Alcohol Withdrawal Syndrome: Outpatient Management

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Approximately one-half of patients with alcohol use disorder who abruptly stop or reduce their alcohol use will develop signs or symptoms of alcohol withdrawal syndrome. The syndrome is due to overactivity of the central and autonomic nervous systems, leading to tremors, insomnia, nausea and vomiting, hallucinations, anxiety, and agitation. If untreated or inadequately treated, withdrawal can progress to generalized tonic-clonic seizures, delirium tremens, and death. The three-question Alcohol Use Disorders Identification Test–Consumption and the Single Alcohol Screening Question instrument have the best accuracy for assessing unhealthy alcohol use in adults 18 years and older. Two commonly used tools to assess withdrawal symptoms are the Clinical Institute Withdrawal Assessment for Alcohol Scale, Revised, and the Short Alcohol Withdrawal Scale. Patients with mild to moderate withdrawal symptoms without additional risk factors for developing severe or complicated withdrawal should be treated as outpatients when possible. Ambulatory withdrawal treatment should include supportive care and pharmacotherapy as appropriate. Mild symptoms can be treated with carbamazepine or gabapentin. Benzodiazepines are first-line therapy for moderate to severe symptoms, with carbamazepine and gabapentin as potential adjunctive or alternative therapies. Physicians should monitor outpatients with alcohol withdrawal syndrome daily for up to five days after their last drink to verify symptom improvement and to evaluate the need for additional treatment. Primary care physicians should offer to initiate long-term treatment for alcohol use disorder, including pharmacotherapy, in addition to withdrawal management. (Am Fam Physician. 2021;104(3):253-262. Copyright © 2021 American Academy of Family Physicians.)

Alcohol-related disorders cause significant physical, psychological, and societal harm. Diagnostic criteria have integrated alcohol abuse and dependence into a single disorder: alcohol use disorder (AUD; Table 1). AUD has an estimated 12-month and lifetime prevalence of 13.9% and 29.1%, respectively.1 Key management principles include promptly recognizing and evaluating for alcohol withdrawal syndrome (AWS), establishing a treatment and monitoring plan, and providing medications and resources to support long-term abstinence.

Recognizing Patients at Risk for AUD
The U.S. Preventive Services Task Force recommends that primary care physicians screen patients 18 years and older for unhealthy alcohol use and offer appropriate behavioral counseling as indicated.3 Several screening instruments can be used to identify hazardous and harmful drinking behaviors. The three-question Alcohol Use Disorders Identification Test–Consumption (AUDIT-C; https://www.mdcalc.com/audit-c-alcohol-use) and the Single Alcohol Screening Question instrument have the best accuracy for assessing unhealthy alcohol use in adults 18 years and older. Two commonly used tools to assess withdrawal symptoms are the Clinical Institute Withdrawal Assessment for Alcohol Scale, Revised, and the Short Alcohol Withdrawal Scale. Patients with mild to moderate withdrawal symptoms without additional risk factors for developing severe or complicated withdrawal should be treated as outpatients when possible. Ambulatory withdrawal treatment should include supportive care and pharmacotherapy as appropriate. Mild symptoms can be treated with carbamazepine or gabapentin. Benzodiazepines are first-line therapy for moderate to severe symptoms, with carbamazepine and gabapentin as potential adjunctive or alternative therapies. Physicians should monitor outpatients with alcohol withdrawal syndrome daily for up to five days after their last drink to verify symptom improvement and to evaluate the need for additional treatment. Primary care physicians should offer to initiate long-term treatment for alcohol use disorder, including pharmacotherapy, in addition to withdrawal management. (Am Fam Physician. 2021;104(3):253-262. Copyright © 2021 American Academy of Family Physicians.)

WHAT’S NEW ON THIS TOPIC

Alcohol Use Disorder and Withdrawal
In a randomized controlled trial, patients who started taking gabapentin (Neurontin) after 3 days of abstinence had fewer heavy drinking days (defined as 5 or more drinks for men and 4 or more drinks for women) and greater rates of abstinence than those who received placebo.

