product Characteristics

<b>Color</b>	WHITE (WHITE)	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	6 mm
<b>Flavor</b>		<b>Imprint Code</b>	E7
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53808-0977-1	30 in 1 BLISTER PACK		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA016320	07/01/2009	

**Labeler** - State of Florida DOH Central Pharmacy (829348114)

### Establishment

Name	Address	ID/FEI	Business Operations
State of Florida DOH Central Pharmacy		829348114	repack