



# Low-threshold, Complex Outpatient Withdrawal Management in Resource Limited Communities

**Sarah Spencer DO, FASAM Ninilchik Tribal Community Clinic**

**Workshop Session, ASAM 2024 Annual Conference, Dallas, TX April 2024**



# Financial Disclosures



I have no financial conflicts of interest to disclose

I am currently employed by the Ninilchik Tribal Council

I work as a treatment consultant for the Opioid Response Network, the Alaska Native Tribal Health Consortium, as well as for other non-profit agencies.

# Learning Objectives

- ☀ Explore challenges in providing stabilizing care to individuals with complicated use disorders in communities with limited resources.
- ☀ Work as a team to problem-solve cases for patients from vulnerable populations who are using multiple substances, whose goals may not be abstinence-based
- ☀ Compile information from evidenced-based resources as well as clinical experience and local knowledge to create patient-centered, harm-reduction focused care plans.

# Off Label Disclosure

**This workshop discusses many off label medication uses.**

**Not all patients fit neatly in a box;**

**That's why medicine is an art.**

We will discuss utilizing evidence-based practices along with local resources to address complex cases when “standard of care” treatment is not accessible. Some off label uses discussed include:

- ☀ Managed alcohol use and outpatient alcohol withdrawal management
- ☀ Rapid outpatient benzodiazepine tapers
- ☀ Rapid initiation of long-acting injectable buprenorphine
- ☀ Alternative buprenorphine dosing strategies

