

CEFUROXIME

BRAND NAME	Cefuroxime SXP
DRUG CLASS	Cephalosporin antibiotic
AVAILABILITY	Vial contains 750 mg of cefuroxime as cefuroxime sodium. ¹ Contains 1.77 mmol (40.65 mg) of sodium. ¹
WARNINGS	Contraindicated in patients with severe immediate (IgE mediated) or severe delayed (T-cell mediated) hypersensitivity to penicillins. Seek specialist advice for patients with non-severe immediate hypersensitivity to penicillins.
pH	6–8.5 when reconstituted ²
PREPARATION	For IM injection: reconstitute the vial with 3 mL of water for injections. Shake gently to form an off-white suspension. The concentration is approximately 216 mg/mL. ³ For IV injection: reconstitute the vial with 6 mL of water for injections. The solution is clear and colourless to yellow. The concentration is approximately 116 mg/mL. ^{1,3} Powder volume – 0.45 mL. ^{1,4}
STABILITY	Vial: store below 25 °C. Protect from light. ¹ Reconstituted solution: stable for 5 hours at 25 °C or 24 hours at 2 to 8 °C. ¹ Infusion solution: stable for 6 hours at 25 °C or 24 hours at 2 to 8 °C. ¹
ADMINISTRATION	
IM injection	Not recommended. ¹ Suitable for doses up to 750 mg only, for higher doses use the IV route. ⁵
SUBCUT injection	Not recommended
IV injection	Inject over 3 to 5 minutes. ¹
IV infusion	Dilute the dose in 50–100 mL of a compatible fluid and infuse over 30 to 60 minutes. ¹
COMPATIBILITY	
Fluids	Glucose 5%, ^{1,3} glucose and sodium chloride solutions, ^{1,3} Hartmann's, ^{1,3} Ringers, ^{1,3} sodium chloride 0.9% ^{1,3}
Y-site	No information
INCOMPATIBILITY	
Fluids	No information
Drugs	No information
SPECIAL NOTES	May cause injection site reactions such as pain and thrombophlebitis. ¹

REFERENCES

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3. Cefuroxime sodium for injection 750mg. Summary of product characteristics. Dublin, Ireland: Flynn Pharma. Approved 01/06/2005. Updated 16/08/2018. Available from www.medicines.org.uk. Accessed 29/04/2021.
4. Scientific affairs. Cefuroxime SXP 750mg injection [email]. Hawthorn, Vic.: Southern Cross Pharma; 25/05/2021.
5. Joint Formulary Committee. BNF 80 (British National Formulary). Basingstoke, UK: Pharmaceutical Press; September 2020-March 2021.