

# Naltrexone for treatment of long-acting opioid poisoning



<b>Areas where Protocol/Guideline applicable</b>	SESLHD Inpatients > 16 years
<b>Authorised Prescribers:</b>	SESLHD Toxicologists or on approval by Toxicologists
<b>Indication for use</b>	Off-label use as opiate antagonist for the treatment of long-acting opioid poisoning (e.g. methadone, buprenorphine, carfentanil) in opioid naïve patients
<b>Clinical condition</b>	<ul style="list-style-type: none"> <li>Opioid naïve patients who present to ED with a history of ingestion of a long-acting opioid (eg: methadone, buprenorphine, carfentanil) or symptoms of opioid toxicity</li> <li>Clinical response to naloxone, but require a naloxone infusion</li> </ul>
<b>Formulary Status</b>	Naltrexone is not listed on the SESLHD Medicines Formulary for this indication. Retrospective Individual Patient use (IPU) applications must be submitted on the IPU platform.
<b>Proposed Place in Therapy</b>	To be used in patients who are known to be opioid naïve, have taken a long-acting opioid and developed respiratory depression, predicted to need to have a prolonged naloxone infusion and have responded clinically to an initial naloxone injection. Naltrexone will be used as second-line therapy in these patients following clinical response to initial naloxone injection.
<b>Adjunctive Therapy</b>	Naloxone
<b>Contra-indications</b>	<ul style="list-style-type: none"> <li>Patients who are opioid dependent</li> <li>Patients being treated with an opioid analgesic for a painful condition or if a painful procedure is planned (since naltrexone will reduce analgesia)</li> <li>Contraindicated in acute hepatitis, liver failure or when liver enzymes &gt;3 times ULN</li> <li>Known allergy to naltrexone</li> </ul>
<b>Precautions</b>	Chronic hepatitis B and/or C or raised baseline liver enzymes – monitor liver function (especially total bilirubin) regularly to ensure naltrexone does not exacerbate condition. Avoid use in pregnancy if possible, contact Mothersafe if required (Ph: 93826539)
<b>Important Drug Interactions</b>	<u>Naltrexone + Opioids:</u> Naltrexone reversibly blocks opioid receptors and reduces effects of opioids; in opioid dependence may precipitate withdrawal symptoms at start of naltrexone treatment
<b>Dosage</b>	Adult and Children > 16 years: <b>50 mg</b> orally once only
<b>Duration of therapy</b>	Single dose. An additional dose may be required after 24 hours to treat respiratory depression, but only on the recommendation of a toxicologist.
<b>Prescribing Instructions</b>	Prescribed on the eMR. In the absence of eMM systems, the appropriate paper medication chart may be used

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<b>Administration Instructions</b>	<p>Initial treatment with naloxone if required for immediate treatment as naltrexone may take up to an hour to exert its peak effect.</p> <p>The patient will be given one 50mg dose of oral naltrexone, and then monitored for 4 hours in the ED. Patients will then be stepped down to a normal ward bed for 12 hours observations</p> <p>Patient to be discharged 12 hours post last naltrexone dose</p> <p>Patients should be discharged home during daylight hours with a carer. Instructions should be provided to return to the ED should the patient develop symptoms of opioid toxicity (cyanosis, reduced level of consciousness, bradypnoea or apnoea).</p>
<b>Adverse effects</b>	Possible GI upset but rare
<b>Monitoring requirements</b> Safety Effectiveness (state objective criteria)	<p>Heart rate, GCS, respiratory rate and O2 saturation</p> <ol style="list-style-type: none"> <li>1. Monitor for 4 hours in the ED</li> <li>2. Stepdown to normal ward bed for observation</li> <li>3. Discharge home 12 hours post last naltrexone dose, with instructions to re-present to ED if symptoms of opioid toxicity develop.</li> </ol>
<b>Management of Complications</b>	Supportive care if patients develop symptoms of opioid toxicity
<b>Basis of Protocol/Guideline:</b> (including sources of evidence, references)	<p>Aghabiklooei A, Hassanian-Moghaddam H, Zamani N, et al. Effectiveness of naltrexone in the prevention of delayed respiratory arrest in opioid- naive methadone-intoxicated patients. <i>BioMed research international</i> 2013; <b>2013</b>: 903172</p> <p>Australian Medicines Handbook – Naltrexone monograph (accessed via CIAP 30/05/2022)</p> <p>Naltrexone Product Information – last amended 1<sup>st</sup> December 2021</p>
<b>Groups consulted in development of this guideline</b>	<p>Dr Zeff Koutsogiannis (Toxicologist – Austin Hospital, Melbourne)</p> <p>Amy Minett, Acting SESLHD QUM Lead Pharmacist</p>

## AUTHORISATION

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## GOVERNANCE

Enactment date	June 2019
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**Medicine Guideline**

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