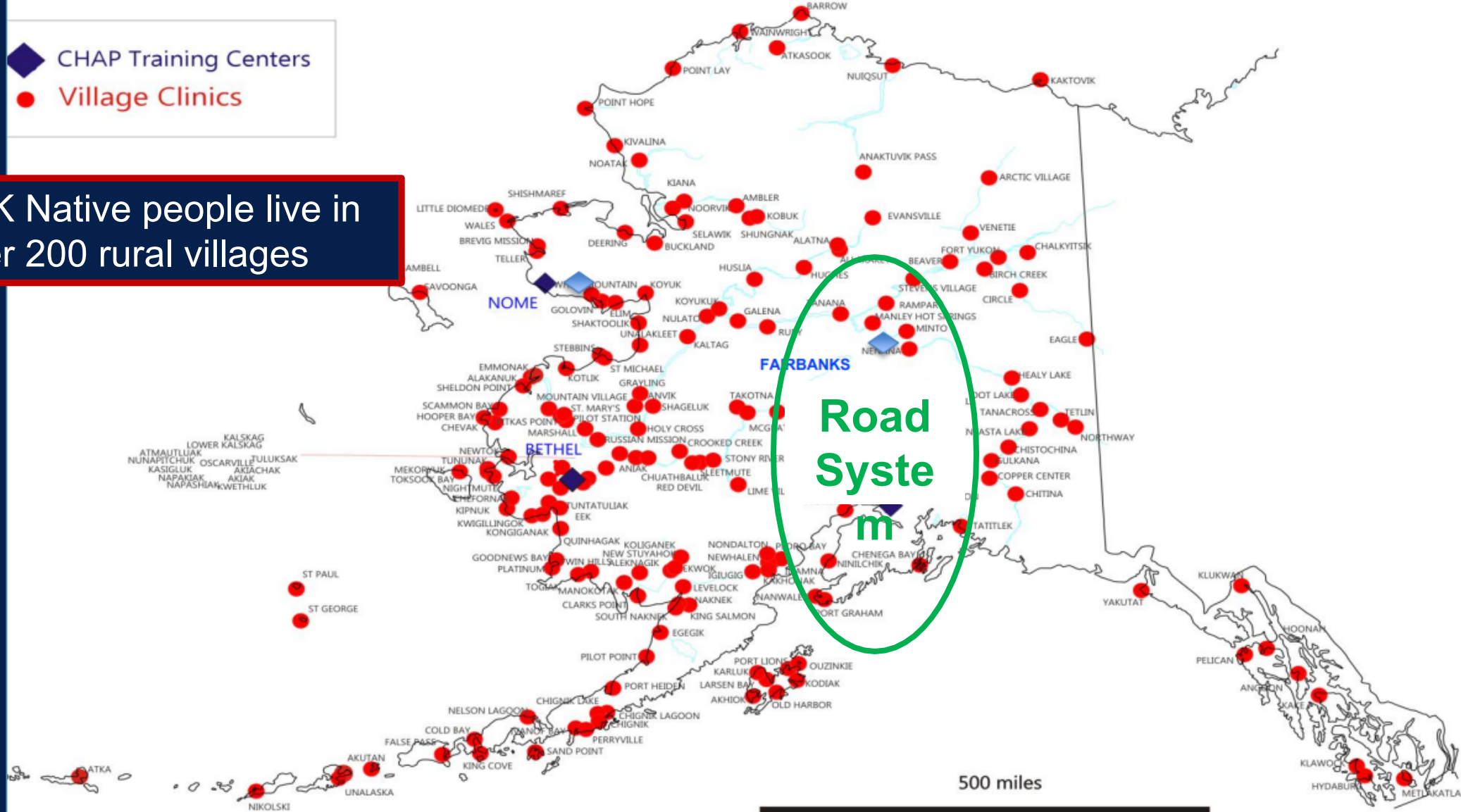
# Access to Care in Rural Alaska



# Community Health Aide/Practitioner Village Clinics

- ◆ CHAP Training Centers
- Village Clinics

90K AK Native people live in over 200 rural villages



# Barriers to Treatment Access in Rural AK



Travel costs (over \$1,000 per trip to ANC) and time

No local pharmacies, weather holds/ Rx delayed in the mail

No local licensed medical/BH providers (only CHAPs/BHAs)

No local OTP or inpatient withdrawal management

Lack of anonymity, STIGMA

# Community Resources in Rural Alaska





# Community Health Aides and Community Health Practitioners (CHA/Ps)



- Local people
- Initially described as “the eyes, ears and hands of the physician”
- Approximately 180 villages utilize CHA/Ps
- Conduct more than ½ of all yearly patient encounters in the State of Alaska
- Includes emergency, acute, chronic, and preventive health components
- Training is “skills” based
- Does not include differential diagnosis
- Under medical supervision of a licensed physician

# Community Health Aides and Community Health Practitioners (CHA/Ps)

Clinical skills include:

- Taking a history
- Performing a physical exam
- Performing lab skills
- Use of the Community Health Aide Manual (CHAM) to make Assessments
- Report
- Following plans per the CHAM
- Giving patient education
- Administering medicines
- Performing certain treatment procedures
- Documenting patient encounters



**CHAPs and BHAs  
can do home  
visits**

# Growing Our Own:



**A Grassroots Approach to Increasing Behavioral Health Access for American Indian and Alaska Native People**



<https://www.anthc.org/behavioral-health-aide-program/>



*BHAs are an integral part of our substance abuse programs. They assist in facilitating sessions for our Intensive Outpatient Substance Abuse program that is delivered via televideo to the villages and, recently, they volunteered to fill-in at our recovery camp program, which is currently experiencing a staffing shortage.*

