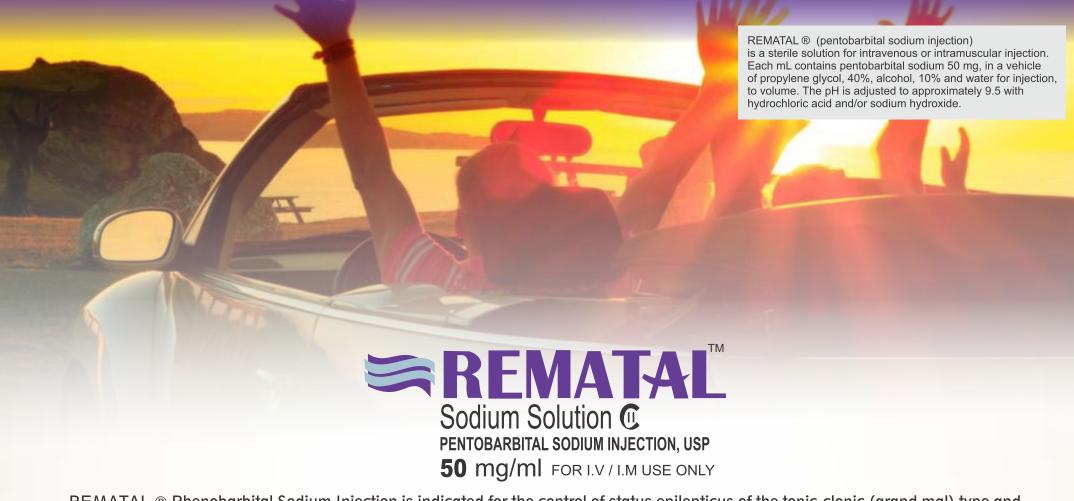
## www.tajpharma.com

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 ® 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:

Lucii iii Comanis.	
Pentobarbital Sodium, derivative of barbituric acid	50 mg
Propylene glycol40	1% v/v
Alcohol	10%
Water for Injection	qs
(pH adjusted to approximately 9.5 with hydrochloric acid and,	
sodium hydroxide.)	
Vial stoppers are latex free	

