



WOODSTOCK HOSPITAL

Department/Category	Nursing -General			
Policy Name/ Unit Number	Clinical Opiate Withdrawal Scale (COWS) Symptom Measurement for Buprenorphine-Naloxone Induction-Unit 5			
Location	Clinical Practice Manual			
Approval Committees	<input checked="" type="checkbox"/> CPC: June 2, 2020 <input type="checkbox"/> MAC: Click here to enter a date. <input type="checkbox"/> Board of Directors: Click here to enter a date. <input type="checkbox"/> Patient and Family Advisory: Click here to enter a date. <input type="checkbox"/> Other: Click here to enter a date.		<input checked="" type="checkbox"/> NAC: June 25, 2020 <input type="checkbox"/> P and T: Click here to enter a date. <input type="checkbox"/> Senior Team: Click here to enter a date.	
Signature (if applicable)				
Document Owner Staff Development	Original Date April 2020	Reviewed Date	Revision Date	Page 1 of 5

Background:

Opioid dependence is an increasing problem for both Public Health and clinical settings in Canada. Public access issues, as well as patient disinterest limit Methadone treatment. Buprenorphine-naloxone (Suboxone) has demonstrated safety in the primary care setting and improved access in the community. Its use will reduce opioid withdrawal symptoms while discouraging misuse of drugs. For patients expressing the desire to stop using opioids, Woodstock Hospital (WH) may offer the use of Suboxone to support the patient in a timely manner. Administration through the Emergency Department (ED) will facilitate early treatment and may prevent admission to hospital. In order to accomplish this, WH will implement an assessment tool, the Clinical Opiate Withdrawal Scale (COWS), to monitor patients receiving Suboxone, either through the ED or inpatient units.

Purpose:

The COWS tool is a standardized tool used to assess for opiate withdrawal symptoms based on a rating system (See Appendix A). The resulting score provides clinicians with information on the stage and severity of opioid withdrawal and can be used for dosing of Suboxone. Withdrawal symptoms occur in stages depending on when the opioid was last used, the dose, the route, and the half-life of the medication. Withdrawal is most often identified during ED encounter, but can occur at any time during the patient hospital stay.



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The summed score for the completed scale can help clinicians determine the stage or severity of opiate withdrawal and assess the level of physical dependence on opioids.

Policy:

1. This screening tool is to be used to assess for and reassess the patient's response to Suboxone administration
2. As directed within the pathway, the Most Responsible Physician (MRP) is to be notified of any positive score, and assess the need for interventions (e.g. Suboxone administration)
3. Documentation of the COWS tool will be completed within the electronic health record (eHR)

Practice Guidelines

1. The COWS tool can be used for any patient requesting treatment for withdrawal or if the nurse suspects opioid withdrawal
2. Perform two client identifiers
3. Provide patient with explanation of the purpose of the COWS screening, frequency of assessments, medications and expected outcomes
4. Assess the symptoms of withdrawal, which can occur six hours after last dose of short acting opioid, peak at two to three days, and resolve by five to seven days. The screening of symptoms includes the following physical assessments:
 - a. Resting heart rate
 - b. GI Upset
 - c. Sweating
 - d. Tremor
 - e. Restlessness
 - f. Yawning
 - g. Pupil Size
 - h. Anxiety or Irritability
 - i. Bone or Joint Pain
 - j. Gooseflesh skin
5. Document the screening results in the eHR. In Powerchart, use the adhoc -folder above the demographic bar, then search screening tools; or in Firstnet, the ED General Information and Screening Tools Band during an ED encounter
6. Once the COWS assessment is complete, click on the TOTAL to obtain a calculated score
7. Notify the MRP of possible withdrawal symptoms and the score from the COWS tool
8. The MRP may choose to order the power plan if the patient meets the following inclusion criteria:



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- i. At least twelve hours since last dose of short acting opioid
 - ii. At least seventy-two hours since last dose of methadone
 - iii. Patient consents to taking Suboxone
9. The power plan is contraindicated if any of the following exclusion criteria are present:
 - i. Allergy to Suboxone
 - ii. Decreased level of consciousness
 - iii. Prescribed methadone or Buprenorphine/naloxone
 - iv. Inability to provide informed consent
 - v. Severe liver dysfunction
 - vi. Acute alcohol intoxication or withdrawal
 - vii. Acute respiratory distress
 - viii. Paralytic ileus
 - ix. Pregnancy
10. If COWS score is less than 12, Suboxone is not required
11. If COWS score is twelve or higher; give the Suboxone dose ordered by the MRP
 - a) Instruct patient to moisten mouth with water and swallow
 - b) Instruct patient to place Suboxone under tongue (sublingually)
 - c) Instruct patient to avoid sucking on tablet(s) or swallowing saliva during this time
 - d) Patient must be supervised while dose dissolves under the tongue for one to five minutes
12. Reassess the patient hourly using the COWS tool for a maximum of 4 hours until the max dose of 8 mg has been administered OR if score is below ten
13. Physician to refer patient to Rapid Access Addiction Medicine Clinic (RAAM)
Provide pamphlet and link to website <http://www.oxfordraam.ca> . If able, complete the intake form with the patient that is available on the website. Fax the intake form with the final Suboxone dose amount and time of discharge to the RAAM clinic.
14. If patient is stable and discharged, provide a prescription for Suboxone until next available date of RAAM clinic drop in date; with no dose provided for this date
15. Fax prescription to local pharmacy of patient's choice, and instruct patient that dose(s) will be dispensed and supervised
16. Nurse is to complete discharge Suboxone therapy checklist
17. Provide patient with RAAM clinic pamphlet
18. Provide and review the Suboxone discharge handout with the patient or family member before the leave the hospital
19. Patient may require a maintenance dose of Suboxone if remaining in hospital and a referral to RAAM clinic on discharge, with a prescription for doses until the next available clinic drop in date



