

## **Mefloquine:**

**Class:** Antimalarial Agent.

### **Indications:**

Treatment of mild-to-moderate acute malarial infections and prevention of malaria caused by *Plasmodium falciparum* (including chloroquine-resistant strains) or *P. vivax*.

**Available dosage form in the hospital:** 250MG TAB.

### **Trade Names:**

### **Dosage:**

-Malaria: Oral (dose expressed as mg of mefloquine hydrochloride):

*-Mild-to-moderate, treatment:* 1250 mg (5 tablets) as a single dose. **Note:** If clinical improvement is not seen within 48-72 hours, an alternative therapy should be used for retreatment.

*-Uncomplicated, treatment (unlabeled dose):* 750 mg (3 tablets) as initial dose, followed 6-12 hours later by 500 mg (2 tablets) (CDC, 2011)

*-Uncomplicated, chloroquine-resistant P. vivax malaria treatment (unlabeled use):* 750 mg (3 tablets) as initial dose, followed 6-12 hours later by 500 mg (2 tablets) with concomitant primaquine (CDC, 2011)

*-Chemoprophylaxis:* 250 mg weekly starting 1 week (CDC, 2012:  $\geq 2$  weeks) before arrival in endemic area, continuing weekly during travel and for 4 weeks after leaving endemic area. **Note:** Prophylaxis may begin 2-3 weeks prior to travel to ensure tolerance.

### **Renal Impairment :**

No dosage adjustment necessary; only a small amount of mefloquine is renally eliminated.

Hepatic impairment: No dosage adjustment provided in manufacture's labeling; however half-life may be prolonged and plasma levels may be higher in patients with hepatic impairment

### **Common side effect:**

Central nervous system: Chills, dizziness, fatigue, fever, headache

Dermatologic: Rash

Gastrointestinal: Vomiting (3%), abdominal pain, appetite decreased, diarrhea, nausea

Neuromuscular & skeletal: Myalgia

**Pregnancy Risk Factor:** B