

Clinical Institute Withdrawal Assessment for Alcohol–Revised might be an unreliable tool in the management of alcohol withdrawal

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The Clinical Institute Withdrawal Assessment for Alcohol–Revised (CIWA–Ar) protocol (**Figure 1**)¹ is the most common method of treating alcohol withdrawal in our institution and it is frequently used by family physicians. Although various rating scales for alcohol withdrawal have been described, the CIWA–Ar protocol managing withdrawal with benzodiazepines is well established.^{2–4} Symptom-triggered benzodiazepine dosing has been demonstrated to lead to shorter duration of treatment and lower medication use compared with fixed-schedule dosing.⁵ Although the CIWA–Ar protocol was validated in medically cleared patients in an alcohol detoxification setting, it has also been evaluated in hospital settings.^{1,6–8} However, the application of the CIWA–Ar needs to be carefully considered, and inappropriate use of the protocol has been documented.³ This article describes a case in which an objective alcohol withdrawal scale (OAWS) was more useful for treatment, as the CIWA–Ar could not be applied.

EDITOR'S KEY POINTS

- Because accurate application of the Clinical Institute Withdrawal Assessment for Alcohol–Revised requires taking a detailed medical history, it should not be used when a substantial language barrier exists, or when patients cannot provide a reliable history because of delirium, dementia, psychosis, etc.
- An objective alcohol withdrawal scale can be tailored to comorbidities and severity of withdrawal, but it has not been validated as an alternative to the Clinical Institute Withdrawal Assessment for Alcohol–Revised protocol. It is intended as an approach to treatment that can be useful when validated protocols cannot reliably be applied.

POINTS DE REPÈRE DU RÉDACTEUR

- Étant donné que l'application exacte de l'échelle des symptômes de sevrage de l'alcool (Clinical Institute Withdrawal Assessment for Alcohol–Revised ou CIWA–AR) exige une anamnèse détaillée, cette échelle ne devrait pas être utilisée s'il existe des barrières de langue importantes ou lorsque les patients ne peuvent pas expliquer leurs antécédents médicaux de manière fiable à cause d'un delirium, d'une démence, d'une psychose et ainsi de suite.
- Une échelle objective des symptômes de sevrage de l'alcool peut être adaptée aux comorbidités et à la gravité des symptômes, mais elle n'a pas été validée comme solution de rechange au protocole CIWA–AR. Elle a pour but de servir d'approche thérapeutique susceptible d'être utile lorsque les protocoles validés ne peuvent pas être appliqués de manière fiable.

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Case

A 57-year-old Polish man presented to an urban hospital at 12:18 PM after falling while intoxicated. His Glasgow Coma Scale score was 11 and his serum ethanol level was 75 mmol/L at 3:53 PM. An eyebrow laceration was sutured, and a computed tomography scan of his head showed a trace subarachnoid hemorrhage. After review by the neurosurgery department, the patient was kept for observation and began to exhibit signs of alcohol withdrawal. A CIWA–Ar protocol using lorazepam was initiated at 8:50 PM with an initial score of 13. Upon reassessment at 7:41 AM, the patient had received a total of 10 mg of oral lorazepam and a consultation with the internal medicine department was initiated. The patient's withdrawal continued to worsen, and lorazepam was switched to diazepam. When seen in the internal medicine department at 3:45 PM, a regular 10-mg dose of oral diazepam 3 times a day was added, and a consultation with the Addiction Medicine Consult Team (AMCT) was requested.

The patient was seen by the AMCT at 6:00 PM, 30 hours into the withdrawal process and 21 hours since starting the CIWA–Ar protocol. The patient had received a total of 18 mg of oral lorazepam, 40 mg of intravenous diazepam, and 20 mg of oral diazepam, and he continued to exhibit signs of severe alcohol withdrawal including agitation, diaphoresis, hypertension, tachycardia, and tremor. He was unable to converse in English, although he was able to speak Polish when a telephone translation service was briefly available. He was confused and disoriented to time and place. There was minimal collateral history, with no previous admissions, no pharmacy records, no next of kin available, and a retired family physician on record. He had signs of chronic liver disease including clubbing, palmar erythema, and a palpable liver. Given the patient's inability to converse, the AMCT discontinued the CIWA–Ar protocol and constructed an OAWS (**Box 1**).

