

CEFAZOLIN- cefazolin injection, powder, for solution

WG Critical Care, LLC

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use CEFAZOLIN FOR INJECTION safely and effectively. See full prescribing information for CEFAZOLIN FOR INJECTION.*

CEFAZOLIN for injection, for intravenous use.

Initial U.S. Approval: 1973

RECENT MAJOR CHANGES

Dosage and Administration (2.1) 3/2024

INDICATIONS AND USAGE

Cefazolin for Injection is a cephalosporin antibacterial indicated for perioperative prophylaxis in adult patients (1.1). (1)

To reduce the development of drug-resistant bacteria and maintain the effectiveness of Cefazolin for Injection and other antibacterial drugs, Cefazolin for Injection should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria. (1.2) (1)

DOSAGE AND ADMINISTRATION

- For intravenous infusion only administered over approximately 30 minutes. (2.1)
- Not for intravenous bolus administration or intramuscular administration. (2.1)

Table 1: Recommended Dosage for Perioperative Prophylaxis in Adults with CLcr Equal to 55 mL/min or Greater (2.2) (2)

Dose administered (2) ½ hour to 1 hour prior to the start (2) of surgery (2)	Additional dose during lengthy operative procedures (e.g., 2 hours or more) (2)	Dose for 24 hours postoperatively (2)
1 gram (g) to 2 g (2)	500 mg to 1 g (2)	500 mg to 1 g every 6 hours to 8 hours (2)

- Dosage adjustment is required for adult patients with CLcr that is less than 55 mL/min. (2.3 and 8.6)
- See full prescribing information for preparation and administration instructions. (2.4)

DOSAGE FORMS AND STRENGTHS

- For Injection: 2 grams or 3 grams of cefazolin as a powder in a single-dose vial for reconstitution. (3)

CONTRAINDICATIONS

- Hypersensitivity to cefazolin or other cephalosporin class antibacterial drugs, penicillins, or other beta-lactams (4.1)

WARNINGS AND PRECAUTIONS

- Hypersensitivity Reactions: Cross-hypersensitivity may occur in up to 10% of patients with a history of penicillin allergy. If an allergic reaction occurs, discontinue the drug. (5.1)
- Clostridioides difficile-associated diarrhea (CDAD): May range from mild diarrhea to fatal colitis. Evaluate if diarrhea occurs. (5.3)
- Prothrombin Activity: May be associated with a fall in prothrombin activity. Prothrombin time should be monitored in patients at risk and exogenous vitamin K administered as indicated (5.5)

ADVERSE REACTIONS

- Adult Patients: Most common adverse reactions: gastrointestinal (nausea, vomiting, diarrhea), and allergic reactions (anaphylaxis, urticaria, skin rash). (6)

To report SUSPECTED ADVERSE REACTIONS, contact WG Critical Care, LLC at 1-866-562-4708 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Probenecid: The renal excretion of cefazolin is inhibited by probenecid. Co-administration of probenecid with Cefazolin for Injection is not recommended. (7)

See 17 for PATIENT COUNSELING INFORMATION.

* Pediatric use information is approved for B. Braun Medical's Cefazolin for Injection and Dextrose for Injection. However, due to B. Braun Medical's marketing exclusivity rights, this drug product is not labeled with that pediatric information.

Revised: 3/2024

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

1.1 Perioperative Prophylaxis

Cefazolin for Injection is indicated for perioperative prophylaxis in adult patients [see *Dosage and Administration* (2.1, 2.2, 2.3, 2.4)].

The perioperative use of Cefazolin for Injection is indicated in adult surgical patients in whom infection at the operative site would present a serious risk (e.g., during open-heart surgery and prosthetic arthroplasty).

The prophylactic administration of Cefazolin for Injection preoperatively, intraoperatively and postoperatively may reduce the incidence of certain postoperative infections in patients undergoing surgical procedures which are classified as contaminated or potentially contaminated (e.g., vaginal hysterectomy, and cholecystectomy in high-risk patients such as those older than 70 years, with acute cholecystitis, obstructive jaundice or common duct bile stones).

Pediatric use information is approved for B. Braun Medical's Cefazolin for Injection and Dextrose for Injection. However, due to B. Braun Medical's marketing exclusivity rights, this drug product is not labeled with that pediatric information.

