Midazolam Newborn use only

Alert	High risk medication causing significant patient harm when used in error.					
Indication	Sedation during ventilatio					
	Treatment of refractory seizure.					
Action	Intensify the physiological inhibitory mechanisms mediated by gamma-aminobutyric acid (GABA) by					
	accumulation and occupation of benzodiazepine receptors. Anti-anxiety properties are related to					
	increasing the glycine inhi					
Drug type	Short acting benzodiazepi					
Trade name	Hypnovel, Midazolam Alphapharm,, Midazolam Pfizer,, Midazolam-Baxter, B.Braun Midazolam, Midazolam Accord, Midazolam Apotex.					
Presentation	5 mg/mL, 5mg/5mL, 150r	mg/10mL and 15mg/3m	L ampoules used for IV and oral use			
Dose						
	Method	Dose				
	IV infusion for sedation	0.2–1 microgram/kg/n	ninute			
	IV infusion for seizures	_) microgram/kg over 3–5 minutes 7 microgram/kg/minute			
			e very 2 hours when required			
	IV bolus	(Dose range: 50–150 microgram/kg/dose)				
		50 microgram/kg/dose every 4 hours when required				
	IM injection	(Dose range: 50–150 microgram/kg/dose)				
	Oral	250 microgram/kg as a	a single dose			
	Sublingual	200 microgram/kg as a				
	Intranasal	200 microgram/kg per	-			
		(Dose range: 200–300	microgram/kg/dose)			
Dose adjustment						
Maximum dose						
Total cumulative						
dose Route	IV IN Oral Sublingual In	tranacal				
Preparation	IV, IM, Oral, Sublingual, In	li di lasai.				
Freparation	IV Sedation_using 5 mg/1 mL strength					
	Infusion		Descerite ad any suret			
	Infusion strength		Prescribed amount			
	1 mL/hour = 1 microgram/kg/minute3 mg/kg midazolam and make up to 50 mLDraw up 0.6 mL/kg (3 mg/kg of midazolam) and add glucose 5%, glucose 10% or sodium chloride					
	0.9% to make final volume 50 mL. Infuse at a rate of 1 mL/ hour = 1 microgram/kg/minute.					
	Sedation using 5mg/5 mL strength					
	Infusion strength		Prescribed amount			
	<u>1 mL/hour = 1 microgram/kg/minute</u>		3 mg/kg midazolam and make up to 50 mL			
	Draw up 3 mL/kg (3 mg/kg of midazolam) and add glucose 5%, glucose 10% or sodium					
	to make final volume 50 mL. Infuse at a rate of 1 mL/ hour = 1 microgram/kg/minute.					
	Seizures using 5 mg/1 mL strength					
	Infusion strength		Prescribed amount			
	1mL/hour = 5microgram/kg/minutes		15mg/kg midazolam and make up to 50mL			
	Draw up 3 mL/kg (15 mg/kg of midazolam) and add glucose 5%, glucose 10% or sodium chloride 0 to make final volume 50 mL. Infuse at a rate of 1 mL/hour = 5 microgram/kg/minute.					
	Seizures using 5 mg/5 mL strength (not to be used for babies over 3.3 Kg)					
	Infusions	strength	Prescribed amount			

