

**NURSING CARE PLAN No 23**  
**EPIDURAL INFUSION**

Full Name: .....

Address: **Addressograph** .....

HCR:.....

(All care plans must be used in conjunction with care plan 1)

<b>Care Plan No 23</b>	<b>EPIDURAL INFUSION</b>	<b>Issue Date:</b> April 2020	
<b>Problem</b>		<b>Review Date:</b> April 2023	
<p>..... has an epidural infusion of ..... in situ and so is potentially at risk from: a) inadequate pain management; b) dense motor block c) excessive sedation d) respiratory depression; e) urinary retention; f) pruritus (itching); g) nausea and vomiting ; h) impaired mobility ; i) neurological damage (j) technical complications (k ) Infection</p>			
<b>GOALS</b>			
<p>a) .....’s pain will be managed individually so that the level of discomfort experienced by him/her is judged to be acceptable by ..... and / or his/her family.</p> <p>b) Any adverse side effects from the epidural will be identified promptly and appropriate action taken.</p> <p>c) .....’s safety will be maintained at all times, problems identified and appropriate action taken.</p>			
	<b>NURSING INTERVENTION</b> (Refer to hospital guidelines for epidural infusions, regional anaesthetic observation chart and regional analgesia prescription sheet)	<b>Commencement, Date, Signature, Time, Grade</b>	<b>Discontinued, Date, time, Signature, Grade</b>
<b>1</b>	<b>INFUSION</b>		
	<ul style="list-style-type: none"> <li>• Epidural Infusions are administered via dedicated infusion pump (CADD Solis©).</li> <li>• Nursing staff will answer any questions that.....and his/her family ask and will provide the epidural parent information sheet to the child/family.</li> <li>• Two staff nurses check infusion pump programme settings, against the prescription.</li> <li>• Ensure.....has IV access while the epidural is in progress &amp; for 6 hours afterwards.</li> <li>• The hourly volume infused, will be recorded on the regional analgesia observation chart.</li> <li>• Nursing staff will independently double check, sign &amp; document: any changes to the infusion: rate, change of bag or giving set on the prescription chart.</li> <li>• The infusion will be changed <b>every 24 hours</b> if it <b>includes an additive</b> e.g. clonidine or fentanyl. If there is <b>no additive</b> the infusion and set will be <b>changed every 48 hours</b>.</li> <li>• Any technical problems with the infusion pump will be reported to the biomedical engineering department / Pain Service and the pump will be removed from service.</li> <li>• The infusion, will be maintained for a <b>minimum of 48 hours</b> and maximum of 96 hours for non-tunnelled catheter.</li> </ul>		
<b>2</b>	<b>DETECTING AND MANAGING SIDE EFFECTS</b>		
	<ul style="list-style-type: none"> <li>• Monitor and observations as per regional analgesia observation sheet.</li> <li>• Ensure continuous pulse oximetry is in situ <b>at all times</b> until the infusion has been discontinued (plus Apnoea monitor &lt; 1yr).</li> <li>• Heart rate and blood pressure will be recorded <b>hourly</b> for <b>four hours</b> following an increase in infusion rate and <b>four hourly</b> until epidural is discontinued.</li> <li>• .....will be monitored whilst receiving an epidural Infusion for signs of local anaesthetic toxicity as per epidural guideline.</li> <li>• The epidural infusion will be stopped and the Pain Control Team/Anaesthesiologist on call will be contacted if .....is experiencing signs of epidural complications.</li> <li>• Nursing observations will be more frequent if any side effects are detected.</li> </ul>		
<b>2a</b>	<b>PAIN ASSESSMENT AS PER CARE PLAN 30</b>		
	<ul style="list-style-type: none"> <li>• Pain will be assessed and managed as per care plan 30.</li> </ul> <p>Additional analgesia will be administered if.....pain is not well controlled. Other measure include increasing the rate of infusion, repositioning, or requesting the pain service to review .....</p>		
<b>2b</b>	<b>MOTOR BLOCK</b>		
	<ul style="list-style-type: none"> <li>• The level of Motor block will be assessed using the modified Bromage score <b>4 hourly</b> as per regional observation sheet and guidelines.</li> <li>• Where the level of motor block is <math>\geq 2</math> the anaesthesiologist or pain service will be informed and the infusion may be decreased by 10%.</li> </ul>		

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<ul style="list-style-type: none"> <li>Where the motor block is <math>\geq 3</math> follow the algorithm for high (dense) motor block.</li> </ul>		
<b>2c</b>	<b>SENSORY BLOCK</b>	
<ul style="list-style-type: none"> <li>Sensory dermatome level will be assessed as per epidural guideline <b>4 hourly</b> or more often if .....is experiencing pain or the level of the block is too high or too low.</li> <li>A bolus of epidural and/or rate increase will be administered where the sensory block is too low as per epidural guideline.</li> </ul>		
<b>2d</b>	<b>HEADACHE</b>	
<ul style="list-style-type: none"> <li>.....will be monitored for signs of severe frontal headache that is worse when sat forward. This may be indicative of a leak of cerebro-spinal fluid.</li> <li>Follow epidural guideline if headache is present.</li> </ul>		
<b>2e</b>	<b>PRURITIS</b>	
<ul style="list-style-type: none"> <li>Assess for pruritis at least 4 hourly.</li> <li>Pruritus will be managed with an antihistamine or low dose Naloxone.</li> <li>Where pruritus persists, Fentanyl can be removed as per guideline.</li> </ul>		
<b>2f</b>	<b>INFECTION</b>	
<ul style="list-style-type: none"> <li>The epidural filter will remain in place at all times.</li> <li>If Temperature exceeds 38.5<sup>o</sup>, the Pain Service or anaesthetist on call will be informed.</li> <li>All changes to epidural infusion bag or giving set will be done using <b>ANTT</b></li> </ul>		
<b>2g</b>	<b>EPIDURAL SITE</b>	
<ul style="list-style-type: none"> <li>The bacterial filter and connection to the epidural catheter will be reinforced with a clear dressing (Tegaderm) to reduce the risk of accidental disconnection.</li> <li>The epidural site will be checked at least four hourly for redness, tenderness, leakage or problems with dressing and correct position.</li> <li>Where leaking is detected, the infusion rate will be reduced.</li> </ul>		
<b>3</b>	<b>MOBILISATION / PRESSURE AREAS</b>	
<ul style="list-style-type: none"> <li>A risk assessment re the need for a pressure-relieving device will be carried out.</li> <li>Pressure points will be assed for redness, .....s' position will be changed 2-3 hourly.</li> <li>.....will be encouraged to mobilise as his/her condition allows. Ensure he/she is accompanied at all times when mobilising.</li> </ul>		
<b>4</b>	<b>REMOVING EPIDURAL CATHETER</b>	
<ul style="list-style-type: none"> <li>.....is receiving anticoagulant therapy, the epidural catheter will be removed as per the epidural guideline <b>12 hours after</b> the last dose of LMWH and a minimum of <b>2 hours before</b> the next dose of anticoagulant is due to be given.</li> <li>Observations of pulse, respirations, pain score and Levobupivacaine toxicity must be continued for 4-6 hours after an epidural infusion has been discontinued.</li> <li>The epidural catheter will be removed using ANTT level three principles on advice from the Pain Service or anaesthetist ensuring supplemental analgesia has been administered.</li> <li>On removal of epidural, the catheter tip will be inspected for intactness and the removal will be documented.</li> </ul>		

Created by Nursing Department, Reviewed by V Kilcullen 2020

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