A Cochrane review found that Alcoholics Anonymous and other 12-step facilitation programs that follow a specific manual or syllabus are more effective at increasing rates of abstinence at 1, 2, and 3 years than motivational interviewing or cognitive behavior therapy, with substantial cost savings.
ALCOHOL WITHDRAWAL SYNDROME

TABLE 1

**DSM-5 Diagnostic Criteria for Alcohol Use Disorder**

A. A problematic pattern of alcohol use leading to clinically significant impairment or distress, as manifested by at least two of the following, occurring within a 12-month period:

1. Alcohol is often taken in larger amounts or over a longer period than was intended.
2. There is a persistent desire or unsuccessful efforts to cut down or control alcohol use.
3. A great deal of time is spent in activities necessary to obtain alcohol, use alcohol, or recover from its effects.
4. Craving, or a strong desire or urge to use alcohol.
5. Recurrent alcohol use resulting in a failure to fulfill major role obligations at work, school, or home.
6. Continued alcohol use despite having persistent or recurrent social or interpersonal problems caused or exacerbated by the effects of alcohol.
7. Important social, occupational, or recreational activities are given up or reduced because of alcohol use.
8. Recurrent alcohol use in situations in which it is physically hazardous.
9. Alcohol use is continued despite knowledge of having a persistent or recurrent physical or psychological problem that is likely to have been caused or exacerbated by alcohol.
10. Tolerance, as defined by either of the following:
   a. A need for markedly increased amounts of alcohol to achieve intoxication or desired effect.
   b. A markedly diminished effect with continued use of the same amount of alcohol.

Specify if:

**In early remission:** After full criteria for alcohol use disorder were previously met, none of the criteria for alcohol use disorder have been met for at least 3 months but for less than 12 months (with exception that Criterion A4, “Craving, or a strong desire or urge to use alcohol,” may be met).

**In sustained remission:** After full criteria for alcohol use disorder were previously met, none of the criteria for alcohol use disorder have been met at any time during a period of 12 months or longer (with the exception that Criterion A4, “Craving, or a strong desire or urge to use alcohol,” may be met).

Specify if:

**In a controlled environment:** This additional specifier is used if the individual is in an environment where access to alcohol is restricted.

Specify current severity:

- **Mild:** Presence of 2 to 3 symptoms
- **Moderate:** Presence of 4 to 5 symptoms
- **Severe:** Presence of 6 or more symptoms

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**TABLE 2**

**SASQ**

The SASQ has a sensitivity of 73% to 88% and specificity of 74% to 100% for detecting unhealthy alcohol use. The question, “How many times in the past year have you had X or more drinks in a day?” where X is five for men and four for women, is used in the screening. Any response greater than one is considered positive. In comparison, the full AUDIT is less sensitive (73.9%) but more specific (82.8%) at detecting unhealthy alcohol use.

Despite the high prevalence of AUD, many patients are undertreated partly because of the stigma associated with the diagnosis. For patients who are reluctant to tell their physician about their alcohol consumption, the National Institute on Alcohol Abuse and Alcoholism has a website, Rethinking Drinking (https://www.rethinkingdrinking.niaaa.nih.gov/), that provides assessment and motivational tools for moderation and abstinence treatment resources.

**Diagnostic Criteria**

Approximately one-half of patients with AUD who abruptly reduce or abstain from alcohol use experience signs or symptoms of AWS. When patients abruptly stop drinking or reduce their alcohol intake after a prolonged period (more than two weeks) of heavy use, withdrawal symptoms begin within six to 24 hours. Withdrawal effects are primarily due to the unmasking of the adaptive responses to chronic alcohol use, including decreased inhibitory activity of alpha-2 receptors, resulting in increased catecholamine levels on presynaptic neurons. Table 2 outlines the diagnostic criteria for alcohol withdrawal.
Assessing Severity

