Indication¹

PALIVIZUMAB

 Prevention of serious lower respiratory tract disease caused by respiratory syncytial virus (RSV) in children at high risk of RSV disease²

Group	Eligibility criteria ¹			
1	Infants who have severe chronic lung disease of prematurity, defined as supplemental oxygen requirement at term equivalent or on discharge from neonatal unit; and are likely to:			
(a) be on oxygen at the start of the RSV season*; and				
	(b**) be less than one year of age at the start of the RSV season*			
2*	Infants with severe chronic lung disease of different aetiologies and who are on home oxygen and are less than one year of age at the start of the RSV season*			
3**	Infants less than six months with congenital heart disease with significant left to right shunt awaiting surgical correction or who have pulmonary hypertension			
4**	Babies less than one year of age at the start of the RSV season* who are at risk of RSV because of a severe immunodeficiency (congenital, acquired or immunosuppressed) and not on intravenous immunoglobulin			
5**	Children having a bone marrow transplant with active RSV			
Groups 2 to 5 unlikely to be relevant to neonatal population * as determined by local protocols and location ** at the discretion of consultant paediatrician or paediatric cardiologist				

	Presentation	Vial: 50 mg in 0.5 mL 100 mg in 1 mL			
	Dosage ^{3,4}	 15 mg/kg A course of five IM injections at one monthly intervals throughout the RSV season 			
Σ	Preparation⁵	 Not required. Do not dilute⁴. Do not shake the viale⁴ 			
=	Administration	 Draw up prescribed dose Do not shake vial or medicine in syringe⁴ Intramuscular injection into thickest part of the vastus lateralis in the anterolateral thigh Divide volumes greater than 1 mL into 2 (or more) injections^{1,3} and administer into separate sites 			
	 Contraindications Known hypersensitivity to palivizumab or other humanised monoclonal ant Special RSV season may vary by location. Check local protocols Not indicated for children who have had RSV infection¹ Discontinue monthly prophylaxis if breakthrough RSV hospitalisation occurs¹ Second episode unlikely in the same RSV season³ 				
	Monitoring • Not required				
	Compatibility • Do not mix with other medications ⁵				
	Incompatibility • Do not mix with other medications ⁵				
	Interactions • Not applicable				
	Stability	• Store at 2–8 °C ⁴ . Refrigerate ⁴ . Do not freeze ⁴			
	Side effects	 Common: fever, rash, rhinitis, wheeze, cough, diarrhoea, injection site reaction, cyanosis (in children with congenital heart disease)² Rare: anaemia, elevated liver enzymes, hypersensitivity (including anaphylaxis)² 			
	Actions	Humanised managinal antibody that neutralises and inhibits fusion of respiratory			
	Abbreviations ID infectious diseases, IM: intramuscular, RSV: respiratory syncytial virus, QCH: Queensland Children's Hospital,				



Keywords	Monoclonal antibody, antiviral, RSV, respiratory syncytial virus, synagis, home oxygen,
Reywords	respiratory virus, chronic lung disease of prematurity, supplemental oxygen, vaccine

The Queensland Clinical Guideline *Neonatal Medicines* is integral to and should be read in conjunction with this monograph. Refer to the disclaimer. Destroy all printed copies of this monograph after use.

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