Prescribing Protocol SESLHDPR/658 Take-home naloxone under the Opioid Overdose Response and Take Home Naloxone (ORTHN) intervention (Nyxoid® or Prenoxad®)



Prescribing Protocol		
Title	Take-home naloxone under the Opioid Overdose Response and Take Home Naloxone (ORTHN) intervention (Nyxoid® or Prenoxad®)	
Areas where Protocol/ Guideline applicable	District – Outpatient and Community-based services (e.g. Drug and Alcohol, Kirketon Road Centre, Medically Supervised Injecting Centre)	
Areas where Protocol/ Guideline not applicable	Inpatient areas, 'on-site' administration by clinicians within healthcare facilities	
Authorised Prescribers	Medical Officers, Nurse Practitioners Under the NSW Health ORTHN Policy, take home naloxone can be supplied to patients as part of an intervention by an appropriately trained and credentialed health worker without a pharmacist or medical officer being present	
Indication for use	Emergency treatment for known or suspected opioid overdose, with respiratory and/or central nervous system depression *Supply restricted to non-admitted patients at risk of opioid overdose	
Clinical condition	Non-admitted patients/clients at risk of opioid overdose, assessed as engaged in recent opioid use or at significant risk of future opioid use Potential witnesses to an opioid overdose	
Contra-indications	Hypersensitivity to naloxone or any excipients	
Precautions	 Take home naloxone products are not a substitute for emergency medical care and cannot replace intravenous injection of naloxone Patients and/or their carers must receive full counselling and written information on the use of the take home naloxone product. Opioid withdrawal syndrome: naloxone administration can lead to a rapid opioid reversal and acute withdrawal syndrome. Patients receiving opioids for the relief of chronic pain may experience pain and opioid withdrawal symptoms when Nyxoid® or Prenoxad® is administered. Paediatrics: absorption may be erratic or delayed. For supply under the NSW Health ORTHN Policy, clients must be 16 years and over. Use in Pregnancy (Category B1) and breastfeeding Reversal of respiratory depression by partial agonists or mixes agonists/antagonists (e.g. buprenorphine) may be incomplete Patients who receive Nyxoid® or Prenoxad® should not drive or operate machinery or engage in other activities demanding physical or mental exertion for at least 24 hours. 	
Place in Therapy	First line	
Dosage	Nyxoid® nasal spray: 1.8 mg (1 nasal spray) administered into one nostril Prenoxad® 2mg/2mL intramuscular injection: 0.4mg (0.4mL) Further doses may be given every 2 to 3 minutes if needed until emergency care is available.	
Important Drug Interactions	Naloxone reverses the analgesic and other effects of opioid agonist analgesics, and may precipitate acute withdrawal symptoms if used concurrently with these medicines in physically dependent patients.	

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Administration instructions	primed or tested before administration of the state of the primed or tested before administration of the primed of the primed or tested before administration of the primed or tested before administration of the primed of the pri	ostril. Press firmly on the plunger until it see after 2 to 3 minutes readminister into vice. If more naloxone is available 3 minutes until person recovers or
	Give 0.4mL of Prenoxad® Injection	(to first black line on syringe) into the he patient does not respond, repeat every 2 to 3 minutes until person
Monitoring requirements	can lead to reoccurrence of respirate responded, patients and their carers	s should be advised to continue to tory depression until emergency care
Management of complications	Naloxone is NOT a substitute for emoverdose is suspected, emergency immediately.	nergency medical care. If an opioid medical assistance should be called
	(body aches, diarrhoea, tachycardia piloerection, sweating, yawning, naurestlessness or irritability, shivering	an cause an acute withdrawal syndrome in fever, runny nose, sneezing, usea or vomiting, nervousness, or trembling, abdominal cramps, ssure). If these signs and symptoms,
	Patients who are receiving opioids for experience pain and opioid withdraw Prenoxad® is administered.	•
Basis of Protocol Groups consulted in	Nyxoid Product Information, last updated 18 Nyxoid Consumer Medicine Information, upon Prenoxad Product Information last updated Prenoxad Consumer Medicines Information Consumer Information Sheet - Prenoxad® - Consumer Information Sheet - Nyoxoid® - Nyo	dated September 2018 11 December 2012 updated January 2018 NSW Ministry of health 2019
	NSW Health PD2019 036 Opioid Overdose Kirketon Road Centre	Response & Take Home Naloxone Policy
development of this protocol	Prof Nick Lintzeris, Director, SESLHD D Man Cho Leung, Senior Pharmacist, SE Amy Minett, Acting Quality Use of Medic	ESLHD Drug & Alcohol Services

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AUTHORISATION			
Author (Name)	Dr Lucy Cho		
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	GOVERNANCE		
Enactment date:	October 2019		
Updated:	July 2020 (Prenoxad information added)		
Expiry date:	July 2023		
Ratification date by SESLHD QUM Committee	2 July 2020		
Chairperson, QUM Committee	Dr Jo Karnaghan		
Version Number	2.0		

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