



Australian Government
Department of Health
 Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: 50310 CEFOXITIN JUNO cefoxitin (as sodium) 1 g powder for injection vial

ARTG entry for Medicine Registered
Sponsor Juno Pharmaceuticals Pty Ltd
Postal Address Level 2 6 Bond Street, South Yarra, VIC, 3141
 Australia
ARTG Start Date 14/10/1994
Product category Medicine
Status Active
Approval area Drug Safety Evaluation Branch

Conditions

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.

Products

1. CEFOXITIN JUNO cefoxitin (as sodium) 1 g powder for injection vial

Product Type	Single Medicine Product	Effective date	8/09/2016
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Permitted Indications

No Permitted Indications included on Record

Indication Requirements

No Indication Requirements included on Record

Standard Indications

No Standard Indications included on Record

Specific Indications

Cefoxitin Sodium for Injection is indicated for the treatment of the following infections when due to susceptible organisms (see Microbiology): peritonitis and other intra-abdominal and intra-pelvic infections, female genital tract infections, septicaemia, endocarditis, urinary tract infections, respiratory tract infections, bone and joint infections, and skin and skin structure infections. Cefoxitin Sodium for Injection has a high degree of stability against beta-lactamase and therefore effective against beta-lactamase producing organisms resistant to penicillins or cephalosporins. It can also be used in mixed infections provided that the organisms are sensitive to it. Cefoxitin Sodium can be used as adjunctive therapy in the surgical treatment of infections including abscesses, infection complicating hollow viscus perforations, cutaneous infections and infections of serous surfaces whether caused by aerobes, mixed aerobes and anaerobes or anaerobes. Cefoxitin Sodium for Injection is also indicated for the prevention of post-operative infections associated with certain surgical procedures of the gastrointestinal, biliary and genital tracts.

Warnings

See Product Information and Consumer Medicine Information for this product

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Vial	Not recorded	3 Years	Store below 25 degrees Celsius	Not recorded	Protect from Light

Pack Size/Poison information

Pack Size	Poison Schedule
5 x 1g vial	(S4) Prescription Only Medicine
10 x 1g vials	(S4) Prescription Only Medicine

Components

1. CEFOXITIN JUNO cefoxitin (as sodium) 1 g powder for injection vial

Dosage Form	Injection, powder for
Route of Administration	Intramuscular Intravenous
Visual Identification	A white to slightly yellow powder

Active Ingredients

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cefoxitin sodium

1.051 g

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