BEVACIZUMAB (or biosimilar)

	Indication	 Acute aggressive ROP¹-³ Type 1 ROP¹,³ in posterior zone 2 Any stage in zone 1, 2 or 3 with plus disease, or Stage 3 in zone 1 without plus disease³ If unsuitable for ROP laser treatment³, due to Previous unsuccessful laser attempt³ Inability to tolerate procedure³ Poor view of retina³ 			
OPHTHALMIC	Presentation	Prefilled syringe: 3 mg in 0.12 mL	O		
	Dosage ^{1,2,4,5}	Single dose: 0.313 mg or 0.625 mg	_		
ΙŽ	Preparation	Nil required			
OPH	Administration	 Intravitreal injection (under sterile conditions) Performed by ophthalmologist or ophthalmology fellow ONLY 			
	Special considerations				
	Monitoring	Red reflex check, day after injection Performed by clinician skilled in assessing red reflex			
	Compatibility	Not applicable			
	Incompatibility	patibility • Not applicable			
	Interactions	• Nil known			
	Stability ⁶	• Refrigerate 2–8°C			
Side effects detact		 Ocular: ocular inflammation^{2,6}, ocular infection⁶, ocular haemorrhage⁷, retinal detachment^{2,4,8}, cataract⁷, loss of vision (rare)⁸ Respiratory: apnoea⁶ 			
	Actions	 Anti-VEGF monoclonal antibody^{2,4,8} Blocks VEGF formation and inhibits formation of new blood vessels^{1,2,4,9} 			
	Abbreviations	Anti-VEGF: anti-vascular endothelial growth factor, ROP: retinopathy of prematurity, VEGF: vascular endothelial growth factor			
	Keywords	anti-VEGF, antivascular endothelial growth factor, avastin, bevacizumab, mvasi, neo medicine, neonatal monograph, ROP, retinopathy of prematurity, VEGF	natal		

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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