Substance abuse assessment &  
treatment



ALASKA NATIVE  
TRIBAL HEALTH  
CONSORTIUM

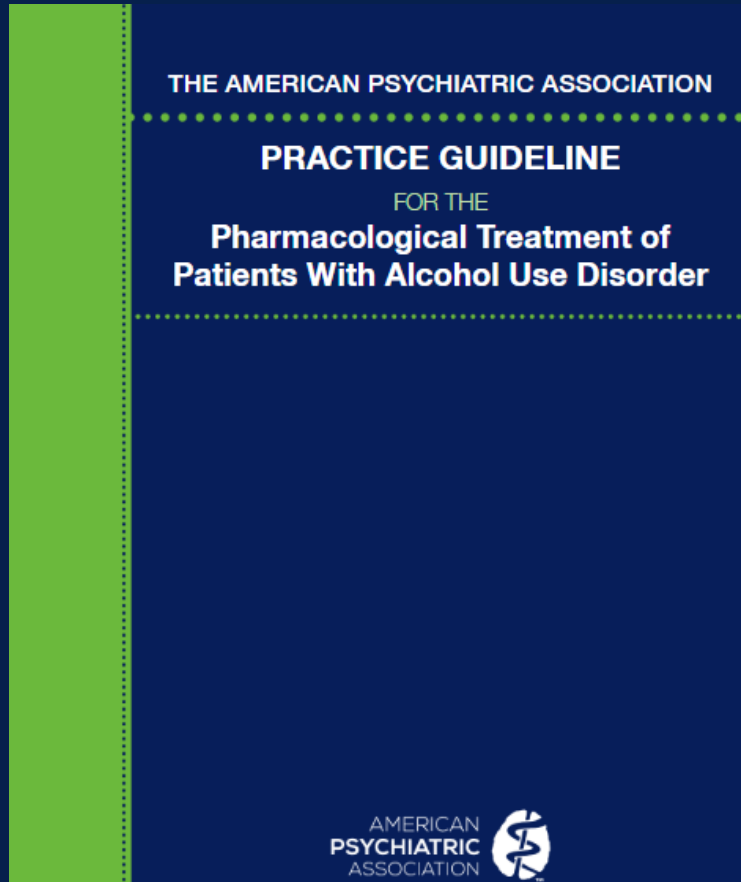
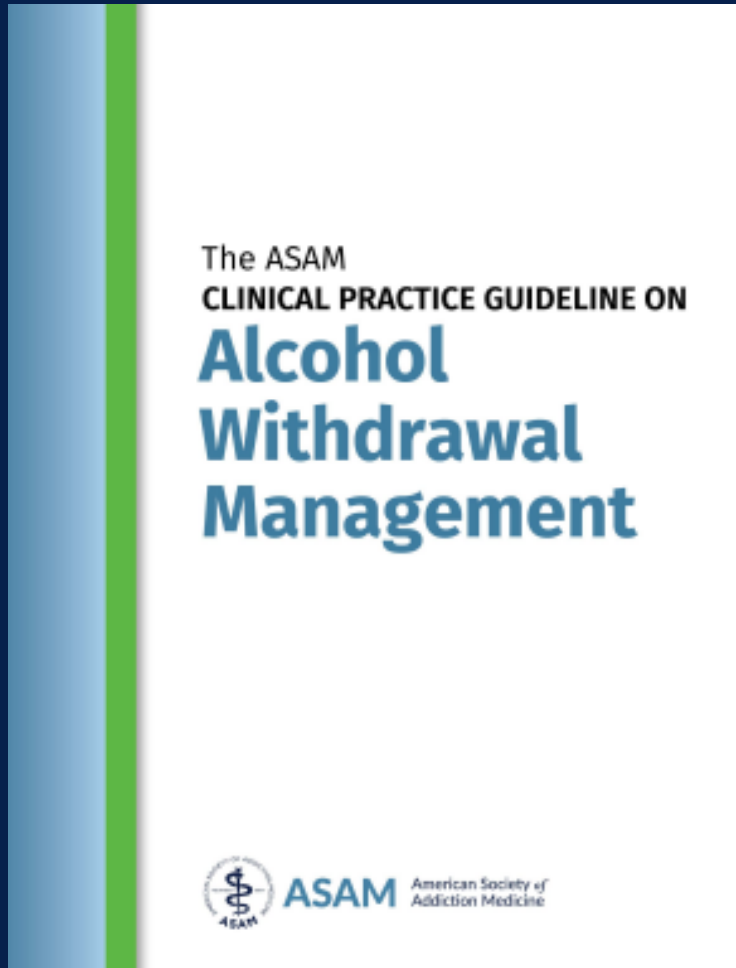
# Peer Support/Behavioral Health Aide Roles

- Available by cell phone (patients can call/text)
- Connect with patients at medical appointments to check in with them
- Support groups (Virtual)
- Recreational opportunities (crafting/drumming groups, beach cleanups, fish camp)
- Connect patients to resources (doing the research, providing and helping fill out paperwork-housing, employment, treatment)
- Providing transportation (to medical/behavioral health appointments, mutual support meetings, court, residential treatment)
- Volunteer and connect with participants at our local syringe access program
- Warm handoff from ED/Hospital or jail (check court records)  
• Sit down for a chat of a cup of coffee
- Crisis Intervention

# Outpatient Alcohol Withdrawal Management



# NATIONAL PRACTICE GUIDELINES



# WHO CAN BE MANAGED IN THE CLINIC



Support system



Only mild or moderate symptoms



Frequent check-ins



No history of severe withdrawal



No significant comorbidities



**Figure 3. Prediction of Alcohol Withdrawal Severity Scale (PAWSS)**

**PART A: THRESHOLD CRITERIA:**

(“Y” or “N”,  
no point)

Have you consumed any amount of alcohol (i.e., been drinking) within the last 30 days?  
OR did the patient have a “+” blood alcohol level (BAL) on admission?

*IF the answer to either is YES, proceed with test:*

**PART B: BASED ON PATIENT INTERVIEW:**

(1 point each)

1. Have you been recently intoxicated/drunk within the last 30 days?
2. Have you ever undergone alcohol use disorder rehabilitation treatment or treatment for alcoholism? (i.e., inpatient or outpatient treatment programs or AA attendance)
3. Have you ever experienced any previous episodes of alcohol withdrawal, regardless of severity?
4. Have you ever experienced blackouts?
5. Have you ever experienced alcohol withdrawal seizures?
6. Have you ever experienced delirium tremens, or DT?
7. Have you combined alcohol with other “downers” like benzodiazepines or barbiturates during the last 90 days?
8. Have you combined alcohol with any other substance of abuse during the last 90 days?