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Stability	(thiopentone), tramadol, trimethoprim-sulfamet Diluted solution: Store at 2–8°C and use within 2	
Stability		
	I (thiopentone), tramadol, trimethonrim-sultamet	
		-
	piperacillin-tazobactam (EDTA-free), potassium a	
	succinate, imipenem-cilastatin, indomethacin, or	
		de (frusemide), ganciclovir, hydrocortisone sodium
	azithromycin, cefepime, ceftazidime, chloramph	
	Y-site: Fat emulsion. Aciclovir, albumin, aminoph	vlline amoxicillin amnicillin azathionrine
meompationity		
Incompatibility	Fluids: No information.	, vəcurunnunn.
	nitroprusside, tirofiban, tobramycin, vancomycir	
		otassium chloride, ranitidine, remifentanil, sodium
	methadone, methylprednisolone, metronidazole	
		peridol lactate, hydromorphone, labetalol, linezolid,
	clindamycin, digoxin, dopamine, doripenem, ept	-
	bivalirudin, calcium gluconate, caspofungin, cefc	-
	Y-site: Amino acid solutions. Abciximab, amikaci	n amiodarone anidulafungin atracurium
compationity		
Compatibility	Fluids: Glucose 5%, glucose 10%, sodium chlorid	
	Seizure-like myoclonus (more common in prema	ture neonates receiving via intravenous route)
	Nasal discomfort (with intranasal route).	
	Hypersalivation.	
Adverse reactions	Hypotension and reduced cardiac output, particularly when used in combination with fentanyl. Respiratory depression and apnoea.	
Adverse reactions	with adding or withdrawing caffeine or aminoph	
		-
Drug mileractions	Concurrent administration with erythromycin promotes accumulation. Xanthines may decrease the anaesthetic/sedative effect of benzodiazepines. Care needs to be taken	
Drug interactions		
	Rapid IV infusion may result in hypotension, resp	-
	system (CNS) effects; use doses at lower end of t	-
	Caution in neonates with renal and hepatic impa	irment - increased sensitivity to central nervous
	precipitate withdrawal seizures.	mistration as abrupt discontinuation may
	hypotension. Withdraw slowly after chronic adm	
	system depressants and may increase the risk of	
	Caution when concurrently used with opioids – r	
	term neonates up to 22 hours in premature infai	
Precautions		n, midazolam half-life is increased from 4–6 hours in
Contraindications	Known hypersensitivity to midazolam.	
	Level of sedation.	
2	Blood pressure	
Monitoring	Apnoea, respiratory depression	
	IM: Inject deep into a large muscle.	
	hours. May be irritating to nasal mucosa.	
	nostrils over 15 seconds. Absorption is rapid; ma	ximum effect in 10 minutes and duration up to 2
	Intranasal: IV ampoules may be used for intrana	
	Oral: Plastic IV ampoules may be used for oral ac	Iministration.
	IV bolus: slow push over 10 minutes. ⁹	
Administration	IV infusion: continuous infusion via a syringe pump. Change solution every 24 hours.	
	chloride 0.9% to make final volume of 5 mL with	
		nicrogram of midazolam) and add 4 mL of sodium
	chloride 0.9% to make final volume of 10 mL wit	
		microgram of midazolam) and add 9.6 mL of sodium
	IV bolus, IM, oral, sublingual and intranasal	
	0.9% to make final volume 50 mL. Infuse at a rat	
	<u>1mL/hour = 5microgram/kg/minutes</u> Draw up 15 mL/kg (15 mg/kg of midazolam) and	15mg/kg midazolam and make up to 50mL add glucose 5% glucose 10% or sodium chloride

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Storage	Midazolam Apotex, Midazolam-Baxter: Store below 30°C. Protect from light. B. Braun Midazolam, Hypnovel, Midazolam Alphapharm: Store below 25°C. Protect from light.	
	Midazolam Pfizer: Store below 25°C. Protect from light. Unopened ampoules will be suitable for use	
	for up to 8 months after the foil sachet has been opened, if protected from light.	
	Schedule 4D (S4D) medication. Store in dangerous drug safe and record use in S4D register.	
Excipients	Sodium chloride, hydrochloric acid, sodium hydroxide, water for injections.	
Special comments	Flumazenil is a specific benzodiazepine antagonist and may be used (very limited experience in the neonate) to rapidly reverse respiratory depression – 10 microgram/kg/dose IV push.	
	May repeat every minute for up to 4 more doses.	
Evidence	 Efficacy There are insufficient data to promote the use of intravenous midazolam infusion as a sedative for neonates undergoing intensive care. Although all studies included in the review reported better sedation, none of the scales used had been validated in preterm infants and thus the effectiveness could not be evaluated [1] (Level 1, Grade B). Midazolam was effective in neonates with refractory seizures that did not respond to phenobarbital (phenobarbitone), phenytoin or pentobarbital (pentobarbitone) [2] (Level IV, Grade D). 	
	Safety One study showed a statistically significant higher incidence of adverse neurological events (death, grade III or IV IVH, PVL) and meta-analysis of data from two studies showed a statistically significant longer duration of NICU stay in the midazolam group compared to the placebo group [1] (Level1, Grade B).	
	Administration of midazolam in ventilated premature infants causes significant changes in cerebral oxygenation and hemodynamics, which might be harmful [3] (Level III, Grade C). Intravenous bolus doses of midazolam in association with fentanyl should be used with great caution in the newborn, especially if very premature or with unstable blood pressure [4] (Level IV, Grade D). Sedation with midazolam has a transient effect on the background aEEG activity [5] (Level III, Grade C).	
	Pharmacokinetics Midazolam is highly protein bound with an elimination half-life of 4–6 hours in term neonates and a variable half-life (up to 22 hours) in premature neonates and those with impaired hepatic function. Bioavailability is approximately 36% with oral administration and 50% with sublingual and intranasal administration [6] (Level III, Grade C).	
Practice points		
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