

Meropenem & Vaboractam (Vaborem®)

SALAD Contains a PENICILLIN-like structure May be appropriate in penicillin-allergic patient. Refer to CUH Antimicrobial Guidelines on Eolas for further information before administration		
Restricted Antimicrobial Please contact Microbiology/ID/Antimicrobial pharmacist for further information		
Form	Vial contains meropenem 1g and vaboractam 1g Powder for concentrate for solution for infusion Prescribed as combination i.e. 1g/1g, 2g/2g etc	
Reconstitution	Reconstitute each 1g/1g vial with 20mL sodium chloride 0.9% Mix gently Final volume 21.3mL Dilute further prior to administration Use immediately once reconstituted	
Compatibility & Stability	Sodium chloride 0.9% only	
Administration	 IV infusion only Add required dose to 250ml sodium chloride 0.9% infusion bag. Administer over 3 hours 	
	Dose of Meropenem/Vaboractam	Volume of reconstituted injection
	2g/2g 1g/1g 0.5g/0.5g	42.6 mL (two vials) 21.3 mL(one vial) 10.5 ml (half vial)
Monitoring	Monitor: for hypersensitivity and infusion site reactions. Monitor LFTs during treatment due to the risk of hepatotoxicity.	
Adverse reactions	Hypersensitivity reaction (in particular if patient is penicillin allergic), Infusion site phlebitis, pyrexia, hypokalaemia, hypoglycaemia, hypotension, headache, diarrhoea, nausea and vomiting.	
Additional Information	Decreases in blood levels of valproic acid have been reported when it is co- administered with carbapenem agents resulting in a 60-100 % decrease in valproic acid levels in about two days. In exceptional circumstances, where treatment options are extremely limited for a patient, following discussion with Microbiology/Infectious Diseases consultant, a carbapenem may be considered the only/best available treatment option In this case, the consultant with primary responsibility for the patient may decide to proceed with carbapenem treatment for a patient on sodium valproate treatment based on a risk/benefit analysis and following consultation with a consultant neurologist Consultant neurologist advice should be sought regarding the potential requirement for adjunct anticonvulsant therapy if the indication for valproate use is seizure control, and advice on clinical monitoring and therapeutic drug monitoring of anticonvulsant drug serum concentrations Information provided relates to Valorem® (Menarini)	

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