**PART C: BASED ON CLINICAL EVIDENCE:**

(1 point each)

9. Was the patient’s BAL on presentation  $\geq$  200?
10. Is there evidence of increased autonomic activity? (e.g., HR > 120 bpm, tremor, sweating, agitation, nausea)

**TOTAL SCORE:** \_\_\_\_\_

*Notes: Maximum score = 10. This instrument is intended as a SCREENING TOOL. The greater the number of positive findings, the higher the risk for the development of AWS. A score of  $\geq$  4 suggests HIGH RISK for moderate to severe (complicated) AWS; prophylaxis and/or treatment may be indicated.*

**Source:** Adapted from Maldonado JR, Sher Y, Ashouri JF, et al. The “prediction of alcohol withdrawal severity scale” (PAWSS): systematic literature review and pilot study of a new scale for the prediction of complicated alcohol withdrawal syndrome. *Alcohol*. 2014;48(4):375-390.

**PAWSS score >4  
= high risk for  
complicated  
AWS**



## Harm Reduction Tip:

Some patients may decline to go to the hospital for AWS management, and for those certain individuals, it may be safer to trial supervised ambulatory withdrawal than receive no medical care at all.



## TRIAGING PEOPLE APPROPRIATE FOR AMBULATORY MANAGEMENT OF ALCOHOL WITHDRAWAL SYNDROME (AWS)



Evaluate with history/physical  
(Labs not necessary)



Currently in moderate or severe alcohol withdrawal?

No

Yes



Recent history of severe alcohol withdrawal (seizure or delirium tremens)?

No

Yes



Relative contraindications:  
comorbidities (4Cs<sup>^</sup>), pregnant,  
unstable psychiatric disease, or  
psychosocial complexity?

No

Yes

Patient-centered discussion about hospital-based treatment of AWS\*

Ambulatory AWS treatment

THE CURB SIDERS  
INTERNAL MEDICINE

\*While in these situations hospital withdrawal management is recommended, particularly if recent severe withdrawal, ambulatory management is still safer than no management in a patient who declines hospital evaluation

<sup>^</sup>CHF/heart failure NYHA Class 2+, decompensated cirrhosis, CKD Stage 3+, or COPD on O2.

# **AMBULATORY ALCOHOL WITHDRAWAL TREATMENT**

## **Best Practices**

**1**

**Start in the morning and don't taper alcohol before**

**2**

**Check in every other day, can use telehealth (video preferred) to make more accessible**

**3**

**Recommend hospitalization if:  
seizures, altered mental status, using more PRNs than  
prescribed**

**4**

**Don't forget to address and treat AUD**

# Short Alcohol Withdrawal Scale

Item	None (0 points)	Mild (1 point)	Moderate (2 points)	Severe (3 points)
Anxious				
Feeling confused				
Restless				
Miserable				
Problems with memory				
Tremor (shakes)				
Nausea				
Heart pounding				
Sleep disturbance				
Sweating				

**Short Alcohol Withdrawal Scale to assess severity of alcohol withdrawal.**

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

*Adapted with permission from Elholm B, Larsen K, Hornnes N, et al. A psychometric validation of the Short Alcohol Withdrawal Scale (SAWS). Alcohol Alcohol. 2010;45(4):362.*



# AMBULATORY ALCOHOL WITHDRAWAL REGIMENS

	<b>Diazepam based<sup>^</sup></b>	<b>Gabapentin based</b>
<b>Day 1</b>	10mg q6hrs*	300mg q6hrs*
<b>Day 2</b>	10mg TID	300mg TID
<b>Day 3</b>	10mg BID	300mg BID
<b>Day 4</b>	10mg once	300mg once
<b>Additional PRNs</b>	5 x 10mg pills	5 x 300mg pills

<sup>^</sup>Can substitute chlordiazepoxide 50mg for diazepam 10mg

\*If >10 drinks per day double dose on first day (Dr Holt Expert opinion)

## Oral Medications Used to Treat Mild to Moderate AWS

Medications	Typical dosing	Comments
<b>Nonbenzodiazepine anticonvulsants</b>		Appropriate monotherapy in mild AWS
Carbamazepine (Tegretol)	600 mg to 800 mg	600 mg to 800 mg per day tapered to 200 mg to 400 mg per day over 4 to 9 days

