Salbutamol

Newborn use only

Alert	Use with caution; safety data in newborn infants are limited.
	Evidence in the treatment of respiratory disease and bronchospasm in neonates is poor.
Indication	Hyperkalaemia
	Bronchospasm (evidence for efficacy is lacking)
Action	Stimulates liver and muscle cyclic AMP production causing potassium flow into cells.
Drug type	Sympathomimetic. β ₂ -agonist.
Trade name	IV: Ventolin Injection
	Inhalation: APO-Salbutamol 2.5, Asmol uni-dose 2.5, Butamol 2.5, Chemmart Salbutamol 2.5,
	Pharmacor Salbutamol 2.5, Salbutamol Actavis 2.5, Salbutamol 2.5, Salbutamol Sandoz 2.5, Salbutamol Sterinebs 2.5, Salbutamol-GA 2.5, Salbutamol-GA 2.5, Ventolin Nebules 2.5
Presentation	IV: 500 micrograms/mL ampoule
riesentation	Inhalation: 1 mg/mL (2.5 mg in 2.5 mL) and 2 mg/mL (5 mg in 2.5 mL) inhalation solution ampoules.
Dose	Intravenous:
2030	4–5 microgram/kg over 20 minutes.
	Monitor serum potassium and heart rate (tachycardia) closely. If potassium critical or continues to rise,
	consider repeating dose every 4 hours or use of other strategy (insulin/glucose; addition of rectal
	cation-resin).
	Inhalation:
	400 microgram via nebulisation. Repeat two-hourly as required and titrated to response [serum
	potassium or respiratory status] and heart rate [tachycardia].
Dose adjustment	
Maximum dose	
Total cumulative	
dose	
Route	IV, inhalation
Preparation	IV:
	Draw up 0.4 mL (200 microgram of salbutamol) and add 19.6 mL of water for injection to make a 10
	microgram/mL solution.
	Inhalation: Draw up 0.4 mL (400 micrograms of salbutamol) from the 1 mg/mL inhalation ampoule and add 1.6 mL
	sodium chloride 0.9% to make a final volume of 2 mL with a final concentration of 0.2mg/mL.
	OR.
	Draw up 0.2 mL (400 micrograms of salbutamol) from the 2 mg/mL inhalation ampoule and add 1.8 mL
	sodium chloride 0.9% to make a final volume of 2 mL with a final concentration of 0.2mg/mL.
Administration	IV: Over 15–20 minutes via syringe driver.
	Inhalation: Via nebuliser over 10 minutes and discard remainder
Monitoring	Cardiac rate and rhythm,
	Serum potassium, blood glucose
Contraindications	
Precautions	Infants with tachycardia
Drug interactions	Non-selective beta-blockers may increase serum potassium.
	Diuretics (hydrochlorothiazide, furosemide) increase risk of hypokalaemia and ECG changes.
	Salbutamol decreases digoxin concentrations.
Adverse reactions	Tachycardia, tremor, hypokalaemia. There is some concern that a transient increase in serum
Overdess	potassium may occur in the first few minutes of treatment.8
Overdose	AUSTRALIA Contact the Poisons Information Contro on 12 11 26 for information on the management of everdose
	Contact the Poisons Information Centre on 13 11 26 for information on the management of overdose. NEW ZEALAND
	Contact the National Poisons Centre on 0800 764 766 for information on the management of overdose.
Compatibility	Fluids: Glucose 5%, sodium chloride 0.9%, glucose 5% in sodium chloride 0.9%,, lactated Ringer's
Compatibility	injection
	PN at Y-site: Compatible with 2 in 1 solution (Amino acid-glucose-trace element mixture). No
	information on lipid emulsion. Y-site: Meropenem, metronidazole, naloxone

