Medicine Guideline



Areas where Protocol/Guideline applicable	SESLHD Inpatient settings (including Calvary hospital)
Authorised Prescribers:	Specialist Palliative Care Services
Indication for use	Must be used under the supervision of a Palliative Care Specialist: -
	 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.
Clinical condition	All conditions causing refractory neuropathic pain or pruritus as per the indications for use.
Proposed 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.
Adjunctive Therapy	Can be used alone or in conjunction with other neuropathic pain agents depending on the clinical circumstances.
Contraindications	 Adams-Stokes syndrome, Wolff-Parkinson-White syndrome Severe atrioventricular, sino-atrial or intraventricular heart block not managed with a pacemaker Sensitivity to amide-type local anaesthetics Patients on flecainide
Precautions and Relative Contra- indications	Cardiac monitoring in the palliative care setting is not indicated due to doses not exceeding the threshold of 2,800mg (2.8g) over 24 hours via CSCI.
	Use with caution in patients with known cardiac disease, cerebral palsy, and history of arrhythmia. Where possible, aim to correct electrolyte balances prior to commencement.
	Consider dose modification (up to 50%) in renal or hepatic impairment, and in the context of frailty.
Important Drug	Avoid in patients taking flecainide

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Interactions				
Dosage		as a narrow therap with Palliative Ca		ose is determined by
			00 to 800 mg over	24 hours) via CSCl ⁶ .
	Titration: Increase by effect.	200 to 800 mg ev	ery 24 hours as re	quired; titrate to
	<mark>Maximum d</mark> mg/hr)	ose 2,800 mg (2.	8g) per 24 hours (a	approximately 120
		se modification (u and in the contex	p to 50%) in renal t of frailty.	or hepatic
Prescribing Instructions		Lidocaine must be prescribed on the eMM system. In the absence of eMM systems, the appropriate paper medication chart may be used.		
Administration Instructions	DOSE of lidocaine	VOLUME & recommended FORMULATION of lidocaine		Approx. Volume of Water for Injection (WFI) to make total volume
		Lidocaine 2%	Lidocaine 10%	WFI
	200mg	10mL	-	7 mL
	400mg	-	4 mL	13 mL
	500mg	-	5 mL	12 mL
	600mg	-	6 mL	11 mL
	700mg	-	7 mL	10 mL
	800mg	-	8 mL	9 mL
	900mg	-	9 mL	8 mL
	1000mg	-	10 mL	7 mL
	1100mg	-	11 mL	6 mL
	1200mg	-	12 mL	5 mL
	1300mg	-	13 mL	4 mL
	1400mg	-	14 mL	3 mL
	1500mg	-	15 mL	2 mL
	1600mg	-	16 mL	1 mL

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	to 17 mL.
	Doses > 1600 mg will require a 30 mL syringe
Preparations	Lidocaine (lignocaine) 2% 100 mg/5 mL ampoules
	Lidocaine (lignocaine) 10% 500 mg/5mL ampoules
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.
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
Monitoring requirements Safety	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.
Effectiveness (state objective criteria)	Perform 4-hourly subcutaneous infusion site checks as per Subcutaneous Syringe Driver inpatient management form SES130.021. These paper forms need to be purchased as per local processes.
Practice Points	Lidocaine is only given by continuous subcutaneous infusion via syringe driver. It is NOT to be given by intermittent bolus subcutaneous injections.
Management of Complications	If patient reports side effects, the infusion should be ceased and management discussed with palliative care team. Where appropriate, consider an ECG after discussion with the team.

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Basis of Protocol/Guideline:	 Benowitz NL, Meister W. Clinical pharmacokinetics of lignocaine. Clinical pharmacokinetics. 1978;3(3):177- 201 Pasero CM, M. Pain Assessment and Pharmacologic Management. Missouri: Mosby Elselvier; 2010 Schwartzman RJ, Patel M, Grothusen JR, Alexander GM. Efficacy of 5-day continuous lidocaine infusion for the treatment of refractory complex regional pain syndrome. Pain medicine (Malden, Mass). 2009;10(2):401-12 Swenson BR, Gottschalk A, Wells LT, Rowlingson JC, Thompson PW, Barclay M, et al. Intravenous lidocaine is as effective as epidural bupivacaine in reducing ileus duration, hospital stay, and pain after open colon resection: a randomized clinical trial. Regional anesthesia and pain medicine. 2010;35(4):370-6. Hsu Y-W, Somma J, Newman M, Mathew JP. Population Pharmacokinetics of Lidocaine Administered During and After Cardiac Surgery. Journal of cardiothoracic and vascular anesthesia. 2011;25(6):931-6. Palliative Care Formulary online. In Medicines Complete. Pharmaceutical Press. Available via CIAP accessed April 2025. Palliative Care [December 2024]. In Therapeutic Guidelines Ltd. Available via CIAP, accessed April 2025. CHCK Policy 'Pain Management (Neuropathic – Lignocaine & Ketamine). September 2018 Dickman A, Schneider J. 2016. The Syringe Driver: Continuous subcutaneous infusions in palliative care. Oxford University Press; 2016 Macleod, R Macfarlane, S. 2018 The Palliative Care Handbook. 9th Ed. Hammondcare Media.
Groups consulted in development of this guideline	St George Palliative Care Team SESLHD Palliative Care working party Dr Caitlin Sheehan, Staff Specialist St George & Calvary Hospital

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GOVERNANCE		
Enactment date	November 2020	
Reviewed (Version 2)	November 2022	
Reviewed (Version 3)	December 2023	
Reviewed (Version 3.1)	April 2025	
Expiry date:	April 2027	
Ratification date by SESLHD DTC	3 April 2025	
Chairperson, DTC	Stuart Binns, Acting Chair, SESLHD DTC	
Committee		
Version Number	4.0	