

Title	Adenosine Protocol – Antiarrhythmic		
Areas where applicable	Cardiac, Critical Care, Emergency Medicine Services and Clinical Emergency Response Systems teams as therapeutic treatment or diagnostic aid.		
Areas where <u>not</u> applicable	NOT for use with radionuclide myocardial perfusion imaging or for non- antiarrhythmic use in Cardiac Catheter Laboratory		
Authorised Prescribers	Medical officers familiar with the product.		
Indications for use	Therapeutic: Rapid conversion to a normal sinus rhythm of paroxysmal supraventricular tachycardia (SVT), including those associated with accessory bypass tracts (Wolff-Parkinson-White syndrome). Diagnostic: As an aid to differential diagnosis of narrow or broad complex tachycardia due to the slowing of AV conduction which makes atrial activity more visible on		
Drug Action	ECG.		
Drug Action	 Antiarrhythmic actions: Slows impulse formation of the sino-atrial node Slows conduction time through the atrio-ventricular node Can interrupt re-entry pathways through the atrioventricular node Coronary vasodilator 		
Pharmacokinetics	Onset: Immediate Peak: 10 seconds Duration: 10 – 30 seconds		
Contraindications	 Hypersensitivity to adenosine Second or third degree heart block (unless a functioning artificial pacemaker present) Sinus node dysfunction, such as sick sinus syndrome or symptomatic bradycardia (unless a functioning artificial pacemaker present) Bronchoconstriction or bronchospastic lung disease (e.g. asthma) either known or suspected 		
	Severe hypotension		
Precautions	 Convulsion /seizure history Recent myocardial infarction Recent heart transplant (less than 1 year) First degree AV or bundle branch block Atrial fibrillation, flutter, especially with accessory pathway Heart failure Hypotension, hypertension Bronchoconstriction in patients with asthma Obstructive lung disease not associated with bronchoconstriction e.g. COPD, bronchitis Bradycardia Prolonged QT interval 		
	 Pregnancy and/or breastfeeding. Limited data available about use in first trimester. Use during second and third trimester is considered safe and effective. Higher doses may be required. Monitor foetal heart rate during administration. 		

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Proposed Place in Therapy	Adenosine is first line drug therapy choice (after physical manoeuvres that enhance vagal tone)				
Dosage	Therapeutic: To be administered by rapid bolus (2 seconds), followed by a rapid 20mL sodium chloride 0.9% flush.				
	Dose 1 Adenosine 6 mg rapid peripheral IV bolus OR 3 mg if administered by central venous access				
	Dose 2 If the first dose is ineffective but well tolerated, after 2 minutes give Adenosine 12mg rapid peripheral IV bolus OR				
	6 mg if administered by central venous access				
	Dose 3 If second dose is ineffective but well tolerated after a further 2 minutes, give a further dose of Adenosine 18 mg rapid peripheral IV bolus OR				
	12 mg if administered by central venous access				
	Diagnostic:				
	 The above ascending dosage schedule should be employed until sufficient diagnostic information has been obtained. Patients who develop high level AV block at a particular dose should not be given further dosage increments. 				
	General information:				
	 The initial adenosine dose should be reduced to 3 mg in patients taking dipyridamole or carbamazepine, those with a transplanted heart or if given by central venous access. Dose adjustment is not required for hepatic or renal impairment IV infusion is ineffective in treating supraventricular tachycardia 				
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Administration instructions	 Universal precautions Administer adenosine undiluted by rapid IV bolus (over 2 seconds) followed by a rapid 20 mL sodium chloride 0.9% flush. Adenosine has a very short duration of effect making it necessary to give as a rapid bolus 				
	 Warn patient they may experience anxiety or a feeling of "impending doom" - this will pass quickly. Administer either directly into a large peripheral vein or into an IV line (injected as proximally as possible). 				
	Patients who develop high level AV block at a particular dose should not be given further dosage increments.				

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Important Drug Interactions	Caffeine and theophyllines antagonise the effects of adenosine; a higher dose of adenosine may be required. Dipyridamole inhibits cellular uptake of adenosine, increasing the risk of bradycardia, so that the dose for stopping a tachycardia may be much less than usual. Stop dipyridamole 24 hours before planned use of adenosine or use lower initial dose of adenosine (a quarter to a half). Carbamazepine has been reported to increase the degree of heart block produced, so lower the initial dose of adenosine.		
	The effect of adenosine is not blocked by atropine.		
Presentation	6 mg in 2 mL vial		
Monitoring requirements	The patient should have cardiac monitoring throughout the procedure. A defibrillator and emergency resuscitation equipment must be available for immediate use.		
	Ensure that the monitor printer or 12 lead ECG is set to record as soon as adenosine is injected. Continue to record until rhythm returns to normal. Heart blocks and asystole may occur. These are generally transient due to the short half-life.		
	Monitor vital signs observations pre and post administration and with change of rhythm		
	Blood pressure should be measured in the arm opposite to the adenosine infusion		
Adverse effects and	Adverse effects resolve rapidly on stopping treatment due to the drugs short duration of action.		
Management of complications	Explain possible adverse effects to patient before administration. Ensure patient understands that these effects will be short-lived.		
	Common: flushing, dyspnoea, chest pain/pressure, nausea or abdominal discomfort, headache, dizziness, apprehension, burning sensation, bradycardia, asystole, sinus pause and A-V block		
	Infrequent: transient arrhythmias, recurrence of SVT, hypotension, tingling in arms or legs, metallic taste		
	Rare: bronchospasm, injection site reaction		

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Basis of	1.	eTG complete November 2012 accessed 8/3/2013			
Protocol/Guideline:		Adenocor® TGA approved Product Information 18/2/2008 accessed via			
		MIMS Online 7/3/2013			
	3.	Australian Medicines Handbook January 2013 accessed 7/3/2013			
	4.	Australian Injectable Drugs Handbook, Fifth Edition accessed 7/3/2013			
		Weismuller P, Kattenbeck K, Heinroth KM, et al. [Terminating			
		supraventricular tachycardia with adenosinecomparing the effectiveness			
		of 12 mg and 18 mg]. Dtsch Med Wochenschr. 2000;125(33):961–969.			
	6.	Curtis A, Belardinelli L, Woodard D, Brown C, Conti J. Induction of			
		Atrioventricular Node Reentrant Tachycardia With Adenosine: Differential			
		Effect of Adenosine on Fast and Slow Atrioventricular Node Pathways. J			
		Am Coll Cardiol. 1997;30(7):1778-1784.			
		Domanovits, H., et al., Adenosine for the management of patients with			
		tachycardias:a new protocol. <i>European Heart Journal</i> , 1994. 15 (5): p. 589-			
		593.			
	8.	Elkayam U., Goodwin TM. Adenosine therapy for supraventricular			
		tachycardia during pregnancy. American Journal of Cardiology. 75 (7):521.			
	•	District Clinical Emergency Response System Committee			
Groups consulted in	•	Cardiac and Respiratory Clinical Stream			
development of this	•	Critical Care and Emergency Medicine Clinical Stream			
guideline	Drug and Quality Use of Medicine Committee				
		Pharmacy Departments			
	•	Royal Hospital for Women			

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GOVERNANCE					
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