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Current Reviewer:	
Responsibility:	Director of Patient Care
Reference:	<p>1) Clinical Opiate Withdrawal Scale. (2015). Retrieved February 6, 2020, from https://www.drugabuse.gov/sites/default/files/files/ClinicalOpiateWithdrawalScale.pdf</p> <p>2) Buprenorphine/Naloxone for Opioid Dependence: Clinical Practice Guideline Copyright © 2011 Centre for Addiction and Mental health From: Wesson, DR & Ling, WJ. (2003). <i>Psychoactive Drugs</i> 35(2): 253–9.</p> <p>3) LHSC Emergency Services Program Protocol for COWS and Opiate withdrawal-C Jaramillo Clinical Educator</p> <p>4) Barbosa-Leiker, C., McPherson, S., Mamey, M. R., Burns, G. L., Layton, M. E., Roll, J., & Ling, W. (2015). Examining the factor structure of the Clinical Opiate Withdrawal Scale: A secondary data analysis from the National Drug Abuse Treatment Clinical Trials Network (CTN) 0003. <i>Drug and Alcohol Dependence</i>, 152, 218–223. https://doi.org/10.1016/j.drugalcdep.2015.03.036</p>
Cross Reference:	1) Buprenorphine/Naloxone (Suboxone®), Unit 7, Clinical Practice Manual



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Appendix A: Clinical Opiate Withdrawal Scale (COWS)

Clinical Opiate Withdrawal Scale (COWS)	
For each item, select the response that best describes the patient's signs or symptom. Rate on just the apparent relationship to opiate withdrawal. For example, if the heart rate is increased because the patient was jogging just prior to the assessment, the increased pulse rate would not add to the score.	
Reason for Assessment	
Resting Pulse Rate Measure after patient is sitting or lying for one minute	<input type="radio"/> 0 - Pulse rate 80 bpm or below <input type="radio"/> 1 - Pulse rate 81-100 bpm <input type="radio"/> 2 - Pulse rate 101-120 bpm <input type="radio"/> 4 - Pulse rate greater than 120 bpm
Sweating Over past 1/2 hour not accounted for by room temperature or patient activity	<input type="radio"/> 0 - No report of chills or flushing <input type="radio"/> 1 - Subjective report of chills or flushing <input type="radio"/> 2 - Flushed or observable moistness on face <input type="radio"/> 3 - Beads of sweat on brow or face <input type="radio"/> 4 - Sweat streaming off face
Restlessness Observation during assessment	<input type="radio"/> 0 - Able to sit still <input type="radio"/> 1 - Reports difficulty sitting still, but is able to do so <input type="radio"/> 3 - Frequent shifting or extraneous movements of legs/arms <input type="radio"/> 5 - Unable to sit still for more than a few seconds
Pupil Size	<input type="radio"/> 0 - Pupils pinned or normal size for room light <input type="radio"/> 1 - Pupils possibly larger than normal for room light <input type="radio"/> 2 - Pupils moderately dilated <input type="radio"/> 5 - Pupils so dilated that only rim of the iris is visible
Bone or Joint Aches If patient was having pain previously, only the additional component attributed to opiates withdrawal is scored	<input type="radio"/> 0 - Not present <input type="radio"/> 1 - Mild diffuse discomfort <input type="radio"/> 2 - Patient reports severe diffuse aching of joints/muscles <input type="radio"/> 4 - Rubbing joints/muscles, can't sit still with discomfort
Runny Nose or Tearing Not accounted for by cold symptoms or allergies	<input type="radio"/> 0 - Not present <input type="radio"/> 1 - Nasal stuffiness or unusually moist eyes <input type="radio"/> 2 - Nose running or tearing <input type="radio"/> 4 - Nose constantly running or tears streaming down cheeks
GI Upset Over last 1/2 hour	<input type="radio"/> 0 - No GI symptoms <input type="radio"/> 1 - Stomach cramps <input type="radio"/> 2 - Nausea or loose stool <input type="radio"/> 3 - Vomiting or diarrhea <input type="radio"/> 5 - Multiple episodes of vomiting or diarrhea
Tremor Observation of outstretched hands	<input type="radio"/> 0 - No tremor <input type="radio"/> 1 - Tremor can be felt, but not observed <input type="radio"/> 2 - Slight tremor observable <input type="radio"/> 4 - Gross tremor or muscle twitching
Yawning Observation during assessment	<input type="radio"/> 0 - No yawning <input type="radio"/> 1 - Yawning once or twice during assessment <input type="radio"/> 2 - Yawning three or more times during assessment <input type="radio"/> 4 - Yawning several times/minute
Anxiety or Irritability	<input type="radio"/> 0 - None <input type="radio"/> 1 - Patient reports increasing irritability or anxiousness <input type="radio"/> 2 - Patient obviously irritable or anxious <input type="radio"/> 4 - Patient so irritable or anxious/ participation difficult
Gooseflesh Skin	<input type="radio"/> 0 - Skin is smooth <input type="radio"/> 3 - Piloerection can be felt or hairs standing up on arms <input type="radio"/> 5 - Prominent piloerection
Total Score	<input type="text"/>
Score Indicative Of	<input type="radio"/> Mild withdrawal <input type="radio"/> Moderate withdrawal <input type="radio"/> Moderately severe withdrawal <input type="radio"/> Severe withdrawal

Wesson, D.R., & Ling, W. (2003). The Clinical Opiate Withdrawal Scale (COWS). J Psychoactive Drugs, 35(2), 253-9.