





www.tajpharma.com

REMATAL® Sodium Solution  
(pentobarbital sodium) Injection, USP  
Pentobarbital sodium injection is subject to control by the  
Federal Controlled Substances Act under DEA schedule II.  
DO NOT USE IF MATERIAL HAS PRECIPITATED

 **REMATAL**<sup>TM</sup>  
Sodium Solution   
PENTOBARBITAL SODIUM INJECTION, USP  
50 mg/ml FOR I.V / I.M USE ONLY

REMATAL® (pentobarbital sodium injection) is a sterile solution for intravenous or intramuscular injection. Each mL contains pentobarbital sodium 50 mg, in a vehicle of propylene glycol, 40%, alcohol, 10% and water for injection, to volume. The pH is adjusted to approximately 9.5 with hydrochloric acid and/or sodium hydroxide.

 **REMATAL**<sup>TM</sup>  
Sodium Solution   
PENTOBARBITAL SODIUM INJECTION, USP  
50 mg/ml FOR I.V / I.M USE ONLY

REMATAL® Phenobarbital Sodium Injection is indicated for the control of status epilepticus of the tonic-clonic (grand mal) type and prevention and treatment of seizures occurring during or following neurosurgery and/or severe head injury.

REMATAL® Phenobarbital Sodium Injection, USP - For direct IV injection

ACTIVE: Pentobarbital Sodium, derivative of Barbituric Acid 50 mg;

PRESERVATIVE: None;

INACTIVES: Propylene Glycol 40%, Alcohol 10% and Water for Injection (pH adjusted to approximately 9.5 with Hydrochloric Acid and/or Sodium Hydroxide.)

STORAGE: Exposure of pharmaceutical products to heat should be minimized. Avoid excessive heat. Protect from freezing. It is recommended that the product be stored at 20°-25°C (68°-77°F), however, brief excursions are permitted between 15°-30°C (59°-86°F). See USP controlled room temperature.

#### HOW SUPPLIED

REMATAL Sodium Solution (pentobarbital sodium injection, USP) is available in the following sizes: 20-mL multiple-dose vial, 1 g per vial (NDC 76478-501-20); and 50-mL multiple-dose vial, 2.5 g per vial (NDC 76478-501-50).

Each mL contains:

Pentobarbital Sodium, derivative of barbituric acid.....50 mg

Propylene glycol.....40% v/v

Alcohol .....10%

Water for Injection.....qs

(pH adjusted to approximately 9.5 with hydrochloric acid and/or sodium hydroxide.)

Vial stoppers are latex free



Food and Drug Administration (FDA) approved .Prescription Only (POM)

A Taj Pharma™ India Product