SUXAMETHONIUM (A) (Revised: January 2014)



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TVDE	D	La visita a nace alla malaccant [CA]		
TYPE:	Depolarising muscle relaxant [S4]			
PRESENTATION:	100mg in 2ml – plastic ampoule			
	Tooms in 2111 plastic ampound			
ACTION:	Acts like the neurotransmitter acetylcholine at the neuromuscular junction. Persists for a period long enough to exhaust the motor endplate by prolonged depolarisation.			
	Onset IV: approx 45 seconds. Duration IV: 5 – 7 minutes.			
USE:	ICP	To facilitate airway management in selected patients		
ADVERSE EFFECTS:	1. Bradycardia			
	2. Potassium release			
	3. Increased intraocular and intragastric pressure			
	4. Occasionally, prolonged paralysis			
	5. Has been associated with malignant hyperthermia			
CONTRA-	1. Previous reaction to suxamethonium			
INDICATIONS:	2. Su	spected hyperkalaemia		
	3. Use in children			
PRECAUTIONS:	1. Elderly patient			
	2. Neuromuscular disease			
	3. Hypothermic patient			
	4. Fitting patient			
	5. Pa	tient with reversible pathology		
	Selec	t patients carefully – always have a fallback position!		
DOSES:				
ADULT:				
ICP 1.5mg/kg IV – over 30 – 60 seconds (to a maximum of 150mg)				
PAEDIATRIC:				
Not used				

continues over

SUXAMETHONIUM (A) cont.

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Suxamethonium Dose Table				
This dose table applies to Suxamethonium 100mg in 2mls				
Weight (kg)	Dose (mg)	Volume (mls)		
40	60	1.2		
50	75	1.5		
60	90	1.8		
70	105	2.1		
80	120	2.4		
90	135	2.7		
100	150	3		
Maximum dose 150mg				

SPECIAL NOTES:

- To be used *only* following completion of the ACTAS designated training programme.
- If heart rate less than 50/minute, consider atropine prior to suxamethonium.
- Suxamethonium may cause bradycardia. If patient is bradycardic once ETT is tied in, consider a dose of atropine.
- (NOTE: bradycardia may be a result of a head injury and raised ICP thus, BP will be elevated. In this case there is no requirement for atropine regardless of the degree of bradycardia).