1.2 Usage

To reduce the development of drug-resistant bacteria and maintain the effectiveness of Cefazolin for Injection USP and other antibacterial drugs, Cefazolin for Injection USP should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. When culture and susceptibility information are available, they should be considered in selecting or modifying antibacterial therapy. In the absence of such data, local epidemiology and susceptibility patterns may contribute to the empiric selection of therapy.

2 DOSAGE AND ADMINISTRATION

2.1 Important Administration Instructions

Administer Cefazolin for Injection via intravenous infusion only over approximately 30 minutes. **Not** for intravenous bolus administration or intramuscular administration.

2.2 Dosage for Perioperative Prophylaxis

Dosage for Perioperative Prophylaxis in Adults with Creatinine Clearance (CLcr) Equal to 55 mL/min or Greater

To prevent postoperative infection in contaminated or potentially contaminated surgery, recommended dosages are described in Table 1 below.

Dose administered ½ hour to 1 hour prior to the start of surgery	Additional dose during lengthy operative procedures (e.g., 2 hours or more)	Dose for 24 hours postoperatively
1 gram (g) to 2 g	500 mg to 1 g	500 mg to 1 g every 6 hours to 8 hours

It is important that (i) the preoperative dose be given just prior (1/2 hour to 1 hour) to the start of surgery so that adequate antibacterial concentrations are present in the serum and tissues at the time of initial surgical incision and (ii) cefazolin be administered, if necessary, at appropriate intervals during surgery to provide sufficient concentrations of the antibacterial drug at the anticipated moments of greatest exposure to infective organisms.

The perioperative prophylactic administration of cefazolin should usually be discontinued within a 24-hour period after the surgical procedure. In surgery where the occurrence of infection may be particularly devastating (e.g., open-heart surgery and prosthetic arthroplasty), the prophylactic administration of cefazolin may be continued for 3 to 5 days following the completion of surgery.

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2.3 Preparation of Cefazolin for Injection

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. If particulate matter is evident in reconstituted fluids, the drug solutions should be discarded. Reconstituted solutions may range in color from pale yellow to yellow.

Reconstitution and Dilution

For intravenous infusion, reconstitute Cefazolin for Injection single-dose vials with Sterile Water for Injection according to Table 3 below and *shake well*. After reconstitution further dilute according to Table 3 using the following diluents:

For intermittent or continuous infusion: Dilute reconstituted Cefazolin for Injection in one of the following solutions:

- 0.9% Sodium Chloride Injection, USP
- 5% Dextrose Injection, USP

Discard unused portion.

Cefazolin for Injection Vial Contents	Amount of Sterile Water for Injection for Reconstitution	Approximate Reconstituted Concentrations	Recommended Diluent Volume	Approximate Final Concentrations
2 grams	5 mL	317 mg/mL	50 mL	40 mg/mL
			100 mL	20 mg/mL
3 grams	7.5 mL	319 mg/mL	100 mL	30 mg/mL

Storage of Reconstituted and Diluted Solutions

When reconstituted or diluted according to the instructions above, Cefazolin for Injection is stable for 4 hours at room temperature or for 3 days if stored under refrigeration at 2°C to 8°C (36°F to 46°F).

3 DOSAGE FORMS AND STRENGTHS

Cefazolin for Injection: Supplied as 2 grams or 3 grams of cefazolin as a white to cream powder in a single-dose vial for reconstitution.

4 CONTRAINDICATIONS

4.1 Hypersensitivity to Cefazolin or the Cephalosporin Class of Antibacterial Drugs, Penicillins, or Other Beta-lactams

Cefazolin for Injection is contraindicated in patients who have a history of immediate hypersensitivity reactions (e.g., anaphylaxis, serious skin reactions) to cefazolin or the cephalosporin class of antibacterial drugs, penicillins, or other beta-lactams [see *Warnings and Precautions (5.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity Reactions to Cefazolin, Cephalosporins, Penicillins, or Other Beta-lactams

Serious and occasionally fatal hypersensitivity (anaphylactic) reactions have been reported in patients receiving beta-lactam antibacterial drugs. Before therapy with Cefazolin for Injection is instituted, careful inquiry should be made to determine whether the patient has had previous immediate hypersensitivity reactions to cefazolin, cephalosporins, penicillins, or carbapenems. Exercise caution if this product is to be given to penicillin-sensitive patients because cross-hypersensitivity among beta-lactam antibacterial drugs has been clearly documented and may occur in up to 10% of patients with a history of penicillin allergy. If an allergic reaction to Cefazolin for Injection occurs, discontinue the drug.