Because of the apparent liver disease, diazepam was changed to 1 mg of oral lorazepam (which does not require hepatic oxidation) 4 times a day, with lorazepam as needed based on the OAWS.

The patient's withdrawal improved and his lorazepam requirements gradually declined. On day 4, scheduled lorazepam was decreased to 3 times a day, then to twice a day on day 5. The patient remained hypertensive and received sporadic as-needed doses because of this. On day 6, the OAWS and treatment with lorazepam was discontinued and the patient was discharged in stable condition on day 8. Recommendations were provided to the patient's Polish-speaking family physician regarding relapse prevention medications, with acamprosate being the drug of choice given his liver dysfunction.

Discussion

An important limitation of the CIWA-Ar is its heavily subjective nature. Only 3 of 10 components (tremor, paroxysmal sweats, agitation) can be rated by observation alone. The other 7 components require at least some discussion with the patient. Given that benzodiazepines are provided based on the CIWA-Ar score, there is risk of incorrect dosing when scores are unreliable, which harbours potential for patient harm. There are 2 primary reasons why the CIWA-Ar was unreliable in this case. First, there was a substantial language barrier preventing the discussions necessary for accurate scoring. This became clearer as the patient's withdrawal improved and he was still unable to answer simple questions in English. Even with an interpreter available, CIWA-Ar might remain impractical, as it requires frequent reassessments and would necessitate 24-hour interpreter coverage. The second limitation of the CIWA-Ar was subtler; the patient was confused and disoriented, so even in the absence of a communication barrier, his responses might have been unreliable. In a hospitalized population this might be a common scenario; acute medical issues can contribute to delirium and complicate the clinical picture.

While this case illustrates 2 reasons to use an OAWS, other common reasons exist. These might include patients with a clouded sensorium from acute psychosis or severe dementia, those with mechanical communication problems including severe facial trauma limiting speech and vision, and those with intubation.

Alternative assessment tools and clinical pathways have been proposed for inpatient management of alcohol withdrawal, but like the CIWA-Ar, they often require a reliable history.^{9,10} The approach presented here has proved reliable for treatment of complex alcohol

Box 1. Objective alcohol withdrawal scale

The objective alcohol withdrawal scale is applied as follows:

- Score 1 point for each of
 - systolic blood pressure \geq 160 mm Hg or diastolic blood pressure \geq 90 mm Hg;
 - heart rate \geq 90 beats/min;
 - tremor;
 - diaphoresis; and
 - agitation
- If total \geq 2 give 1 mg oral lorazepam (or 10 mg of diazepam)
- If total \geq 3 give 2 mg oral lorazepam (or 20 mg of diazepam)
- Reassess every hour until score is $<$ 2 for 3 consecutive measures, then reassess every 6 hours for 24 hours, then every 24 hours for 72 hours, then discontinue

Figure 1. The Clinical Institute Withdrawal Assessment for Alcohol–Revised scale

Appendix: Addiction Research Foundation Clinical Institute Withdrawal Assessment for Alcohol (CIWA-Ar)

Patient _____ Date |_|_|_|_| Time _____ : _____
 y m d (24 hour clock, midnight=00:00)

Pulse or heart rate, taken for one minute: _____ Blood pressure: _____/_____

NAUSEA AND VOMITING—As “Do you feel sick to your stomach? Have you vomited?” Observation.
 0 no nausea and no vomiting
 1 mild nausea with no vomiting
 2
 3
 4 intermittent nausea with dry heaves
 5
 6
 7 constant nausea, frequent dry heaves and vomiting

TREMOR—Arms extended and fingers spread apart. Observation.
 0 no tremor
 1 not visible, but can be felt fingertip to fingertip
 2
 3
 4 moderate, with patient’s arms extended
 5
 6
 7 severe, even with arms not extended

