

Prescribing Protocol - Fentanyl 2microg/mL with Adrenaline (epinephrine) 2 microg/mL and Bupivacaine 0.1% epidural infusion		
Areas where Protocol/Guideline applicable e.g. District, Hospital, ITU, Ward	District-wide	
Areas where Protocol/Guideline not applicable	N/A	
Authorised Prescribers	Anaesthetists, Anaesthetic Registrars, Pain Specialists	
Indication for use	Pain Management	
<b>Clinical condition</b> Patient selection: Inclusion criteria (list investigations necessary and relevant results)	Peri-operative pain management Rib fracture analgesia Each individual patient's risk / benefit assessment will be considered by the prescriber	
Contra-indications	Severe coagulation disturbances Patient refusal Allergy to local anaesthetics Localised infection at insertion site	
Precautions	Hypotensive patients Patients with pre-existing neurological disease Anticoagulant and antiplatelet drugs	
<b>Place in Therapy</b> State whether drug to be used as first, second or third line. When not first line, describe therapies to be used first. (Consider using algorithm)	First-line	

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<b>Dosage</b> (Include dosage adjustment for specific patient groups)	Suggested starting dose 8-10mL/hr (range 6-12 mL/hr) Delivered via Continuous Infusion (CI) or Programmed Intermittent Epidural Bolus (PIEB) and/or Patient Controlled Epidural Bolus (PCEA). Refer to <u>Pain</u> <u>Management - Epidural Analgesia (Adult) non-obstetric</u> <u>procedure SESLHD/324.</u> Example prescription (to be prescribed on <u>NSW Health</u> <u>Epidural Analgesia Adult Form SMR130.022</u> ): PIEB 4mL every 30mins + PCEA 3mL every 15mins PRN. Hourly limit 24mL
Duration of therapy	2 to 5 days
Administration instructions	Premixed infusion bags <b>must</b> be used. Administer using a sterile single-use administration set via programmable pump
Availability and supply	<ul> <li>Premixed bupivacaine with fentanyl and adrenaline (epinephrine) bags are a compounded Schedule 8 product prepared by Baxter Healthcare with an expiry date of 90 days.</li> <li>A form C must be completed by the Staff Specialist prior to first order.</li> <li>The supply of bags is maintained and stored in Pharmacy.</li> <li>Nursing staff must monitor expiry dates of the epidural pre- mixed bags on the ward and order from Pharmacy in a timely manner. The turnaround time to order from Baxter via Pharmacy is two days.</li> <li>There is 1 bag available in the Post Acute Care Unit (POWH only).</li> </ul>
Monitoring requirements	Observations and monitoring as per <u>NSW Health</u> <u>Epidural Analgesia Adult Form SMR130.022</u> and <u>Pain</u> <u>Management - Epidural Analgesia (Adult) non-obstetric</u> <u>procedure SESLHD/324.</u>
Management of complications	Intravenous fluid bolus Pharmacological management Contact Acute Pain Service during business hours, and the anaesthetist on call after hours for advice.

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Basis of Protocol/Guideline (including sources of evidence, references)	ANZCA Acute Pain Management: Scientific Evidence Fifth Edition 2020:
	'In postoperative thoracic epidural infusion, the addition of adrenaline to fentanyl and ropivacaine or bupivacaine improved analgesia.' Level II evidence (Evidence obtained from at least one properly designed randomised-controlled trial).
	Sakagutchi et al. Does Adrenaline Improve Epidural Bupivacaine and Fentanyl Analgesia After Abdominal Surgery? Anaesth Intensive Care 2000; 28: 522-526.
	Niemi G, Breivik H. The minimally effective concentration of adrenaline in a low-concentration thoracic epidural analgesic infusion of bupivacaine, fentanyl and adrenaline after major surgery. Acta Anaesthesiol Scand 2003; 47: 439-450.
	Kjonikksen J et al. Stability of an epidural analgesic solution containing adrenaline, bupivacaine and fentanyl. Acta Anaesthesiol Scand 2000; 44: 864–867.
	Brustugun J et al. The stability of a sulphite-free epidural analgesic solution containing fentanyl, bupivacaine, and adrenaline. Acta Anaesthesiol Scand 2013; 57: 1321–1327.
	Priston MJ et al. Stability of an epidural analgesic admixture containing epinephrine, fentanyl and bupivacaine. Anaesthesia 2004; 59: 979-983.
Groups consulted in development of this protocol	Department of Anaesthesia and Pain Management, Prince of Wales Hospital
	Pharmacy Department, Prince of Wales Hospital

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GOVERNANCE		
Renewal date	February 2021	
Expiry date: (maximum 36 months from date of original approval)	29 February 2024	
Ratification date by SESLHD QUM Committee	4 February 2021	
Chairperson, QUM Committee	Dr John Shephard	
Version Number	2.0	