5.2 Seizures in Patients with Renal Impairment

Seizures may occur with the administration of Cefazolin for Injection, particularly in patients with renal impairment when the dosage is not reduced appropriately. Discontinue Cefazolin for Injection if seizures occur or make appropriate dosage adjustments in patients with renal impairment [see *Dosage and Administration (2.3)*]. Anticonvulsant therapy should be continued in patients with known seizure disorders.

5.3 *Clostridioides difficile*-associated Diarrhea

Clostridioides difficile-associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including cefazolin for injection, and may range in severity from mild diarrhea to fatal colitis. Treatment with antibacterial agents alters the normal flora of the colon leading to overgrowth of *C. difficile*.

C. difficile produces toxins A and B, which contribute to the development of CDAD. Hypertoxin-producing isolates of *C. difficile* cause increased morbidity and mortality, as these infections can be refractory to antimicrobial therapy and may require colectomy. CDAD must be considered in all patients who present with diarrhea following antibacterial drug use. Careful medical history is necessary since CDAD has been reported to occur over two months after the administration of antibacterial agents.

If CDAD is suspected or confirmed, ongoing antibacterial drug use not directed against *C. difficile* may need to be discontinued. Appropriate fluid and electrolyte management, protein supplementation, antibacterial drug treatment of *C. difficile*, and surgical evaluation should be instituted as clinically indicated.

5.4 Risk of Development of Drug-resistant Bacteria

Prescribing Cefazolin for Injection in the absence of proven or strongly suspected bacterial infection or a prophylactic indication is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

As with other antimicrobials, prolonged use of Cefazolin for Injection may result in overgrowth of nonsusceptible microorganisms. Repeated evaluation of the patient's condition is essential. Should superinfection occur during therapy, appropriate measures should be taken.

5.5 Prothrombin Activity

Cefazolin for Injection may be associated with a fall in prothrombin activity. Those at risk include patients with renal or hepatic impairment or poor nutritional state, as well as patients receiving a protracted course of antimicrobial therapy, and patients previously stabilized on anticoagulant therapy. Prothrombin time should be monitored in patients at risk and exogenous vitamin K administered as indicated.

5.6 Drug/Laboratory Test Interactions

Urinary Glucose

The administration of cefazolin may result in a false-positive reaction with glucose in the urine when using glucose tests based on Benedict's copper reduction reaction that determine the amount of reducing substances like glucose in the urine. It is recommended that glucose tests based on enzymatic glucose oxidase reactions be

used.

Coombs' Test

Positive direct Coombs' tests have been reported during treatment with cefazolin. In hematologic studies or in transfusion cross-matching procedures when antiglobulin tests are performed on the minor side or in Coombs' testing of newborns whose mothers have received cephalosporin antibacterial drugs before parturition, it should be recognized that a positive Coombs' test may be due to the drug.

6 ADVERSE REACTIONS

The following serious adverse reactions to cefazolin for injection are described below and elsewhere in the labeling:

- Hypersensitivity Reactions to Cefazolin, Cephalosporins, Penicillins, or Other Beta-lactams [*see Warnings and Precautions (5.1)*]
- Seizures in Patients with Renal Impairment [*see Warnings and Precautions (5.2)*]
- *Clostridioides difficile*-associated Diarrhea [*see Warnings and Precautions (5.3)*]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The following adverse reactions were reported from clinical trials:

Gastrointestinal: Diarrhea, oral candidiasis (oral thrush), mouth ulcers, vomiting, nausea, stomach cramps, epigastric pain, heartburn, flatus, anorexia and pseudomembranous colitis. Onset of pseudomembranous colitis symptoms may occur during or after antibacterial treatment [*see Warnings and Precautions (5.3)*].

Allergic: Anaphylaxis, eosinophilia, urticaria, itching, drug fever, skin rash, Stevens-Johnson syndrome.