The patient’s symptom severity should be evaluated using a validated scale to determine the risk of severe or complicated AWS. The syndrome is classified as mild, moderate, severe, and complicated by the most recent guideline from the American Society of Addiction Medicine. Patients with mild AWS tend to have mild to moderate anxiety, sweating, and insomnia, but tremor is absent. Moderately severe AWS causes moderate anxiety, sweating, insomnia, and mild tremor. Those with severe AWS experience severe anxiety and moderate to severe tremor, but they do not have confusion, hallucinations, or seizures. Complicated AWS is identified by seizures or signs and symptoms indicative of delirium, such as the inability to fully comprehend instructions, clouding of the sensorium, confusion, or new onset of hallucinations. Correlating the patient’s symptoms in relation to the time since their last drink is useful in anticipating the progression of symptoms. When not properly treated, AWS can progress to delirium tremens (Table 3). The two most commonly used tools to assess withdrawal symptoms are the Clinical Institute Withdrawal Assessment for Alcohol Scale, Revised (CIWA-Ar; https://www.mdcalc.com/ciwa-ar-alcohol-withdrawal) and the Short Alcohol Withdrawal Scale (SAWS). The CIWA-Ar is a 10-item questionnaire completed by a physician that assesses the signs, symptoms, and severity of alcohol withdrawal to guide benzodiazepine dosing as part of symptom-triggered dosing regimens (Figure 1). The SAWS is a validated 10-item symptom checklist designed to be a self-assessment completed by the patient (Figure 2). The initial assessment can help to determine appropriate treatment setting and to monitor symptom improvement.

Physicians may consider laboratory testing that includes a comprehensive metabolic panel (basic metabolic profile and hepatic panel), a complete blood count with differential, urine drug screen, and blood alcohol level. These results also aid in identifying the risk of severe or complicated withdrawal and determining the appropriate disposition of care.

Treatment

Treatment goals include reducing withdrawal symptoms, preventing seizures, and preventing

### TABLE 2

<table>
<thead>
<tr>
<th>DSM-5 Diagnostic Criteria for Alcohol Withdrawal</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. Cessation of (or reduction in) alcohol use that has been heavy and prolonged.</td>
</tr>
<tr>
<td>B. Two (or more) of the following, developing within several hours to a few days after the cessation of (or reduction in) alcohol use described in Criterion A:</td>
</tr>
<tr>
<td>1. Autonomic hyperactivity (e.g., sweating or pulse rate greater than 100 beats per minute).</td>
</tr>
<tr>
<td>2. Increased hand tremor.</td>
</tr>
<tr>
<td>3. Insomnia.</td>
</tr>
<tr>
<td>4. Nausea or vomiting.</td>
</tr>
<tr>
<td>5. Transient visual, tactile, or auditory hallucinations or illusions.</td>
</tr>
<tr>
<td>6. Psychomotor agitation.</td>
</tr>
<tr>
<td>7. Anxiety.</td>
</tr>
<tr>
<td>C. The signs or symptoms in Criterion B cause clinically significant distress or impairment in social, occupational, or other important areas of functioning.</td>
</tr>
<tr>
<td>D. The signs or symptoms are not attributable to another medical condition and are not better explained by another mental disorder, including intoxication or withdrawal from another substance.</td>
</tr>
</tbody>
</table>

**Specify if:**

With perceptual disturbances: This specifier applies in the rare instance when hallucinations (usually visual or tactile) occur with intact reality testing, or auditory, visual, or tactile illusions occur in the absence of a delirium.

DSM-5 = Diagnostic and Statistical Manual of Mental Disorders, 5th ed.

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### TABLE 3

<table>
<thead>
<tr>
<th>Expected Symptoms of Alcohol Withdrawal Syndrome after Cessation of Alcohol Use</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Symptoms</strong></td>
</tr>
<tr>
<td>Anorexia, diaphoresis, gastrointestinal upset, headache, insomnia, mild anxiety, palpitations, tremulousness</td>
</tr>
<tr>
<td>Alcoholic hallucinosis: auditory, tactile, or visual hallucinations</td>
</tr>
<tr>
<td>Withdrawal seizures: generalized tonic-clonic seizures</td>
</tr>
<tr>
<td>Alcohol withdrawal delirium (delirium tremens): agitation, diaphoresis, disorientation, hallucinations (predominantly visual), hypertension, low-grade fever, tachycardia</td>
</tr>
</tbody>
</table>

*—Symptoms generally resolve within 48 hours.  |
†—Symptoms reported as early as 2 hours after cessation.  |
‡—Symptoms peak at 5 days.  |
Information from references 8-10.
**图1**

