Psychiatric Hospital at Vanderbilt

OPIATE WITHDRAWAL PROTOCOL - PHV: (Pregnant patients should not be withdrawn from opiates)

If patient requires detoxification from multiple CNS depressants and/or alcohol, do not use this protocol until the detoxification from these substances is completed.

- Notify MD if patient meets criteria for initiation of Opiate Withdrawal Protocol, based on the patient's report and/or documentation of opiate dependency (i.e., + history, + UDS, long term use).
- Obtain vital signs prior to each dose of clonidine and/or buprenorphine dose; then every shift after stabilization
 of acute detox.
- · Each time vital signs are obtained, assess patient using COWS Scale and document on scoring flow sheet.
- Score 5 or greater = Administer clonidine 0.1 mg po q 1hr, up to 0.4 mg in 8 hrs or less, while awake (Hold dose for the following: If baseline BP less than or equal to 100/70, hold for BP less than or equal to 80/50; If baseline BP greater than 100/70, hold for BP less than 90/60).
- After pt. has received 0.4 mg clonidine in 8 hrs or less or has scored 15 or > prior to reaching 0.4 mg clonidine = call MD for orders to start buprenorphine dosing schedule (0.3 mg lM q 8hrs, using the hours of 0600, 1400, 2200 x 9 doses).
- Continue clonidine orders even after buprenorphine injections have started, according to the parameters of the protocol.
- If score is 15 or > after dose # 3 of buprenorphine 0.3 mg, call MD for order to increase buprenorphine to 0.6 mg IM q 8hrs for doses # 4, 5, 6; then decrease back to 0.3 mg q 8hrs for doses # 7, 8, 9.
- Stop buprenorphine after dose # 9; If patient continues to score 15 or > after the 9th dose, notify MD for how to
 proceed without restarting the buprenorphine.
- Inform patient that the buprenorphine will not be restarted once it has been stopped.
- 24 hrs of observation is recommended following the last dose of buprenorphine prior to discharge.
- Encourage patient not to take any more buprenorphine than necessary, particularly if COWS score is 0 prior to the 9th dose.
- Observe patient for signs of CNS depression and, if present, notify MD.

PROTOCOL MEDICATIONS:

Clonidine:

- Clonidine 0.1mg po q1hr, up to 0.4mg within and 8 hour period, while awake, for COWS score of 5 or greater.
- <u>Hold dose for the following</u>: If baseline BP less than or equal to 100/70, hold for BP less than or equal to 80/50; If baseline BP greater than 100/70, hold for BP less than 90/60.

Buprenorphine:

- If protocol criteria met: Buprenorphine 0.3mg IM q 8 hours for 9 consecutive doses for pain
 management; on day 1, may alter dose spacing to not less than q 6 hours to adjust to 0600, 1400, 2200 dosing
 schedule.
- <u>If score 15 or > after dose #3 and initial order discontinued</u>: Buprenorphine 0.6mg IM q 8 hours x 3 doses for pain management for doses #4, 5, 6 (continue on the dosing schedule of 0600, 1400, 2200) and then decrease to 0.3mg IM for doses #7, 8, 9.

MC 8308 (3/2006) BACK

Psychiatric Hospital at Vanderbilt

| CENTRAL NERVOUS SYSTEM DEPRESSANT PROTOCOL(CNSDP)- PHV | | | | | | | | | |
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| Signature | Initials | Signature | initials | | | | | | |
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Criteria for initiation of CNSDP Protocol is based on the patient's report and/or documented history of recent or long term use of CNS depressants (hypno-sedatives, benzodiazepines, barbiturates) and/or verification of + UDS of same substances.

(Caution: Phenobarbital is not recommended for use in Pregnancy).