Hematologic: Neutropenia, leukopenia, thrombocytopenia, thrombocythemia.

Hepatic: Transient rise in SGOT, SGPT, and alkaline phosphatase levels has been observed. Reports of hepatitis have been received.

Renal: Reports of increased BUN and creatinine levels, as well as renal failure, have been received.

Local Reactions: Instances of phlebitis have been reported at site of injection. Some induration has occurred.

Other Reactions: Pruritus (including genital, vulvar and anal pruritus, genital moniliasis, and vaginitis). Dizziness, fainting, lightheadedness, confusion, weakness, tiredness, hypotension, somnolence and headache.

Pediatric use information is approved for B. Braun Medical's Cefazolin for Injection and Dextrose for Injection. However, due to B. Braun Medical's marketing exclusivity rights, this drug product is not labeled with that pediatric information.

6.2 Postmarketing Experience

The following adverse reactions have been identified during post approval use of cefazolin. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Immune system disorders: Serum sickness-like reaction

Renal and urinary disorders: Acute tubulointerstitial nephritis (ATIN)

Skin and subcutaneous tissue disorders: Acute generalized exanthematous pustulosis (AGEP)

6.3 Cephalosporin-class Adverse Reactions

In addition to the adverse reactions listed above that have been observed in patients treated with cefazolin, the following adverse reactions and altered laboratory tests have been reported for cephalosporin-class antibacterials: Stevens-Johnson syndrome, erythema multiforme, toxic epidermal necrolysis, renal impairment, toxic nephropathy, aplastic anemia, hemolytic anemia, hemorrhage, a fall in prothrombin activity, hepatic impairment including cholestasis, and pancytopenia.

7 DRUG INTERACTIONS

The renal excretion of cefazolin is inhibited by probenecid. Co-administration of probenecid with Cefazolin for Injection is not recommended.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Available data from published prospective cohort studies, case series and case reports over several decades with cephalosporin use, including cefazolin, in pregnant women have not established a drug-associated risk of major birth defects, miscarriage, or adverse maternal or fetal outcomes. Cefazolin crosses the placenta.

Animal reproduction studies with rats, mice and rabbits administered cefazolin during organogenesis at doses 1 to 3 times the maximum recommended human dose (MRHD) did not demonstrate adverse developmental outcomes. In rats subcutaneously administered cefazolin prior to delivery and throughout lactation, there were no adverse effects on offspring at a dose approximately 2 times the MRHD (*see Data*).

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Data

Human Data

While available studies cannot definitively establish the absence of risk, published data from case-control studies and case reports over several decades have not identified an association with cephalosporin use during pregnancy and major birth defects, miscarriage, or other adverse maternal or fetal outcomes. Available studies have methodologic limitations, including small sample size, retrospective data collection, and inconsistent comparator groups.

Animal Data

Reproduction studies have been performed in rats, mice and rabbits administered cefazolin during organogenesis at doses of 2000, 4000 and 240 mg/kg/day (approximately 1 to 3 times the maximum recommended human dose on a body surface area comparison). There was no evidence of any adverse effects on embryofetal development due to cefazolin. In a peripostnatal study in rats, cefazolin administered subcutaneously up to 1200 mg/kg/day (approximately 2 times the MRHD based on body surface area comparison) to pregnant dams prior to delivery and through lactation caused no adverse effects on offspring.

8.2 Lactation

Data from published literature report that cefazolin is present in human milk but is not expected to accumulate in a breastfed infant. There are no data on the effects of cefazolin on the breastfed child or on milk production.

The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for Cefazolin for Injection and any potential adverse effects on the breastfed child from Cefazolin for Injection or from the mother's underlying condition.

8.4 Pediatric Use

Safety and effectiveness of Cefazolin for Injection for perioperative prophylaxis have not been established for pediatric patients less than 10 years or weighing less than 50 kg.

The safety and effectiveness of Cefazolin for Injection in premature infants and neonates have not been established and is not recommended for use in this age group of pediatric patients.

Pediatric use information is approved for B. Braun Medical's Cefazolin for Injection and Dextrose for Injection. However, due to B. Braun Medical's marketing exclusivity rights, this drug product is not labeled with that pediatric information.