临床机构戒断评估量表（修订版）（CIWA-Ar）用于评估酒精戒断的严重程度。

- **轻度症状**：评分 < 10
- **中度症状**：评分 10 到 18
- **严重症状或复杂情况**：评分 ≥ 19


<table>
<thead>
<tr>
<th>患者：</th>
<th>日期：</th>
<th>时间：</th>
<th>(24小时制，午夜=00:00)</th>
<th>脉搏或心率，每分钟一次：</th>
<th>血压：</th>
</tr>
</thead>
</table>

### 呕吐和恶心

询问“你感觉不舒服吗？有呕吐吗？”观察。

0. 无恶心和呕吐
1. 轻度恶心，无呕吐
2. 间歇性恶心，伴有恶心
5. 恶心，频繁干呕
7. 持续恶心，频繁干呕

### 恐慌

观察。

0. 正常活动
1. 略大于正常活动
2. 3
4. 稍有不安，持续性不安
5. 6
7. 持续不安，持续不安

### 视频的扰乱

询问“周围的光线看起来太亮了吗？颜色不一样吗？眼睛疼吗？你看到什么让你感到不安的吗？你看到的东西你知道不在那里吗？”观察。

0. 无
1. 轻微不适
2. 轻微不适
3. 中度不适
4. 中度严重心理紊乱
5. 严重心理紊乱
6. 极度严重心理紊乱
7. 连续性心理紊乱

### 触觉的扰乱

询问“你有没有觉得痒、刺痛、烧灼或麻木，或者你觉得有虫子在你皮肤上爬？”观察。

0. 无
1. 轻微不适
2. 轻微不适
3. 中度不适
4. 中度严重心理紊乱
5. 严重心理紊乱
6. 极度严重心理紊乱
7. 连续性心理紊乱

### 头痛

询问“你的头感觉不同吗？你的头看起来像头带子吗？”否则，根据严重程度评分。

0. 无
1. 轻微
2. 轻微
3. 中度
4. 中度严重
5. 严重
6. 极度严重
7. 极度严重

### 注意力和思维意识混乱

询问“今天是哪天？我在哪里？我是谁？”

0. 有意识并能做连续的加法
1. 无法做连续的加法或对日期不确定
2. 混淆日期超过两天
3. 混淆地点和人物

### 总CIWA-Ar评分：

### 评语的评语：

总可能得分为67

**临床机构戒断评估量表（修订版）（CIWA-Ar）用于评估酒精戒断的严重程度。**

轻度症状：评分 < 10；中度症状：评分 10到18；严重症状或复杂情况：评分 > 19。

ALCOHOL WITHDRAWAL SYNDROME

progression to delirium tremens, which has a 5% to 10% mortality rate.8-10 In appropriate patients, outpatient treatment for AWS is safe and may be preferred. A randomized trial (n = 164) comparing outpatient and inpatient treatment of male veterans with mild to moderate withdrawal symptoms found the duration of therapy was shorter for outpatients.13 The American Society of Addiction Medicine guidelines define two levels of outpatient care: Level 1 Withdrawal Management and Level 2 Withdrawal Management.8 The two levels differ in their monitoring capabilities, with Level 1 Withdrawal Management being a typical outpatient clinic and Level 2 Withdrawal Management having extended on-site monitoring outside the scope of most primary care clinics. Level 2 Withdrawal Management facilities, including day hospitals, mental health facilities, and addiction treatment facilities, can monitor each patient for several hours each day and have greater access to psychological or psychiatric specialty treatments. Generally, patients experiencing mild symptoms (CIWA-Ar score less than 10) can be managed in Level 1 or Level 2 Withdrawal Management. For those with moderate symptoms, particularly those at increased risk of complicated withdrawal, a Level 2 Withdrawal Management facility may be more appropriate (Figure 3). Those with severe or complicated symptoms or additional risk factors should be treated in an inpatient facility (Table 4).

