

Title	Subcutaneous lidocaine (lignocaine) for refractory neuropathic pain in the palliative care setting		
Area where Protocol/Guideline applicable	SESLHD Inpatient settings (including Calvary hospital)		
Authorised Prescribers	Specialist Palliative Care Services		
Indications for use	Must be used under the supervision of a Palliative Care Specialist.  1. Refractory neuropathic pain not responding to standard analgesic drugs, including optimal use of opioids and adjuvant therapies.  2. Refractory pruritis when the oral route is no longer available		
Place in Therapy	Lidocaine is a systemic local anaesthetic agent and known membrane stabiliser. It is used in the palliative care setting as a third or fourth line drug in the treatment of complex & refractory neuropathic pain.		
Contraindications	<ul> <li>Adams-Stokes syndrome, Wolff-Parkinson-White syndrome</li> <li>Severe atrioventricular, sino-atrial or intraventricular heart block not managed with a pacemaker</li> <li>Sensitivity to amide-type local anaesthetics</li> <li>Patients on flecainide</li> </ul>		
Precautions & Relative Contraindications	Cardiac monitoring in the palliative care setting is not indicated due to doses not exceeding the threshold of 2 g over 24 hours via CSCI.  Caution in patients with known cardiac disease, cerebral palsy, electrolyte imbalance (correct before starting treatment) or a history of arrhythmia		

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Important Drug Interactions	Avoid in patients taking flecainide
Known Adverse Effects	Monitor closely for the following initial signs of systemic toxicity:  Light-headedness, Dizziness Perioral numbness or tingling (around lips) Tinnitis Metallic taste Drowsiness and dysarthria If any of the above are observed, cease infusion immediately and inform Palliative Care Medical Officer. Lidocaine infusion may be restarted at a lower dose.  Worsening toxicity is indicated by the progressive appearance of: Visual changes Muscle spasm Seizures Coma Cardiorespiratory depression and arrest
Preparations	Lidocaine (lignocaine) 2% 100 mg/5 mL ampoules Lidocaine (lignocaine) 10% 500 mg/5mL ampoules
Dosing	Lidocaine has a narrow therapeutic index and dose is determined by consultation with Palliative Care Specialist.  Starting dose: Lidocaine 0.5 mg/kg/hr (i.e. 200 - 800 mg over 24 hours) via CSCI <sup>6</sup> Titration: Increase by 200- 800 mg every 24 hours as required; titrate to effect.  Maximum dose 2,800 mg per 24 hours (~120 mg/hr)

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	DOSE of lidocaine	VOLUME & recommended FORMULATION		Approx. Volume of WFI to make total volume	
		of lidocaine			
		Lidocaine 2%	Lidocaine 10%	WFI	
	200 mg	10 mL	-	7 mL	
	400 mg	-	4 mL	13 mL	
	500 mg	-	5 mL	12 mL	
Administration	600 mg	-	6 mL	11 mL	
Administration	700 mg	-	7 mL	10 mL	
	800 mg	-	8 mL	9 mL	
	900 mg	-	9 mL	8 mL	
	1000 mg	-	10 mL	7 mL	
	1100 mg	-	11 mL	6 mL	
	1200 mg	-	12 mL	5 mL	
	1300 mg	-	13 mL	4 mL	
	1400 mg	-	14 mL	3 mL	
	1500 mg	-	15 mL	2 mL	
	1600 mg	-	16 mL	1 mL	
	For doses < 1600 mg; use a 20 mL syringe and make the volume up to 17 mL.  Doses > 1600 mg will require a 30 mL syringe.				
Diluents	Water for Inject	tion (WFI)			
Drug Compatibility	Lidocaine should not be mixed in a syringe with any other medication due to lack of robust compatibility data. Lignocaine may be given in conjunction with ketamine but NOT in same syringe driver.				
Monitoring requirements	Monitor for signs of adverse effects (as above) and if any of the initial signs of toxicity occur cease the infusion and report to the Palliative Care consultant immediately.				
	Perform 4-hourly subcut infusion site checks as per Subcutaneous Syringe Driver inpatient management form SES130.021				
Practice Points	Lidocaine is only given by continuous subcutaneous infusion via syringe driver. It is NOT to be given by intermittent bolus subcutaneous injections.				

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	St George Palliative Care Team		
Consultation	SESLHD Palliative Care working party Dr Caitlin Sheehan, Staff Specialist St George & Calvary Hospital		

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