8.5 Geriatric Use

Of the 920 subjects who received cefazolin in clinical studies, 313 (34%) were 65 years and over, while 138 (15%) were 75 years and over. No overall differences in safety or effectiveness were observed between these subjects and younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients, but greater sensitivity of some older individuals cannot be ruled out.

This drug is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in

dose selection, and it may be useful to monitor renal function [see *Dosage and Administration (2.3) and Warnings and Precautions (5.2)*].

8.6 Renal Impairment

When Cefazolin for Injection is administered to adult patients with low urinary output because of impaired renal function (creatinine clearance less than 55 mL/min. for adult patients), lower daily dosage is required [see *Dosage and Administration (2.3) and Warnings and Precautions (5.2)*].

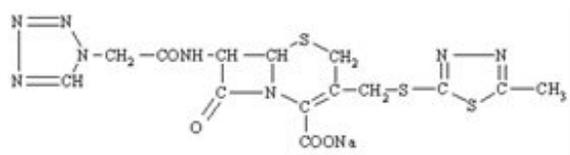
10 OVERDOSAGE

Accidental overdosage resulting in seizures may occur in patients with renal impairment who receive doses greater than the recommended dosage of Cefazolin for Injection [see *Warnings and Precautions (5.2)*]. If seizures associated with accidental overdosage occur, discontinue Cefazolin for Injection and give supportive treatment.

11 DESCRIPTION

Cefazolin for Injection USP is a semi-synthetic cephalosporin for parenteral administration. It is the sodium salt of 3-[[[(5-methyl-1,3,4-thiadiazol-2-yl)thio]-methyl]-8-oxo-7-[2-(1H-tetrazol-1-yl)acetamido]-5-thia-1-azabicyclo [4.2.0]oct-2-ene-2-carboxylic acid.

Structural Formula:



$C_{14}H_{13}N_8NaO_4S_3$ M.W. 476.5

Cefazolin for Injection is supplied as a sterile powder in single-dose vials.

The 2 g/vial Cefazolin for Injection contains 2 grams of cefazolin (equivalent to 2.097 g of cefazolin sodium). The 3 g/vial Cefazolin for Injection contains 3 grams of cefazolin (equivalent to 3.144 grams of cefazolin sodium).

Each vial contains 48 mg of sodium per gram of cefazolin sodium.

After reconstitution with sterile water for injection, the drug product solution has a pH of 4.0 to 6.0.

Cefazolin for Injection is intended for intravenous infusion.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Cefazolin is an antibacterial drug [see *Microbiology (12.4)*].

12.2 Pharmacodynamics

The pharmacokinetic/pharmacodynamic relationship for cefazolin has not been evaluated in patients.

12.3 Pharmacokinetics

Studies have shown that following intravenous administration of cefazolin to normal volunteers, mean serum concentrations peaked at approximately 185 mcg/mL and were approximately 4 mcg/mL at 8 hours for a 1 gram dose.

In a study of constant intravenous infusion with dosages of 3.5 mg/kg for 1 hour (approximately 250 mg) and 1.5 mg/kg the next 2 hours (approximately 100 mg) in healthy volunteers, cefazolin serum concentrations at the third hour were approximately 28 mcg/mL.

Plasma pharmacokinetic parameters of cefazolin in healthy volunteers (N=12) following a single 15-minute IV infusion of 2 grams of Cefazolin for Injection are summarized in Table 4.

	N	C_{max} (mcg/mL)	T_{max}* (h)	AUC_{0-inf} (mcg*h/mL)	t_{1/2} (h)	CL (L/h)	V_z (L)
Single 2 grams Dose as a 15-Minute IV Infusion	12	280.9 (45.9)	0.25 (0.25-0.33)	509.9 (89.3)	2.01 (0.28)	4.03 (0.68)	11.50 (1.53)

*T_{max} reported as median (range)

N= number of subjects observed; C_{max} = maximum plasma concentration; T_{max} = time to maximum plasma concentration; AUC_{0-inf} = area under the plasma concentration-time curve extrapolated to infinity; t_{1/2} = apparent plasma terminal elimination half-life; CL = total clearance; V_z = volume of distribution

Studies in patients hospitalized with infections indicate that cefazolin produces mean peak serum concentrations approximately equivalent to those seen in normal volunteers.