MILD SYMPTOMS (CIWA-AR SCORE LESS THAN 10 OR SAWS SCORE LESS THAN 12)

For patients with mild symptoms who are at minimal risk of developing severe or complicated alcohol withdrawal, treatment options include supportive care with or without pharmacotherapy (Table 5).8,14 Patients at minimal risk have none of the following factors: history of alcohol withdrawal–related delirium or seizures, multiple prior withdrawal episodes, comorbid illness, age older than 65 years, long duration of alcohol consumption (e.g., heavy alcohol use five or more days in the past month),15 seizures during current withdrawal episode, marked autonomic hyperactivity on presentation, and physiologic dependence on GABAergic agents.8

Supportive care includes educating patients on the course of withdrawal, monitoring for severe withdrawal, instructing them on how to maintain low-stimulation home environments, consuming noncaffeinated fluids, recommending a daily multivitamin containing 400 mcg of folic acid, and prescribing thiamine (typical dosage of 100 mg daily for three to five days).16,17 If medications are used, carbamazepine (Tegretol) and gabapentin (Neurontin) are appropriate options for monotherapy but do not reliably prevent withdrawal seizures or delirium tremens.18 Gabapentin is effective in treating AUD; patients already taking it should continue during treatment of AWS.18,19

MEDIUM SYMPTOMS (CIWA-AR SCORE OF 10 TO 18 OR SAWS SCORE GREATER THAN 12)

Benzodiazepines are a first-line therapy for patients experiencing moderate withdrawal symptoms, reducing the risk of seizure and the development of delirium tremens.20,21 (Table 5). Benzodiazepine dosing can be either fixed or symptom triggered. Fixed dosing sets a specific dose and time and is gradually tapered on a set schedule. Symptom-triggered dosing is given as needed based on specific CIWA-Ar or SAWS scores. Symptom-triggered use of benzodiazepines is preferred when the patient or caregiver can reliably assess symptoms and follow the dosing guidelines.22

Long-acting benzodiazepines, such as clorazepate (Librium) and diazepam (Valium), can help control and minimize breakthrough symptoms and are preferred.

FIGURE 2

<table>
<thead>
<tr>
<th>Item</th>
<th>None (0 points)</th>
<th>Mild (1 point)</th>
<th>Moderate (2 points)</th>
<th>Severe (3 points)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anxious</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Feeling confused</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Restless</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Miserable</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Problems with memory</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tremor (shakes)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nausea</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heart pounding</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sleep disturbance</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sweating</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

Short Alcohol Withdrawal Scale to assess severity of alcohol withdrawal.

Mild symptoms: score < 12; moderate to severe symptoms: score > 12.

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over short-acting benzodiazepines. Patients should be monitored for oversedation and respiratory depression, especially if concomitant liver disease is present. Physicians should consider benzodiazepines with less hepatic metabolism, such as lorazepam (Ativan) and oxazepam (Serax), in patients with liver disease. If contraindications to

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**FIGURE 3**

Suspected AWS and/or meets diagnostic criteria (Table 2)

Risk factors for developing severe or complicated withdrawal (Table 4)?

Yes

No

Consider laboratory tests as clinically indicated

Laboratory results within normal limits

Determine withdrawal severity

Laboratory results:
- Abnormal electrolytes
- Elevated AST/ALT
- Elevated BUN/Cr ratio
- Elevated blood alcohol level
- Positive urine drug screen

Consider inpatient management

Suspected AWS and/or meets diagnostic criteria (Table 2)

Risk factors for developing severe or complicated withdrawal (Table 4)?

Yes

No

Consider laboratory tests as clinically indicated

Laboratory results within normal limits

Determine withdrawal severity

Laboratory results:
- Abnormal electrolytes
- Elevated AST/ALT
- Elevated BUN/Cr ratio
- Elevated blood alcohol level
- Positive urine drug screen

Consider inpatient management

Mild (CIWA-Ar score < 10)

Moderate (CIWA-Ar score 10 to 18)

Severe or complicated (CIWA-Ar score ≥ 19)

Inpatient management

Level 2 Withdrawal Management criteria

Presence of:
- Physiologic dependence on opioids or opioid use disorder
- Severe withdrawal < 1 year ago
- Older age or history of epilepsy
- Mild or stable psychiatric symptoms

Presence of the following:
- Medical or psychiatric condition requiring inpatient treatment
- Unstable chronic condition
- Inability to tolerate oral drugs
- Severe psychiatric/cognitive impairment
- Imminent risk of harm

