

| Title | Subcutaneous furosemide (frusemide) for end stage heart failure with fluid overload in the dying patient |
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| Area where Protocol/Guideline applicable | SESLHD Inpatient settings (including Calvary hospital) |
| Indications for use | For management of refractory congestive cardiac failure and/or pulmonary oedema in end stage cardiac failure, when the oral route is no longer possible and the IV route is not appropriate or desirable. |
| Place in Therapy | When oral route for furosemide therapy no longer available and patient in terminal phase of illness. |
| Clinical condition | Continuing diuretic therapy is crucial to decrease the pulmonary fluid overload in a patient with end-stage heart failure on regular furosemide to minimise exacerbation of breathlessness and other symptoms associated with end stage heart failure. |
| If part of combination therapy, list other drugs | Frequently used in conjunction with opioids for dyspnoea management in end stage heart failure |
| Contraindications | Allergy to furosemide or sulfonamides Fluid depletion, anuria |
| Known adverse effects | Electrolyte imbalance, dehydration, metabolic alkalosis Orthostatic hypotension, dizziness Rash Tinnitus |
| Preparations | Furosemide 20mg/2mL ampoules Furosemide 250mg/25mL ampoules |
| Dose conversion for oral to subcutaneous route | A ratio of 1:1 between oral and subcutaneous routes should be used |
| Dosage | Intermittent dosing: initial dose 20mg once or twice daily subcutaneously Note doses greater than 20mg are unsuitable for intermittent dosing due to volume. Continous Subcutaneous Infusion (CSCI): Adjust dose according to clinical response to a maintenance dose of 40-250mg daily (maximum 1g daily) |

| Administration | Dilute to a maximum concentration of 10mg/mL: Maximum 200mg in 20mL syringe Maximum 300mg in 30mL syringe (30mL syringes available from Palliative Care team) If higher doses required two 12 hour syringe drivers may be necessary |
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| Diluents | Sodium chloride 0.9% |
| Drug Compatibility | Furosemide injection is alkaline and there is a high risk of incompatibility when mixed with acidic drugs. Because of this and the lack of compatibility data, furosemide should not be mixed in the same syringe with any other drugs |
| Monitoring requirements | Monitor level of fluid overload and titrate dose accordingly. Monitor for injection site reactions. If administered via CSCI perform 4 hourly infusion site checks as per Subcutaneous Syringe Driver inpatient management form SES130.021 |
| Practice Points | Administer alone in a separate syringe driver |
| Basis of Protocol/Guideline (including sources of evidence, references) | Palliative Care Formulary 7 th Ed, 2020 p 67-72 Therapeutic Guidelines – Palliative Care Version 4, 2016, Dickman A, Schneider J. The syringe driver: continuous subcutaneous in palliative care. Oxford University Press; 2016 Domenic, A.S et al, <i>Subcutaneous Furosemide in</i> <i>Heart Failure,</i> JACC: Basic to Translational Science VOL, 3 NO 1, Feb 2018 pp 25 -34 |
| Consultation | St George Palliative Care Team SESLHD Palliative Care working party: Dr Jan Maree Davis, Medical Director, Palliative Care. |

| AUTHORISATION | | |
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| Author (Name) | Dr Jan Maree Davis | |
| Position | Medical Director, Palliative Care | |
| Department | SESLHD Southern Sector | |
| Department Contact (for ongoing maintenance of Protocol/Guideline) | JanMaree.davis@health.nsw.gov.au | |

| GOVERNANCE | | |
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| Chairperson, QUM Committee | Dr John Shephard | |
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