Distribution

Bile concentrations in patients without obstructive biliary disease can reach or exceed serum concentrations by up to five times; however, in patients with obstructive biliary disease, bile concentrations of cefazolin are considerably lower than serum concentrations (less than 1 mcg/mL).

In synovial fluid, the cefazolin concentration becomes comparable to that reached in serum at about 4 hours after drug administration.

Studies of cord blood show prompt transfer of cefazolin across the placenta. Cefazolin

is present in very low concentrations in the milk of nursing mothers.

Elimination

The serum half-life for cefazolin is approximately 1.8 hours following IV administration.

Excretion

Cefazolin is excreted unchanged in the urine. In the first 6 hours approximately 60% of the drug is excreted in the urine and this increases to 70% to 80% within 24 hours.

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12.4 Microbiology

Mechanism of Action

Cefazolin is a bactericidal agent that acts by inhibition of bacterial cell wall synthesis.

Resistance

Predominant mechanisms of bacterial resistance to cephalosporins include the presence of extended-spectrum beta-lactamases and enzymatic hydrolysis.

Antimicrobial Activity

Cefazolin has been shown to be active against most isolates of the following microorganisms, both *in vitro* and in clinical infections [see *Indications and Usage (1)*]:

Aerobic bacteria

Gram-Positive Bacteria

Staphylococcus aureus
Staphylococcus epidermidis
Streptococcus agalactiae
Streptococcus pneumoniae
Streptococcus pyogenes

Methicillin-resistant staphylococci are uniformly resistant to cefazolin.

Gram-Negative Bacteria

Escherichia coli
Proteus mirabilis

Most isolates of indole positive *Proteus* (*Proteus vulgaris*), *Enterobacter* spp., *Morganella morganii*, *Providencia rettgeri*, *Serratia* spp., and *Pseudomonas* spp. are resistant to cefazolin.

Susceptibility Testing

For specific information regarding susceptibility test interpretive criteria and associated test methods and quality control standards recognized by FDA for this drug, please see: <https://www.fda.gov/STIC>.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis and Mutagenesis

Mutagenicity studies and long-term studies in animals to determine the carcinogenic potential of Cefazolin for Injection have not been performed.

Impairment of Fertility

Fertility studies conducted in rats subcutaneously administered cefazolin at doses of 2000 mg/kg/day (approximately 3 times the maximum recommended human dose based on body surface area comparison) showed no impairment of mating and fertility.

16 HOW SUPPLIED/STORAGE AND HANDLING

Cefazolin for Injection is available in a single-dose vial containing 2 grams or 3 grams of cefazolin as a white to cream powder for reconstitution. Each vial contains cefazolin sodium equivalent to 2 grams or 3 grams of cefazolin, and is supplied as follows:

2 grams per Single-Dose Vial: NDC 44567-840-01

Carton of 25: NDC 44567-840-25

3 grams per Single-Dose Vial: NDC 44567-845-01

Carton of 25: NDC 44567-845-25

Store Cefazolin for Injection vials at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature]. Protect from Light.

Storage conditions for the reconstituted and diluted solutions are described elsewhere in labeling [see *Dosage and Administration (2.4)*].

17 PATIENT COUNSELING INFORMATION

Serious Allergic Reactions

Advise patients that allergic reactions, including serious allergic reactions could occur and that serious reactions require immediate treatment and discontinuation of Cefazolin for Injection. Patients should report to their health care provider any previous allergic reactions to cefazolin, cephalosporins, penicillins, or other similar antibacterials [see *Warnings and Precautions (5.1)*].

Seizures

Advise patients that seizures could occur with Cefazolin for Injection. Instruct patients to inform a healthcare provider at once of any signs and symptoms of seizures, for immediate treatment, dosage adjustment, or discontinuation of Cefazolin for Injection [see *Warnings and Precautions (5.2)*].

Diarrhea

Advise patients that diarrhea is a common problem caused by antibacterials including cefazolin for injection, which usually ends when the antibacterial is discontinued.