Inpatient management

Low risk of developing severe symptoms and does not meet Level 2 Withdrawal Management criteria except for mild or stable psychiatric symptoms (e.g., having a limited nonsignificant effect on daily functioning)

Level 1 Withdrawal Management

Supportive care

Can prescribe gabapentin (Neurontin) or carbamazepine (Tegretol)

Level 2 Withdrawal Management

Prescribe first-line benzodiazepines: chlordiazepoxide (Librium), diazepam (Valium), lorazepam (Ativan)

Prescribe gabapentin, carbamazepine, and valproate (Depacon) as adjuncts if symptoms persist despite adequate benzodiazepine use

Algorithm for AWS management.

Information from reference 8.
ALCOHOL WITHDRAWAL SYNDROME

benzodiazepines exist or if the risk of use outweighs the benefits, gabapentin, carbamazepine, and phenobarbital may be considered as alternative monotherapies.\textsuperscript{8,18}

SEVERE OR COMPLICATED SYMPTOMS (CIWA-AR \textsuperscript{19} Score of 19 or More)

Those with severe or complicated symptoms should be referred to the nearest emergency department for inpatient hospitalization.

OTHER MEDICATIONS

Gabapentin, carbamazepine, and valproate (Depo) may be used adjunctively with benzodiazepines if symptoms persist despite adequate use.\textsuperscript{19} Valproate is currently not recommended as monotherapy for the treatment of alcohol withdrawal.\textsuperscript{26} Alpha-adrenergic agonists (e.g., clonidine) and beta-blocker agonists (e.g., atenolol, metoprolol) may also be used adjunctively with benzodiazepines for persistent hypertension or tachycardia.\textsuperscript{27,28} Evidence does not support the use of oral or intravenous alcohol, butalbocen (Lioresal), or magnesium in prophylaxis and treatment of AWS.\textsuperscript{21,29,30} Table 5 provides a list of several medications used to treat moderate AWS.\textsuperscript{8,14}

Monitoring and Follow-up

The frequency and setting for outpatient monitoring of AWS should be guided by symptom severity, risk of complications, and social factors, including reliable social support and a safe home environment. Most patients will require daily evaluations for up to five days after their last drink, but evaluations may increase or decrease in frequency as necessitated by changes in symptom severity.\textsuperscript{8} These visits can be with any health care professional. Face-to-face visits are preferred, but telemedicine appointments can alternate with in-person visits when necessary, for instance during treatment in a viral pandemic.\textsuperscript{8} Evaluation should include multiple indicators of symptom severity and overall health, including mental status, hydration, sleep, mood, suicidality, and substance use. Blood pressure, pulse, and alcohol breath analysis should be obtained whenever possible. The assessment should also include a validated measure of withdrawal symptom severity, ideally with the same instrument as the initial assessment.

Continued symptoms despite multiple doses of the prescribed medication, worsening or severe symptoms (persistent vomiting, hallucinations, confusion, or seizure), signs of oversedation, worsening psychiatric symptoms, or unstable vital signs should prompt transfer to a higher level of care. Symptoms outside of the anticipated withdrawal period or resumption of alcohol use also warrants referral to an addiction specialist or inpatient treatment program.

Supporting Long-Term Abstinence

Long-term treatment of AUD should begin concurrently with the management of AWS.\textsuperscript{8} Successful long-term treatment includes evidence-based community resources and pharmacotherapy. Primary care physicians should offer to initiate appropriate medications.

Currently, three medications are approved by the U.S. Food and Drug Administration for AUD treatment: acamprosate, naltrexone (Revia), and disulfiram (Antabuse). Acamprosate and naltrexone have the best evidence for use in AUD and should be initiated in patients who wish to reduce or abstain from alcohol use.\textsuperscript{31} Neither of the medications has been shown consistently to be more effective than the other, and thus the choice should be individualized based on patient and physician preference.\textsuperscript{31} Disulfiram should be considered for patients who have not responded to acamprosate or naltrexone.\textsuperscript{32}

