

Hepaclin™

L-Ornithine L-Aspartate

Presentation

Hepaclin™ Granules for solution: Each sachet contains L-Ornithine L-Aspartate INN 3 gm.

Hepaclin™ Injection (Infusion Concentrate): Each 10 ml injectable solution contains L-Ornithine L-Aspartate INN 5 gm.

Description

Hepaclin is a stable combination of two important endogenous Amino Acids, L-Ornithine and L-Aspartate. After administration, it quickly breaks down into L-Ornithine and L-Aspartate. L-Ornithine being a substrate of urea cycle, converts toxic ammonia into non-toxic urea which is eliminated via kidneys, helping the diseased liver to carry out its normal function smoothly (detoxification). The process lowers the elevated level of ammonia in blood (hyperammonemia) which is a common problem in most of the liver diseases. L-Aspartate is an essential component of citric acid cycle which liberates energy (ATP), and thus helps in regeneration of damaged liver cells.

Indications

Hepaclin is indicated for the treatment of Hepatic Encephalopathy, Acute Liver Failure, Hyperammonemia due to Acute and Chronic Hepatitis, Fatty Liver, Alcoholic Liver damage, Adjunct to hepatotoxic drugs, Liver Cirrhosis, Post hepatitis convalescence etc.

Dosage and Administration

Sachet: 1-2 sachets of Hepaclin granules is dissolved in a large amount of fluid (e.g. in a glass of water or juice) and is taken orally 3 times a day during or after meals.

Injection (Infusion Concentrate): The recommended dose is up to 20 gm (4 ampoules daily). In case of loss of consciousness (pre-coma) and clouding of consciousness (coma) up to 8 ampoules within 24 hours, depending on the severity of the condition.

Administration

- 1) Hepaclin infusion concentrate can be added to any of the conventional intravenous fluids (0.9% Sodium Chloride, 5% Dextrose and Ringer's lactate etc.)
- 2) The normal dilution rate is 1 ampoule per 100 ml of I.V. fluid. However, the dose should not exceed 6 ampoules per 500 ml infusion. The solution mixtures were stable up to 24 hours.
- 3) Infusion Rate: 5 gm/h at maximum.

Adverse Effects

Very rarely side effects like nausea and vomiting occur. These side effects are usually transient and do not necessitate the withdrawal of the drug.

Contra-Indications

Hepaclin is contraindicated in severe renal insufficiency (serum creatinine value > 3 mg/100 ml).

Precautions

Monitoring of serum and urinary urea levels at regular intervals should be done.

Pregnancy and lactation

The administration in pregnancy and lactation should be avoided.

Drug Interactions

No data regarding the interactions of L-Ornithine L-Aspartate was found.

Storage

Store in a cool and dry place, keep away from light.

Commercial pack

Hepaclin™ Granules for solution: Each box contains 10 sachets of granules for solution.

Hepaclin™ Injection (Infusion Concentrate): Each box contains 1 Ampoule of 10 ml infusion concentrate.

Manufactured by
 **Incepta Pharmaceuticals Ltd**
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V.N.01
HPC