Sometimes after starting treatment with antibacterials, including Cefazolin for Injection, patients can develop watery and bloody stools (with or without stomach cramps and fever) even as late as two or more months after having taken the last dose of the antibacterials. If this occurs, patients should contact a physician as soon as possible [see *Warnings and Precautions (5.3)*].

Antibacterial Resistance

Patients should be counseled that antibacterial drugs, including Cefazolin for Injection should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When Cefazolin for Injection is prescribed to treat a bacterial infection, patients should be told that although it is common to feel better early in the course of therapy, the medication should be taken exactly as directed. Skipping doses or not completing the full course of therapy may (1) decrease the effectiveness of the immediate treatment and (2) increase the likelihood that bacteria will develop resistance and will not be treatable by Cefazolin for Injection or other antibacterial drugs in the future [see *Warnings and Precautions (5.4)*].

Manufactured for:

WG Critical Care, LLC

Paramus, NJ 07652

Made in Italy

PACKAGE/LABEL PRINCIPAL DISPLAY PANEL

NDC 44567-840-01

Cefazolin for Injection, USP 2 grams per vial

For Intravenous Infusion Only.

LOT

EXP

F50000403182

3 44567 84001 4

NDC 44567-840-01 Single-Dose Vial
Discard Unused Portion

**CEFAZOLIN
for Injection, USP**

2 grams per vial

FOR INTRAVENOUS INFUSION ONLY.

Each vial contains cefazolin sodium equivalent to 2 grams of cefazolin. The sodium content is approximately 48 mg (2.1 mEq) per gram of cefazolin sodium.

Sterile, Nonpyrogenic, Preservative free. **Recommended Dosage:** See Prescribing Information. **Before reconstitution store at 20°C to 25°C (68°F to 77°F) (see USP Controlled Room Temperature).** **PROTECT FROM LIGHT.** **After reconstitution:** Stable for 4 hours at room temperature or 3 days if refrigerated 2°C to 8°C (36°F to 46°F). **Preparation:** See Prescribing Information.

Manufactured for: **WG Critical Care, LLC**
Paramus, NJ 07652
Made in Italy

Rx only Rev. 01/2025

PACKAGE/LABEL PRINCIPAL DISPLAY PANEL

NDC 44567-845-01

Cefazolin for Injection, USP 3 grams per vial

For Intravenous Infusion Only.

LOT

EXP

F50000403475



NDC 44567-845-01 Single-Dose Vial Discard Unused Portion

CEFAZOLIN
for Injection, USP

3 grams per vial

FOR INTRAVENOUS INFUSION ONLY.

Each vial contains cefazolin sodium equivalent to 3 grams of cefazolin. The sodium content is approximately 48 mg (2.1 mEq) per gram of cefazolin sodium.

Sterile, Nonpyrogenic, Preservative free. **Recommended Dosage:** See Prescribing Information. **Before reconstitution store at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature]. PROTECT FROM LIGHT. After reconstitution:** Stable for 4 hours at room temperature or 3 days if refrigerated 2°C to 8°C (36°F to 46°F). **Preparation:** See Prescribing Information.

Manufactured for: **WG Critical Care, LLC**
Paramus, NJ 07652
Made in Italy

Rx only Rev. 03/2025



CEFAZOLIN

cefazolin injection, powder, for solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:44567-840
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CEFAZOLIN SODIUM (UNII: P380M0454Z) (CEFAZOLIN - UNII:IHS69LOY4T)	CEFAZOLIN	2 g

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
Does not contain	NATURAL LATEX RUBBER (UNII: 2LQ0UUW8IN)	0

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44567-840-25	25 in 1 CARTON	05/18/2023	
1	NDC:44567-840-01	1 in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA211413	05/18/2023	

CEFAZOLIN

cefazolin injection, powder, for solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:44567-845
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CEFAZOLIN SODIUM (UNII: P380M0454Z) (CEFAZOLIN - UNII:IHS69LOY4T)	CEFAZOLIN	3 g

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
Does not contain	NATURAL LATEX RUBBER (UNII: 2LQ0UUW8IN)	0

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44567-845-25	25 in 1 CARTON	05/10/2024	
1	NDC:44567-845-01	1 in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA211413	05/10/2024	

Labeler - WG Critical Care, LLC (829274633)

Revised: 3/2024

WG Critical Care, LLC