Gabapentin and topiramate (Topamax), though not approved for this use, may be considered as second-line treatments.\textsuperscript{33} In a randomized controlled trial, patients who started taking gabapentin after three days of abstinence had fewer heavy drinking days (defined as five or more drinks for men and four or more drinks for women) and greater

\begin{table}[h]
\centering
\begin{tabular}{|l|}
\hline
Factors for Hospitalization in Patients with AWS \\
\hline Absence of caregiver support, unstable dwelling situation \\
Active psychiatric conditions \\
Consumption of > 8 drinks per day \\
Failure to benefit from ambulatory treatment \\
History of severe alcohol withdrawal < 1 year ago \\
Medical conditions (e.g., non-alcohol-related seizure disorder, clinically significant abnormal results on laboratory testing, unstable chronic condition, inability to tolerate anything by mouth, suspected head injury) \\
Physiologic dependence on opioids or other substances \\
Risk of imminent relapse, harm to self or others, or low commitment and questionable cooperativeness \\
Severe and complicated withdrawal symptoms (CIWA-Ar ≥ 19) \\
Unstable housing or transportation situation \\
\hline
AWS = alcohol withdrawal syndrome; CIWA-Ar = Clinical Institute Withdrawal Assessment for Alcohol Scale, Revised. \\
\textit{Information from reference 8.} \\
\end{tabular}
\end{table}
### Oral Medications Used to Treat Mild to Moderate AWS

<table>
<thead>
<tr>
<th>Medications</th>
<th>Typical dosing</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Nonbenzodiazepine anticonvulsants</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Carbamazepine (Tegretol)</td>
<td>600 mg to 800 mg</td>
<td>Appropriate monotherapy in mild AWS</td>
</tr>
<tr>
<td>Gabapentin (Neurontin)</td>
<td>Loading dose: 1,200 mg Days 1 through 3: 600 mg to 1,200 mg per day Days 4 through 7: taper to 300 mg to 600 mg per day</td>
<td>Adjunctive therapy dosing: 300 mg to 500 mg every 6 to 8 hours Consider in those with continuing treatment for AUD (1,200 mg per day)</td>
</tr>
<tr>
<td><strong>Benzodiazepines</strong></td>
<td></td>
<td>First-line treatment for moderate AWS. Longer-acting types are preferred; if concern for liver disease, use benzodiazepines with less hepatic metabolism</td>
</tr>
<tr>
<td>Chlordiazepoxide (Librium)</td>
<td>50 mg to 100 mg</td>
<td>Single dose of 50 mg to 100 mg or symptom-triggered dose every 4 to 6 hours</td>
</tr>
<tr>
<td>Diazepam (Valium)</td>
<td>10 mg to 20 mg</td>
<td>10 mg to 20 mg every 6 to 12 hours for the first 24 hours, then reduce to 5 mg to 10 mg every 6 to 12 hours for the next 3 to 5 days Alternative front-loading regimen of 20 mg every 1 to 2 hours for 3 doses, then proceed to symptom-triggered regimen</td>
</tr>
<tr>
<td>Lorazepam (Ativan)</td>
<td>0.5 mg to 2 mg</td>
<td>0.5 mg to 1 mg every 6 to 8 hours on a scheduled basis, plus 1 mg every 4 hours if needed for mild symptoms or plus 2 mg every 2 hours if needed for moderate symptoms</td>
</tr>
<tr>
<td>Oxazepam (Serax)</td>
<td>15 mg to 30 mg</td>
<td>15 mg to 30 mg every 6 to 8 hours</td>
</tr>
<tr>
<td>Phenobarbital</td>
<td>60 mg to 260 mg*</td>
<td>Narrow therapeutic window, should be used by physicians with extensive experience or in Level 2 Withdrawal Management facility</td>
</tr>
<tr>
<td><strong>Adjunctive therapy with benzodiazepines</strong></td>
<td></td>
<td>Used if symptoms persist despite adequate benzodiazepine use</td>
</tr>
<tr>
<td>Beta blockers</td>
<td>Atenolol: 25 mg to 50 mg daily Metoprolol: 25 mg to 50 mg every 12 hours</td>
<td>For persistent hypertension and tachycardia</td>
</tr>
<tr>
<td>Carbamazepine</td>
<td>200 mg every 8 hours or 400 mg every 12 hours</td>
<td>For additional control; reduces craving</td>
</tr>
<tr>
<td>Clonidine</td>
<td>0.2 mg</td>
<td>For autonomic hyperactivity or anxiety</td>
</tr>
<tr>
<td>Gabapentin</td>
<td>400 mg every 6 to 8 hours</td>
<td>For additional control; reduces craving</td>
</tr>
<tr>
<td>Valproate (Depacon)</td>
<td>300 mg to 500 mg every 6 hours</td>
<td>Contraindicated in pregnancy and in patients with liver disease; should not be used as monotherapy for withdrawal</td>
</tr>
</tbody>
</table>

