SUXAMETHONIUM CHLORIDE

Alert	Intubation, suction and ventilation equipment MUST be ready prior to administration of	
	suxamethonium. A medical officer/nurse practitioner (preferably two personnel) experienced in	
	advanced neonatal airway management techniques should be present when the medication is	
	being administered.	
	Risk of cardiac arrest from hyperkalemic rhabdomyolysis.	
	There are two preparations.	
	Chloride anhydrous salt (SAS product) equates to 110mg in 2 mL of suxamethonium chloride which	
	is 10% more suxamethonium than suxamethonium chloride dihydrate salt (Australian TGA	
1 1 1	registered product)	
Indication	Elective endotracheal intubation.	
Action	Short-acting, depolarising neuromuscular blocker. It acts as an acetylcholine antagonist at nicotinic	
	acetylcholine receptors at neuromuscular junctions, resulting in persistent depolarisation of the	
Davis Times	motor end plate.	
Drug Type	Neuromuscular blocking agent (depolarising)	
Trade Name	Suxamethonium Chloride (dihydrate) Injection BP, Succinolin Chloride (anhydrous) Injection,	
Duccontation	MercuryPharma Suxamethonium Chloride (dihydrate) Injection	
Presentation	100 mg/2 ml ampoule. *See "Alert" section above to account for brand difference.	
Dosage	IV (preferred): 2 mg/kg (up to 3 mg/kg)	
	IM (only if IV is not accessible): 3–4 mg/kg ⁹ (onset of action can be delayed up to 3 minutes and duration of action is up to 15 minutes)	
Doce adjustment	Therapeutic hypothermia: No information on the dose adjustment, but has been used.	
Dose adjustment	ECMO: Not applicable.	
	Renal impairment: use with caution as use associated with hyperkalaemia.	
	Hepatic impairment: may prolong duration of action. Avoid repeated doses.	
Route	IV, IM	
Maximum Dose	IV: 3 mg/kg/dose; IM: 4 mg/kg/dose	
Preparation	IV:*	
Preparation	Draw up 2 mL (100 mg of suxamethonium) and add 8 mL sodium chloride 0.9% to make final	
	volume 10 mL with a concentration of 10 mg/mL.	
	Volume 10 m2 with a concentration of 10 mg/ m2.	
	*Dilution for both dihydrate and anhydrous salts is kept the same as the difference is insignificant.	
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	IM: Administer undiluted.	
Administration	IV: Rapid injection at proximal cannula site.	
	IM: Administer in anterior thigh muscle.	
Monitoring	Continuous cardiorespiratory monitoring. Monitor temperature, blood pressure, oxygenation and	
	assisted ventilator status.	
Contraindications	Hyperkalaemia	
	Family history of malignant hyperthermia	
	Skeletal muscle myopathy	
	Hypersensitivity to suxamethonium	
Precautions	Anaphylaxis: Severe anaphylactic reactions (some life-threatening and fatal) have been reported.	
	Cross-sensitivity with other neuromuscular-blocking agents may occur; use extreme caution in	
	patients with previous anaphylactic reactions.	
	Bradycardia: May increase vagal tone. Risk of bradycardia may be increased with second dose and	
	may occur more often in children. Occurrence may be reduced by pre-treating with anticholinergic agents (e.g. atropine).	
	May Increase intraocular pressure.	
	May cause a transient increase in intracranial pressure.	
	May increase intragastric pressure, which could result in regurgitation and possible aspiration of	
	stomach contents.	
	Malignant hyperthermia: Use may be associated with acute onset of malignant hyperthermia; risk	
	may be increased with concomitant administration of volatile anaesthetics.	
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SUXAMETHONIUM CHLORIDE