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	Inhalation: Sodium chloride 0.9%
Incompatibility	Fluids: No information.
	PN at Y-site: No information on lipid emulsions.
	Y-site: Pantoprazole. Ketamine (variable)
	Inhalation: No information
Stability	IV: Ampoules should be used immediately after opening. Any unused solution should be discarded.
	Diluted solution stable for 24 hours below 25°C.
	Inhalation: Ampoules should be used immediately after opening. Any unused solution should be
Chaman	discarded.
Storage	IV ampoule: Store at room temperature below 30°C. Protect from light.
	Inhalation ampoule: Store at room temperature below 25°C. Protect from light.
Excipients	Cross should the correct strength of calbutamed introvenous and inhalation among les
Special comments	Cross-check the correct strength of salbutamol intravenous and inhalation ampoules.
Evidence	Efficacy and safety Treatment of hyperbolic Acceptance is review identified and study (Circle at al., 2003) of 10 infents.
	Treatment of hyperkalaemia: A systematic review identified one study (Singh et al., 2002) of 19 infants which compared inhaled salbutamol [albuterol] versus placebo for non-oliguric hyperkalaemia (serum
	K ⁺ 5–7.5 mmol/L) in premature newborns. Inhaled salbutamol 400 microgram, repeated 2-hourly as
	required, reduced serum K ⁺ from baseline at 4 hours (mean difference 0.69 mmol/L) and 8 hours
	(mean difference 0.59 mmol/L).¹ All-cause mortality was not reduced and cardiac arrhythmia did not
	occur in either study group. There was no significant difference in severe IVH, tremor, hyperglycaemia
	or pulmonary haemorrhage. ¹
	A number of case reports and case series have been published documenting the efficacy of salbutamol
	by infusion for treatment of hyperkalaemia in the newborn. Greenhough et al reported the use of IV
	salbutamol 4 microgram/kg over 20 minutes in 10 consecutive neonates with hyperkalaemia. ² The
	potassium fell in 7 of the 10 infants (range 0.7–1.8 mmol/L) but continued to rise in 3 infants, all of
	whom had a persistent metabolic acidosis. ²
	Murdoch et al reported on the use of IV salbutamol 4 microgram/kg over 20 minutes in 13 children
	(ages 0.01–16.7 years) with hyperkalaemia. ³ The mean reduction in plasma potassium concentration
	was 1.48 mmol/L at 40 minutes and 1.64 mmol/L at 120 minutes. ³
	was 1.46 minory East 46 minutes and 1.64 minory East 126 minutes.
	Kemper et al reported on the use of IV salbutamol at 5 microgram/kg over 20 minutes in 15 children
	(ages 0.1–16 years) with hyperkalaemia. ⁴ The mean reduction in plasma potassium concentration was
	0.87 mmol/L at 30 minutes and 1.69 mmol/L at 120 minutes. Transient tachycardia was detected in
	three patients. ⁴
	Recommendation: Salbutamol (either inhaled or intravenously administered) may be used in the
	treatment of hyperkalaemia in the neonate. Salbutamol may be useful in settings where
	hypoglycaemia limits the use of insulin. Salbutamol may have additive effects when used with insulin
	and glucose. Salbutamol appears to be generally safe with limited risk of tachycardia. (LOE II – III, GOR
	B).
	Treatment of respiratory disease: Systematic review of 3 trials including 140 infants comparing
	salbutamol versus placebo in near term or term infants less than three days of age with transient
	tachypnoea of the newborn found a reduction in the duration of oxygen therapy (MD -43.10 hours,
	95% CI -81.60 to -4.60), but no difference in the need for CPAP, mechanical ventilation or duration of
	hospital stay and tachypnoea. At present there is insufficient evidence to determine the efficacy and
	safety of salbutamol in the management of transient tachypnoea of the newborn. ⁵
	Systematic review ⁶ found a single study that reported prophylaxis of preterm infants at risk of chronic
	lung disease with salbutamol led to no difference in mortality (RR 1.08, 95% CI 0.50 to 2.31) or CLD (RR
	1.03, 95% CI 0.78 to 1.37). There is no evidence for the use of salbutamol for prevention of chronic lung
	disease. ⁶
	Recommendation: There is insufficient evidence to recommend use of nebulised salbutamol in
	newborn infants with respiratory disease. (LOE I GOR C)

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	Pharmacokinetics		
	Reports describing the pharmacokinetics of intravenous salbutamol in neonates and children are		
	limited. Kirpalani et al studied the pharmacokinetics of a single dose of intravenous salbutamol in six		
	preterm infants (GA 24 to 28 weeks), postnatal age 54 to 105 days, with bronchopulmonary dysplasia. ⁷ The elimination half-life of salbutamol was 118 minutes (range 69 to 162 minutes), volume of		
	distribution was 1291 mL/kg (range 246 to 2997) and clearance 7.5 mL/kg/min (range 2.46 to 20.1).		
	The authors noted that the elimination half-life in their neonates was slightly shorter than that of		
	healthy adults. ⁷		
Practice points			
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