Perth Children's Hospital

Children's Antimicrobial Management Program (ChAMP)

MONOGRAPH				
Inhaled Vancomycin Monograph - Paediatric				
Scope (Staff):	Clinical Staff – Medical, Nursing, Pharmacy			
Scope (Area):	Perth Children's Hospital (PCH)			

Child Safe Organisation Statement of Commitment

The Child and Adolescent Health Service (CAHS) commits to being a child safe organisation by meeting the National Child Safe Principles and National Child Safe Standards. This is a commitment to a strong culture supported by robust policies and procedures to ensure the safety and wellbeing of children at CAHS.

This document should be read in conjunction with this **DISCLAIMER**

DESCRIPTION High Risk Drug	 Vancomycin is a glycopeptide antibiotic. (1-3) It inhibits bacterial cell wall synthesis by preventing the formation of peptidoglycan polymers, alters the bacterial cell membrane permeability and inhibits RNA synthesis. (4-6) Vancomycin is a <u>High Risk Medicine</u>. 		
INDICATIONS AND RESTRICTIONS	Inhaled vancomycin is used in the eradication and treatment of Staphylococcus aureus from the respiratory tract in patients with cystic fibrosis. (3)		
	Inhaled: Monitored (orange) antibiotic		
	 If the use is consistent with a standard approved indication, this must be communicated by documenting that indication on all prescriptions (inpatient and outpatient). 		
	 The ChAMP team will review if ongoing therapy is required and/or if the order does not meet <u>ChAMP Standard Indications</u>. 		
	 If use is not for a standard approved indication, phone approval must be obtained from ChAMP before prescribing. 		
	 Refer to the <u>IV vancomycin</u> and <u>oral vancomycin</u> monograph for further information on different routes of administration. 		
CONTRAINDICATIONS	Hypersensitivity to vancomycin or any component of the formulation. (4, 6)		
	 A history of significant bronchospasm on inhalation of vancomycin.⁽⁷⁾ 		
PRECAUTIONS	Bronchospasm may occur - ensure a short acting bronchodilator is administered prior to administration of the first and subsequent doses. (3, 7)		

	Care must be taken in patients with a history of hypersensitivity to teicoplanin as cross reactivity has occurred. (3)				
EODMIII ATIONS	Available at PCH:				
FORMULATIONS	500mg vials				
	1gram vials (Only for Pharmacy Compounding Service use)				
	 125mg capsules (for oral administration only- not covered in this monograph) 				
	 250mg capsules (for oral administration only- not covered in this monograph) 				
	Other preparations available:				
	Multiple generic preparations				
DOSAGE	The doses listed below fall within the standard range.				
	 Higher doses may be prescribed for certain situations in consultation with an Infectious Diseases or Clinical Microbiology consultant. 				
	Neonates (less than 30 days of age):				
	Not routinely used in neonates less than 30 days old, contact Infectious Diseases or Clinical Microbiology for advice.				
	Children:				
	Inhalation must only be administered after a short acting bronchodilator				
	 4 years to 18 years: 4mg/kg/dose (to a maximum of 250mg) four times daily for 10 to 14 days. (3, 7-10) 				
	For children less than 4 years old dosing is not established, contact Infectious Diseases or Clinical Microbiology for advice.				
DOSAGE ADJUSTMENT	As systemic absorption of vancomycin following nebulisation is low, dose adjustment in renal or hepatic impairment is not necessary, however caution should still be taken. (8, 11)				
RECONSTITUTION		Τ			
	Vial	Volume of 0.9% sodium chloride	Resulting concentration ⁽¹⁾		
	500mg	10mL	50mg/mL		
	Shake the vial well to ensure the vancomycin has dissolved and allow the bubbles to settle before drawing up the required dose.				
ADMINISTRATION	Administer a short acting bronchodilator prior to each dose of inhaled vancomycin. (3)				
	After reconstitution, draw up the required volume and place in				

