Vancolon

Sterile vancomycin HCl injection

Varicolon is a white to off-white lyophilized powder which makes a clear solution with a pH range of 2.5 to 4.5 when reconstituted in water.

Composition

1 vial contains sterile vancomycin HCl (USP) equivalent to 0.5g/ 1g of vancomycin.

Properties

Vancomycin HCI is a purified tricyclic glycopeptide antibiotic derived from Norcardia orientalis. It is effective for the treatment of serious or severe infections especially those caused by susceptible strains of methicillinresistant (beta-lactam resistant) Staphylococci.

Indications

Microbiology

It is active against many Gram-positive organisms including Staphylococci, group A beta-hemolytic Streptococci including Streptococcus pnęumoniae, Enterococci, Corynebacterium, and Clostridium spp. including C. difficile.

Indications

Vancolon is effective in the treatment of:

- Infections due to Staphylococci including staphylococcal endocarditis, septicemia, bone infections, lower respiratory tract infections, and skin and skin structure infections.
- Hypersensitive patients, who can not tolerate or who have failed to respond to other drugs including the penicillins or cephalosporins, and for infections caused by vancomycin susceptible organisms that are resistant to other antimicrobial drugs.

Note: The parenteral form of vancomycin may be administered orally for the treatment of antibiotic-associated pseudomembranous colitis caused by *C. difficile* and for staphylococcal enterocolitis.

Dosage

Adults: The usual daily intravenous dose is 2g divided either into 0.5g every 6 hours or 1g every 12 hours. Each dose should be administered at no more than 10mg/min or over a period of at least 60 minutes, whichever is longer. Other patient factors, such as age or weight, may call for modification of the usual intravenous daily dose.

Children: The usual intravenous dosage of **Vancolon** is 10mg/kg given every 6 hours. Each dose should be administered over a period of at least 60 minutes.

Infants and Neonates: In neonates and young infants, an initial dose of

15mg/kg is suggested, followed by 10mg/kg every 12 hours for neonates in the 1st week of life and every 8 hours thereafter up to the age of 1 month. Each dose should be administered over 60 minutes. Close monitoring of serum concentrations of vancomycin may be warranted in these patients.

Patients with Impaired Renal Function and Elderly Patients: Dosage adjustment must be made in patients with impaired renal function. In premature infants and the elderly, greater dosage reductions than expected may be necessary because of decreased renal function. Measurement of vancomycin serum concentrations can be helpful in optimizing therapy, especially in seriously ill patients with changing renal function.

A recommended regimen is as follows:

Creatinine Clearance mL/min	Vancomycin Dose mg/24h
100	1,545
90	1,390
80	1,235
70	1,080
60	925
50	770
40	620
30	465
20	310
10	155

The initial dose should be not less than 15mg/kg, even in patients with mild to moderate renal insufficiency.

The table is not valid for functionally anephric patients. For such patients, an initial dose of 15mg/kg of body weight should be given to achieve prompt therapeutic serum concentrations. The dose required to maintain stable concentrations is 1.9mg/kg/24h. In patients with marked renal impairment, it may be more convenient to give maintenance doses of 250 to 1,000mg once every several days rather than administering the drug on a daily basis. In anuria, a dose of 1,000mg every 7 to 10 days has been recommended.

If a dose is missed

- . Give the missed dose as soon as possible.
- If it is nearly time for your next dose, wait until then to give the medicine and skip the missed dose.
- . Do not give two doses at one time

Reconstitution and Administration

Vancolon is administered by slow IV infusion for the treatment of

systemic infections. The drug is very irritating to tissue and must not be given IM. Vancolon is reconstituted by adding 10 or 20mL of sterile water for injection to a vial-labeled as containing 0.5g or 1g of vancomycin, respectively, to provide solutions containing 50mg of the drug per mL. For intermittent IV infusion, the reconstituted solutions containing 0.5g or 1g must be diluted further with at least 100mL or at least 200mL, respectively. of a compatible IV solution and administered over a period of at least 1 hour. When intermittent IV infusion is not feasible, Vancolon may be administered by continuous infusion. In this method, 1-2g of reconstituted Vancolon may be added to a sufficient volume of 0.9% sodium chloride or 5% dextrose injection to permit administration of the desired daily dosage over a 24 hour period. Concentrations of no more than 5mg/mL and rates of no more than 10mg/min are recommended in adults. In selected patients in need of fluid restriction, a concentration up to 10mg/mL may be used. Use of such higher concentrations may increase the risk of infusion-related events that may occur, however, at any rate or concentration.

Note: After reconstitution, solutions retain their potency for 24 hours at room temperature and 96 hours if refrigerated.

Contraindication

Vancomycin is contraindicated in patients with known hypersensitivity to this antibiotic.

Precautions

Rapid bolus administration (e.g., over several minutes) may be associated with exaggerated hypotension and rarely cardiac arrest. Vancomycin should be administered in a dilute solution over a period of not less than 60 minutes to avoid such complications. Stopping the infusion usually results in prompt cessation of these reactions.

Vancomycin should be used with caution in patients with renal insufficiency because the risk of toxicity is appreciably increased by high, prolonged blood concentrations.

Vancomycin should be given to a pregnant woman only if clearly needed. Vancomycin HCl' is excreted in human milk. Caution should be exercised when vancomycin is administered to a nursing woman. Because of the potential for adverse events, a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother.

Monitoring of vancomycin serum concentrations is recommended in premature neonates and young infants. Elderly patients are more likely to have an age related decrease in renal function, which may require dosage adjustment to avoid excessive vancomycin serum concentrations.

Side Effects

Rapid infusion rates of vancomycin have been associated with anaphylactoid reactions including hypotension, wheezing, dyspnea, urticaria, or pruritus. These reactions usually resolve within 20 minutes in most cases.

Nephrotoxicity and ototoxicity have been reported in rare cases, mostly in predisposed patients or those taking concomitantly aminoglycosides or ototoxic drugs, respectively.

Hematopoietic reactions including neutropenia, thrombocytopenia and reversible agranulocytosis have been rarely reported.

Drug Interactions

Concomitant administration of vancomycin and anesthetic agents has been associated with erythema, histamine-like flushing, and anaphylactoid reactions.

Concurrent and/or sequential systemic or topical use of other potentially neurotoxic and/or nephrotoxic drugs such as amphotericin B, aminoglycosides, bacitracin, polymyxin B, colistin, viomycin, or cisplatin, when indicated, requires careful monitoring.

Presentation

Vancolon sterile powder for injection is available in vials containing 0.5g or 1g vancomycin.

* Store at room temperature not exceeding 25°C, away from heat and light.

THIS IS A MEDICAMENT

- Medicament is a product which affects your health, and its consumption contrary to instructions is dangerous for you.
- Follow strictly the doctor's prescription, the method of use and the instructions of the pharmacist who sold the medicament.
- The doctor and the pharmacist are experts in medicines, their benefits and risks.
- Do not by yourself interrupt the period of treatment prescribed for you.
- Do not repeat the same prescription without consulting your doctor.
- Keep all medicaments out of the reach of children.

Council of Arab Health Ministers, Union of Arab Pharmacists.

Any Information ? Call Our Toll Free No. (971) 800-4994



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