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Drug Interactions	May enhance the effect of other agents with neuromuscular-blocking properties:
	acetylcholinesterase inhibitors; magnesium, quinidine, quinine, vancomycin, cyclophosphamide
	monohydrate, ciclosporin, esmolol, lincosamide, loop diuretics.
	Aminoglycosides: May enhance the respiratory depressant effect of aminoglycosides.
	Opioid analgesics: Suxamethonium may enhance the bradycardic effect of opioid analgesics.
A -l	Cardiac glycosides: May enhance the arrhythmogenic effect of cardiac glycosides
Adverse	Bradycardia is common in neonates and children, especially after a second dose of suxamethonium.
Reactions	May be prevented by administration of atropine prior to administration of suxamethonium.
	Hyperkalaemia
	Prolonged paralysis in infants with deficiency of pseudocholinesterase.
	Hypersensitivity reactions Malignant hyperthermia
	Management of suxamethonium overdose and/or toxicity is supportive.
	Wanagement of suxamethornum overdose and/or toxicity is supportive.
Compatibility	Dextrose 5%, dextrose 10%, sodium chloride 0.9%, dextrose 5% in sodium chloride 0.9%, dextrose 5%
Companionity	in sodium chloride 0.45%, sodium chloride 0.45%.
	Y-site administration: potassium chloride, propofol, vitamin B complex with C.
Incompatibility	Y site administration: Amino acid solution, lipid emulsion, heparin, alkaline solutions with pH > 8.5.
Stability	Suxamethonium Chloride (dihydrate) Injection BP brand: once removed from fridge, is stable below
	25 °C for 1 month only. Discard any unused product after that time, do not return to the fridge.
	Infusion solution: use within 24 hours
Storage	Refrigeration at 2°C to 8°C. DO NOT FREEZE.
Storage	For Succinolin and MercuryPharma brands: protect from light.
Special	Poorly absorbed from gastrointestinal tract – must be given IM or IV.
Comments	Rapidly and completely hydrolysed by hepatic and plasma pseudocholinesterase.
Comments	Very rapid onset (30–60 seconds) and short duration of action (3–5 minutes) with IV administration.
	Continuous administration over a prolonged period of time may result in irreversible blockade
	(phase II block).
	Should not be used without additional sedation.
Evidence	Efficacy
	Suxamethonium in combination with other drugs (analgesics and vagolytic agents) resulted in
	superior intubation conditions and a shorter procedure duration ¹⁻⁶ . (Level II, Grade A)
	For laparoscopic pyloromyotomy in term infants using propofol, sevoflurane and no intraoperative
	opioid, succinylcholine may be the neuromuscular blocking drug of choice, provided no
	contraindication is present ⁴ . (Level III-3, Grade B)
	Safety
	Suxamethonium has been very widely used, but has several rare side effects and causes an increase
	in blood pressure, simultaneously with depolarisation. ^{1,2} (Level II Grade B)
	Hyperkalaemia may occur, but major elevations are uncommon. It may trigger malignant
	hyperkalaemia, a rare autosomal dominant disorder of skeletal muscles that remain asymptomatic
	unless triggering substances are given. It should not be used in infants with hyperkalaemia or family
	history of malignant hyperthermia. (Level IV Grade D)
	It can cause prolonged neuromuscular blockade requiring ventilation until spontaneous resolution
	occurs in infants with pseudocholinesterase deficiency. ⁷ (Level IV Grade D)
	Pharmacokinetics
	Suxamethonium has a rapid onset of action (30 seconds) and a short duration of action (3 to 6
	minutes) with IV administration. The increased dose (2–3 mg/kg vs. 1 mg/kg in adults) requirement
	of succinylcholine in younger patients is thought to be due to its rapid distribution into an enlarged
	volume of extracellular fluid rather than an altered response to the action of the drug at
Dunation mainte	neuromuscular junction nicotinic acetylcholine receptors.8 (Level III Grade C)
Practice points	Suxamethonium in combination with other drugs (analgesics and vagolytic agents) resulted in
	superior intubation conditions and a shorter procedure duration. 1-6(Level II, Grade A)
	Chloride anhydrous salt equates to 110mg in 2 mL of suxamethonium chloride which is 10% more
	suxamethonium than suxamethonium chloride dihydrate salt.

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Authors Contribution

Original author/s	Dr Himanshu Popat, Dr Srinivas Bolisetty
Evidence Review	Assoc Prof David Osborn
Expert review	
Nursing Review	Ms Eszter Jozsa
Pharmacy Review	Ms Carmen Burman, Ms Thao Tran
ANMF Group contributors	Dr Himanshu Popat, Ms Carmen Burman, Ms Thao Tran, Dr John Sinn, Ms Wendy Huynh, Mr Jing Xiao
Final editing and review of the original	David Osborn, Carmen Burman, Thao Tran, Srinivas Bolisetty
Electronic version	Dr Ian Callander, Ms Cindy Chen
Facilitator	Dr Srinivas Bolisetty

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