	 the nebuliser pot. The patient should be sitting upright or standing and breathing normally through the mouthpiece of the nebuliser. 		
	Refer to <u>Pari Nebuliser Administration</u> Guideline for further information.		
MONITORING	 Measure lung function before and after initial dose of vancomycin and monitor for bronchospasm.⁽³⁾ 		
	 Patients should be monitored for adverse effects as per systemic administration, in particular ototoxicity and nephrotoxicity. Please refer to IV vancomycin monograph for possible systemic side effects. 		
ADVERSE EFFECTS	Common:		
	 Cough, bronchospasm, bitter taste, moderate chest pain after coughing, throat irritation, bronchoconstriction. (7, 12) 		
	Rare:		
	 Studies have shown minimal systemic absorption of vancomycin when administered by inhalation. However, patients should still be monitored for adverse effects as per systemic administration, in particular ototoxicity and nephrotoxicity. 		
COMPATIBLE FLUIDS	Sodium chloride 0.9%		
	Water for injection. (1, 3)		
STORAGE	Vial: Store below 25°C ^(1, 4)		
	For use as an inpatient, a new vial must be used for each dose.		
	 For use at home a vial may be reconstituted each day and the remaining solution stored between 2-8°C for up to 24 hours.^(2, 13) 		
INTERACTIONS	Inhaled vancomycin may interact with other medications; please consult PCH approved references (e.g. Clinical Pharmacology), your ward pharmacist or Pharmacy on extension 63546 for more information.		
	Whilst systemic absorption should be negligible, Vancomycin can cause nephrotoxicity and ototoxicity. Caution should be taken with the concurrent use of nephrotoxic and ototoxic agents. (4, 5)		
MANUFACTURER SAFETY DATA SHEET (SDS)	To access to the Manufacturer SDS for this product, use the following link to ChemAlert.		

Please note: The information contained in this guideline is to assist with the preparation and administration of **inhaled vancomycin. Any variations to the doses recommended should be clarified with the prescriber prior to administration**

Related CAHS internal policies, procedures and guidelines

Antimicrobial Stewardship Policy

ChAMP Empiric Guidelines and Monographs

KEMH Neonatal Medication Protocols

References and related external legislation, policies, and guidelines

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- Clinical Pharmacology [Internet]. Elsvier BV. 2021 [cited 14/01/2021]. Available from: http://www.clinicalpharmacology-

ip.com.pklibresources.health.wa.gov.au/default.aspx.

- 3. Paediatric Formulary Committee. BNF for Children: 2020. London: BMJ Group Pharmaceutical Press: 2021.
- 4. MIMS Australia. MIMS online [full product information]. St Leonards, N.S.W: CMP Medica Australia.; 2020. p. 1v. (various pagings).
- 5. Rossi S, editor. Australian Medicines Handbook. Adelaide, S. Aust.: Australian Medicines Handbook; 2020.
- 6. Taketomo CK, Hodding JH, Kraus DM. Pediatric and Neonatal dosage handbook with international trade names index. 26th edition ed. Ohio: Lexi-comp; 2019-2020.
- 7. Dezube R, Jennings MT, Rykiel M, Diener-West M, Boyle MP, Chmiel JF, et al. Eradication of persistent methicillin-resistant Staphylococcus aureus infection in cystic fibrosis. J Cyst Fibros. 2019;18(3):357-63.
- 8. Hayes et al. Aerosolized Vancomycin for the treatment of MRSA after lung transplantation (abstract) Respirology. 2010;15:182-6.
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- 10. Kiefer A, Bogdan C, Melichar VO. Successful eradication of newly acquired MRSA in six of seven patients with cystic fibrosis applying a short-term local and systemic antibiotic scheme. BMC Pulm Med. 2018;18(1):20.
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- 12. Jennings M, Bucur C, Boyle M, Konstan M. Safety and pharmacokinetics of inhaled vancomycin in individuals with cystic fibrosis. American Cystic Fibrosis Conference; September 2010; Baltimore: Pediatric pulmonology; 2010. p. 320.
- 13. Royal Pharmaceutical Society. Martindale: The Complete Drug Reference. London: The Pharmaceutical Press 2020.

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