Protocol Instructions:

- · Notify MD for initiation of CNSDP Protocol if patient meets above criteria.
- Patient to remain on Seizure Precautions for the duration of the CNSDP Protocol.
- Assess and record vital signs and signs/symptoms q 4 hours (observe for objective signs).
- Document assessments and vital signs on the CNSDP Flowsheet.
- At the first sign of progression from mild to moderate/severe withdrawal signs, begin q 1 hour assessment and vital sign schedule.
 if all signs return to mild status (after a minimum of 2 hourly checks), resume q 4 hour assessment and vital sign schedule.
- If patient does not display symptoms of CNS depressant withdrawal within 72 hours, discontinue the protocol.
- Initiate Phenobarbital Load when patient displays at least 2 moderate to severe signs from the list below.
- Continue to assess patient and record vital signs hourly prior to the administration of each Phenobarbital dose.
- Discontinue Phenobarbital Load when patient displays 2 or more of the following symptoms, indicating intoxication:
 - > Patient asleep/difficult to arouse
 - e > Nystagmus
 - > Ataxic (unsteady gait)
- ➤ Labile Mood
- > Dysarthric (thick tongue)

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| MILD WITHDRAWAL SIGNS | DATE: | | | | | | | | | |
|--------------------------------------|-------------|-----|----------|---------|--|----------|----------|--|--------------|--|
| Mark (+) if Present | TIME: | | | . 34 | | | | | i - | |
| Blood Pressure | | A | | | | | | | - | |
| Pulse: Record beats per minute (bpm) | •••• | | | | | | | , | | |
| Mild agitation, irritability | | 3.7 | | | | | | | | |
| Mild restlessness, anxiety | | | | | İ | | | | | |
| Lightheadedness / dizziness | | | | | | | | <u> </u> | | |
| Paraesthesia (tingling sensation) | | | | | | | | 1 | | |
| Mild tremors | | | | | | | | | | |
| Nausea / anorexia | | | | | · | <u> </u> | <u> </u> | <u> </u> | | |
| Mild diaphoresis | | | | | | | | | - | |
| Insomnia | | | | | 1 | | | | <u> </u> | |
| TOTAL # Mild Sig | ns Present | | ***.**.* | at Hug. | | 177 | | | | |

| MODERATE to SEVERE WITHDRAWAL SIGNS Mark (+) if Present | ASELINI | <u> </u> | ; ; | | | | | | |
|--|---------|----------|----------|----------|--|----------|----------|---|---------------------------------------|
| Hypertension: Increase in DBP greater than or equal to 20 mm Hg in 2 hrs or less | | | | | 1 | | | | |
| Tachycardia: Increase in pulse =greater than or equal to 20 bpm in 2 hrs or less | | | | ŀ | | | | | |
| Marked increase in agitation, iπitability | | | 1 | | | 1 | | | |
| Marked increase in anxiety; very restless | | | | | 1 | <u> </u> | | | |
| Progressive confusion/disorientation | | | | 1 | 1 | | | | |
| Increasingly severe fasciculation (Muscle twitching) | | | | <u> </u> | | | 1 | | |
| Increasingly severe tremors | | | | <u> </u> | † | | | | , , , , , , , , , , , , , , , , , , , |
| Vomiting / dry heaves | | | <u> </u> | 1 | | 1 | <u> </u> | | |
| Increasingly severe diaphoresis | - | | | | | | <u> </u> | | |
| Pre-seizure activity (e.g., aura, bright lights) | | | | | <u> </u> | | | ļ | |
| Increasingly severe visual hallucinations | | | | | | | | | |
| Increasingly severe tactile hallucinations | | | | | <u> </u> | | | | |
| TOTAL # Moderate to Severe Signs Present | | | | | | | | | |
| Initials | | | | | | | | | |

| Patient has met the criteria | for initial | tion of Phenobarbital Load | d by sco | oring at | least 2 l | Moderat | e to Sev | rere Sig | ns at | L | (Tir | ne) |
|------------------------------|-------------|----------------------------|----------|----------|-----------|---------|----------|----------|-------|---|------|-----|
| on | (Date) | Initials: | | | | | | | | | N. | |