AUD = alcohol use disorder; AWS = alcohol withdrawal syndrome.
*—Can also be given intramuscularly.

Information from references 8 and 14.
ALCOHOL WITHDRAWAL SYNDROME

rates of abstinence than those who received placebo. These results were more pronounced in patients with higher self-reported scores on the Alcohol Withdrawal Severity Checklist than those with lower scores. Antidepressants may be beneficial for concomitant mood disorders but are not used for the treatment of AUD. Community-based support and formal therapeutic interventions should accompany pharmacotherapy. Motivational interviewing and cognitive behavior therapy are beneficial. A recent Cochrane review found Alcoholics Anonymous and 12-step facilitation programs that follow a specific manual or syllabus to be more effective at increasing rates of abstinence than one- or two-step programs that follow up than motivational interviewing or cognitive behavior therapy, with substantial cost savings. Regular monthly follow-up visits for at least one year can increase abstinence.

This article updates previous articles on this topic by Muncie, et al.; Blondell; and Bayard, et al.

Data Sources: A PubMed search was completed using the key terms alcohol use disorder (AUD), alcohol withdrawal, alcohol dependence, epidemiology, diagnosis, delirium tremens, screening, outpatient, and management. The search included meta-analyses, systematic reviews, practice guidelines, clinical trials, and original studies. Also searched were the Cochrane database, ECRI Guidelines Trust, Essential Evidence Plus, and the U.S. Preventive Services Task Force. Search dates: March 4 and 14, April 15 and 29, and June 15, 2020.

The opinions and assertions contained herein are the private views of the authors and are not to be construed as official or as reflecting the views of the Uniformed Services University of the Health Sciences, the U.S. Army Medical Department, or the U.S. Army at large.

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<table>
<thead>
<tr>
<th>Clinical recommendation</th>
<th>Evidence rating</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adults should be screened for unhealthy alcohol use and counseled, if appropriate.</td>
<td>B</td>
<td>U.S. Preventive Services Task Force recommendation</td>
</tr>
<tr>
<td>Patients with mild or moderate AWS can be treated safely in the outpatient setting.</td>
<td>C</td>
<td>ASAM guideline</td>
</tr>
<tr>
<td>Carbamazepine (Tegretol) and gabapentin (Neurontin) are appropriate options for treating mild AWS.</td>
<td>C</td>
<td>ASAM guideline recommendation, randomized controlled trial, and review</td>
</tr>
<tr>
<td>Benzodiazepines are the preferred medication for treating moderate AWS.</td>
<td>A</td>
<td>Clinical practice guideline and ASAM meta-analysis</td>
</tr>
<tr>
<td>Gabapentin, carbamazepine, and valproate (Depacon) may be prescribed as adjuncts to benzodiazepines if symptoms persist despite adequate benzodiazepine use.</td>
<td>C</td>
<td>Expert opinion</td>
</tr>
<tr>
<td>Patients successfully treated for AWS should be offered referral to a long-term treatment program to maintain abstinence and adjunctive use of an FDA-approved pharmacotherapy (i.e., acamprosate, naltrexone [Revia], and disulfiram [Antabuse]), meta-analysis of FDA-approved pharmacotherapies and Cochrane review</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

ASAM = American Society of Addiction Medicine; AWS = alcohol withdrawal syndrome; FDA = U.S. Food and Drug Administration.

A = consistent, good-quality patient-oriented evidence; B = inconsistent or limited-quality patient-oriented evidence; C = consensus, disease-oriented evidence, usual practice, expert opinion, or case series. For information about the SORT evidence rating system, go to https://www.aafp.org/afpsort.
ALCOHOL WITHDRAWAL SYNDROME

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References

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