PAROXYSMAL SWEATS—Observation.
 0 no sweat visible
 1 barely perceptible sweating, palms moist
 2
 3
 4 beads of sweat obvious on forehead
 5
 6
 7 drenching sweats

ANXIETY—Ask “Do you feel nervous?” Observation.
 0 no anxiety, at ease
 1 mildly anxious
 2
 3
 4 moderately anxious, or guarded, so anxiety is inferred
 5
 6
 7 equivalent to acute panic states as seen in severe delirium or acute schizophrenic reactions

AGITATION—Observation.
 0 normal activity
 1 somewhat more than normal activity
 2
 3
 4 moderately fidgety and restless
 5
 6
 7 paces back and forth during most of the interview, or constantly thrashes about

TACTILE DISTURBANCES—Ask “Have you any itching, pins and needles sensations, any burning, any numbness or do you feel bugs crawling on or under your skin?” Observation.
 0 none
 1 very mild itching, pins and needles, burning or numbness
 2 mild itching, pins and needles, burning or numbness
 3 moderate itching, pins and needles, burning or numbness
 4 moderately severe hallucinations
 5 severe hallucinations
 6 extremely severe hallucinations
 7 continuous hallucinations

AUDITORY DISTURBANCES—Ask “Are you more aware of sounds around you? Are they harsh? Do they frighten you? Are you hearing anything that is disturbing to you? Are you hearing things you know are not there?” Observation.
 0 not present
 1 very mild harshness or ability to frighten
 2 mild harshness or ability to frighten
 3 moderate harshness or ability to frighten
 4 moderately severe hallucinations
 5 severe hallucinations
 6 extremely severe hallucinations
 7 continuous hallucinations

VISUAL DISTURBANCES—Ask “Does the light appear to be too bright? Is its colour different? Does it hurt your eyes? Are you seeing anything that is disturbing to you? Are you seeing things you know are not there?” Observation.
 0 not present
 1 very mild sensitivity
 2 mild sensitivity
 3 moderate sensitivity
 4 moderately severe hallucinations
 5 severe hallucinations
 6 extremely severe hallucinations
 7 continuous hallucinations

HEADACHE, FULLNESS IN HEAD—Ask “Does your head feel different? Does it feel like there is a band around your head?” Do not rate for dizziness or lightheadedness. Otherwise, rate severity.
 0 not present
 1 very mild
 2 mild
 3 moderate
 4 moderately severe
 5 severe
 6 very severe
 7 extremely severe

ORIENTATION AND CLOUDING OF SENSORIUM—Ask “What day is this? Where are you? Who am I?”
 0 oriented and can do serial additions
 1 cannot do serial additions or is uncertain about date
 2 disoriented for date by no more than 2 calendar days
 3 disoriented for date by more than 2 calendar days
 4 disoriented for place and/or person

Total CIWA-A Score _____
 Rater’s Initials _____
 Maximum Possible Score 67

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withdrawal by a busy AMCT in a tertiary Canadian hospital. The OAWS in **Box 1** is not intended as an alternative to the CIWA-Ar and therefore has not been validated as such. Rather, it is an approach to treatment that can be useful when other validated tools cannot be reliably applied. It is based on objective findings and can be modified to fit the clinical situation. For example, in a patient with poorly controlled hypertension, blood pressure could be excluded or a higher blood pressure cutoff chosen. Similarly, heart rate might be excluded for a patient with uncontrolled atrial fibrillation or sepsis, and tremor excluded for a patient with essential tremor or parkinsonism. Additionally, the OAWS can be modified by changing the cutoff for scores prompting doses of benzodiazepines. In this case, the patient was unwell and required high doses, so we opted for a liberal scale to minimize underdosing. In more moderate withdrawal or where there was concern for benzodiazepine toxicity, cutoffs of 3 or more, or 4 or more, would be more benzodiazepine sparing.

Conclusion

The OAWS can be useful for cases of alcohol withdrawal in which the CIWA-Ar is unreliable. The OAWS can be used

as a framework and tailored to individual cases with consideration of comorbidities and withdrawal severity. 🌿

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Competing interests

None declared

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