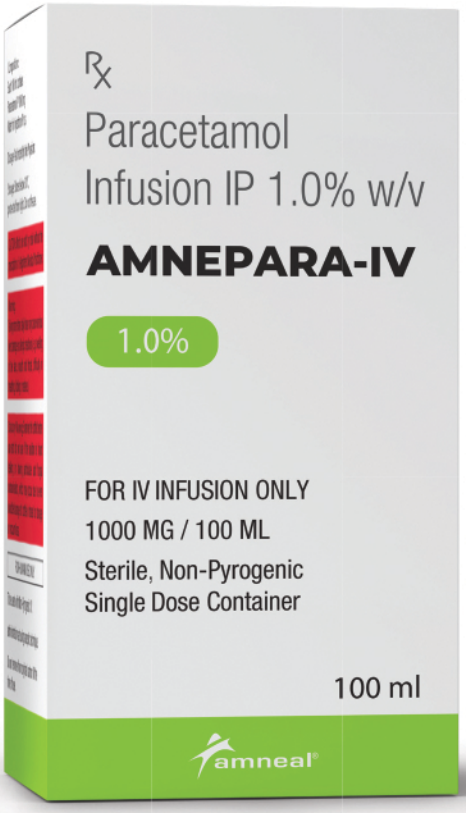
While managing fever & pain in IPD patients*

AMNEPARA-IV®

Paracetamol Infusion IP 1.0% w/v

100 ml



14 USFDA approved facilities across the globe[1]

40+ Global injectable portfolio and expanding[1]

Ref: *As per prescribing information. 1. Data on this page is as of Feb 2025

Abridged Prescribing Information:

AMNEPARA-IV (Paracetamol 2.0% w/v Solution for Infusion (10 mg/mL) (1,000 mg/100 mL)

Presentation: Contains 32.4 mg sodium per 100 mL, which should be taken into account by patients on a low salt diet. Indication: This medicine is an analgesic and an antipyretic. The 100 ml bag is restricted to adults, adolescents and children weighing more than 33 kg. It is indicated for the short-term treatment of moderate pain, especially following surgery, and for the short-term treatment of fever. Dosage and administration: Should not be given more medicine than the label says. Do not exceed the stated dose. Intravenous use. Administered by a healthcare professional by infusion into one of your veins. The dose will be individually adjusted by doctor, based on weight and general condition. Check patient's weight before administration. Contraindications: Allergic (hypersensitive) to paracetamol and propacetamol or to any of the other ingredients of the drug (another analgesic for infusion and a precursor of paracetamol). In a severe liver disease. Warnings & precautions: use a suitable analgesic oral treatment as soon as this administration route is possible. If you suffer from a liver or kidney disease, or from alcohol abuse. If you are taking other medicines containing paracetamol. In cases of nutrition problems (malnutrition) or dehydration. Pregnancy: If necessary, the drug can be used during pregnancy. However, in that case the doctor must evaluate if the treatment is advisable. Patient should use the lowest possible dose that reduces their pain and/or fever and use it for the shortest time possible. Breast-feeding: If necessary, the drug can be used during breast-feeding. Symptoms and treatment of overdosage: a malaise, a drop in blood pressure or changes in laboratory test results: abnormally high levels of hepatic enzymes found during blood checks, a serious skin rash or allergic reaction may occur. Stop the treatment immediately. Changes in laboratory test results have been observed which have necessitated regular blood checks: abnormally low levels of some types of blood cells (platelets, white cells), possibly leading to bleeding from the nose or gums. Cases of redness of the skin, flushing, itching and abnormally rapid beating of the heart. Cases of pain and burning sensation at injection site. Drug interactions: Do not give with any other paracetamol-containing medicine. A dose reduction should be considered for concomitant treatment with Probenecid. - Flucloxacillin (antibiotic), due to a serious risk of blood and fluid abnormality (high anion gap metabolic acidosis) that must have urgent treatment and which may occur particularly in case of severe renal impairment, sepsis (when bacteria and their toxins circulate in the blood leading to organ damage), malnutrition, chronic alcoholism, and if the maximum daily doses of paracetamol are used. Storage: Keep this medicine out of the sight and reach of children. Store below 30°C. Protect from light. Keep the bag in the protective overwrap in order to protect from light and excessive water loss. After opening the overwrapping immediate use is recommended. Amneal Healthcare Private Limited.

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