Adrenaline (epinephrine) nebulised

Newborn use only

Alert	Adrenaline 1:1000 strength should be used for nebulisation.	
Indication	Management of post-extubation stridor [evidence for effectiveness is not clear]. [1-4]	
muication		
	Initial treatment of outpatients with moderate to severe bronchiolitis. [5] Initial treatment of croup.[6]	
Action	Sympathomimetic catecholamine with alpha and beta adrenergic actions. Vasoconstrictor. It	
Action	also induces relaxation of the bronchial smooth muscle by acting on beta-adrenergic	
	receptors to alleviate wheezing and dyspnoea.	
	The effects of nebulised adrenaline for the treatment of croup lasts for 2–3 hours.	
Drug type	Sympathomimetic catecholamine. Inotropic vasopressor.	
Drug type Trade name		
	Adrenaline 1:1,000 Adrenaline Acid Tartrate injection.	
Presentation	1 mg/mL or 1:1,000 ampoule [1000 microgram/mL]	
Dose	0.5 mg/kg (0.5 mL/kg of adrenaline 1:1000 ampoule)	
	Dose may be repeated every 60 minutes if required following medical assessment of previous	
	dose effect.	
Dose adjustment	Not applicable.	
Maximum dose		
Total cumulative		
dose		
Route	Nebulised	
Preparation	Using a 1:1,000 (1000 microgram/1 mL) ampoule	
	Draw up 0.5 mL/kg (0.5 mg/kg) adrenaline and add sodium chloride 0.9% to make a final	
	volume of 4 mL.	
Administration	Deliver final volume of 4 mL via nebuliser over 15 minutes.	
	Driving gas as prescribed by medical staff.	
	Set flow rate at 6 L/minute.	
	There will always be dead space that is not available for nebulisation - it is not possible to	
	nebulise to dryness.	
Monitoring	Administer under close supervision of medical staff.	
	Cardiorespiratory monitoring including respiratory rate, oxygen saturation, heart rate and	
	blood pressure.	
Contraindications		
Precautions	Infants with arrhythmias, hypertension or hyperthyroidism.	
	Infants with dilated or ischaemic cardiac disease (relative).	
	Crosscheck correct adrenaline strength ampoule used.	
	Do not use if the injection is pink or brown or contains a precipitate.	
Drug interactions	No information.	
Adverse reactions	Tachycardia and arrhythmia.	
	Systemic hypertension.	
Compatibility	Fluids: Sodium chloride 0.9%	
	Drugs: No information. Not to be mixed with other drugs in the same nebulisation chamber.	
Incompatibility	Fluids and drugs: No information.	
Stability	Discard remainder after use.	
Storage	Store below 25°C. Protect from light. Do not refrigerate or freeze.	
Excipients	Tartaric acid, sodium metabisulfite, sodium chloride and water for injections.	
Special comments		
Evidence	Efficacy:	
	Nebulised racemic adrenaline for extubation of newborn infants: There are no trials proving	
l		
	the efficacy of nebulised adrenaline compared to placebo or intravenous dexamethasone for	

ANMF consensus group JHCH_NICU_19.069

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	Treatment and prevention of bronchiolitis in newborns and infants: Nebulised adrenaline
	decreases hospitalisations in patients presenting to ER. There is no evidence to support the
	use of epinephrine for inpatients. [5, 8] (LOE I, GOR A)
	Treatment of children with croup: Nebulised epinephrine is associated with clinically and
	statistically significant transient reduction of symptoms of croup 30 minutes post-treatment.
	[6] (LOE I, GOR A) Evidence does not favour racemic epinephrine or L-epinephrine, or IPPB
	over simple nebulization. (LOE II, GOR B)
	Safety: Nebulised adrenaline is associated with increased heart rate and blood pressure. [2, 8]
	Pharmacokinetics: Not reported for nebuliser use in newborns or children. No difference in
	plasma adrenaline levels in asymptomatic children with history of anaphylaxis given
	adrenaline inhaler (10-20 activations) versus children given a placebo.[9]
Dunatica nainta	adrenaine initaler (10-20 activations) versus children given a piacebo.[9]
Practice points	
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