<b>Adjunctive therapy with benzodiazepines</b>		Used if symptoms persist despite adequate benzodiazepine use
Beta blockers	Atenolol: 25 mg to 50 mg daily Metoprolol: 25 mg to 50 mg every 12 hours	For persistent hypertension and tachycardia
Carbamazepine	200 mg every 8 hours or 400 mg every 12 hours	For additional control; reduces craving
Clonidine	0.2 mg	For autonomic hyperactivity or anxiety
Gabapentin	400 mg every 6 to 8 hours	For additional control; reduces craving
Valproate (Depacon)	300 mg to 500 mg every 6 hours	Contraindicated in pregnancy and in patients with liver disease; should not be used as monotherapy for withdrawal

# Outpatient Phenobarbital

*In a Level 2-WM ambulatory setting, phenobarbital monotherapy, managed by a clinician experienced with its use, is appropriate for patients with a contraindication for benzodiazepine use who are experiencing moderate or severe alcohol withdrawal or who are at risk of developing severe or complicated alcohol withdrawal or complication of alcohol withdrawal. Discussion. There is disagreement in the literature regarding the appropriateness of phenobarbital in ambulatory settings, due to the risk of toxicity when used in combination with alcohol or in high doses. In general, phenobarbital should only be used by clinicians experienced with its use in settings that offer close monitoring. Phenobarbital may cause respiratory depression and over-sedation and its narrow therapeutic window makes it challenging to dose correctly compared to other medications used to treat alcohol withdrawal. As with benzodiazepines, effects on the central nervous system are exacerbated when other CNS depressants such as alcohol are also used.*

# Phenobarbital Options for Withdrawal

## Mild

260 mg IV push or 100 mg PO



130 – 260 mg IV push or 100 mg PO  
x 2 prn q60 min



Discharge

## Moderate-Severe

10 mg/kg IV over 30 minutes



130 – 260 mg IV push or  
100 – 200 mg PO  
x 2 prn q60 min



3 day benzodiazepine taper of diazepam  
or chlordiazepoxide

Mild: Oral Taper 60mg qid day1, 60 mg tid day 2,  
60 mg bid day 3, 30 mg bid day 4



## Managed Alcohol Program (MAP) Outcomes

- ☀️ Improvement in quality of life among MAP participants who were **less likely to report acute alcohol-related harms** such as seizures, acute intoxication, trauma, or assault
- ☀️ Participants noted a positive change in their relationship with alcohol, from a decreased focus on alcohol procurement to an increased feeling of control regarding consumption levels...**increased sense of self-determination and motivation for positive change**
- ☀️ **Reported a reduction in feelings of shame and guilt**
- ☀️ MAP participants had greater number of drinking days, with one study finding an average of 27.8 alcohol days per month for MAP participants compared to 22.6 alcohol days for controls. Yet, this drinking pattern resulted **both in a reduction in overall quantity of alcohol and in a less hazardous consumption**

## Create At-Home Alcohol Withdrawal Monitoring and Management Plan

- Managed alcohol intake, dosed by caregiver
- Homer breathalyzer?
- Train caregivers to calculate SOWS
- On-hand meds to treat withdrawal (including alternative to PO route)

# Outpatient Benzodiazepine Withdrawal Management



# Outpatient Benzo Taper Selection

- ☀️ Reliable and motivated to stop
- ☀️ Medical/MH stable
- ☀️ Social support/transport to ED
- ☀️ No/mild use disorder
- ☀️ No h/o seizures
- ☀️ Not pregnant

**Generally,  
Taper  
Slowly!**

## Benzodiazepine Equivalency Chart

Drug	Half-life (hrs)	Dose Equivalent
Chlordiazepoxide (Librium)	5–30 h	25mg
Diazepam (Valium)	20–50 h	10mg
Alprazolam (Xanax)	6–20 h	0.5mg
Clonazepam (Klonopin)	18–39 h	0.5mg
Lorazepam (Ativan)	10–20 h	1mg
Oxazepam (Serax)	3–21 h	15mg
Triazolam (Halcion)	1.6–5.5 h	0.5mg
Phenobarbital (barbituate)	53 – 118 h	30 mg

# Rapid Benzodiazepine Taper

## RAPID TAPER

- 1** Pre-medicate two weeks prior to taper with valproate 500mg BID or carbamazepine 200mg every AM and 400mg every HS. Continue this medication for four weeks post-benzodiazepines. Follow the usual safeguards (lab testing and blood levels) when prescribing these medications.
- 2** Utilize concomitant behavioral supports.
- 3** Discontinue current benzodiazepine treatment and switch to diazepam 2mg BID for two days, followed by 2mg every day for two days, then stop. For high doses, begin with 5mg BID for two days and then continue as described.



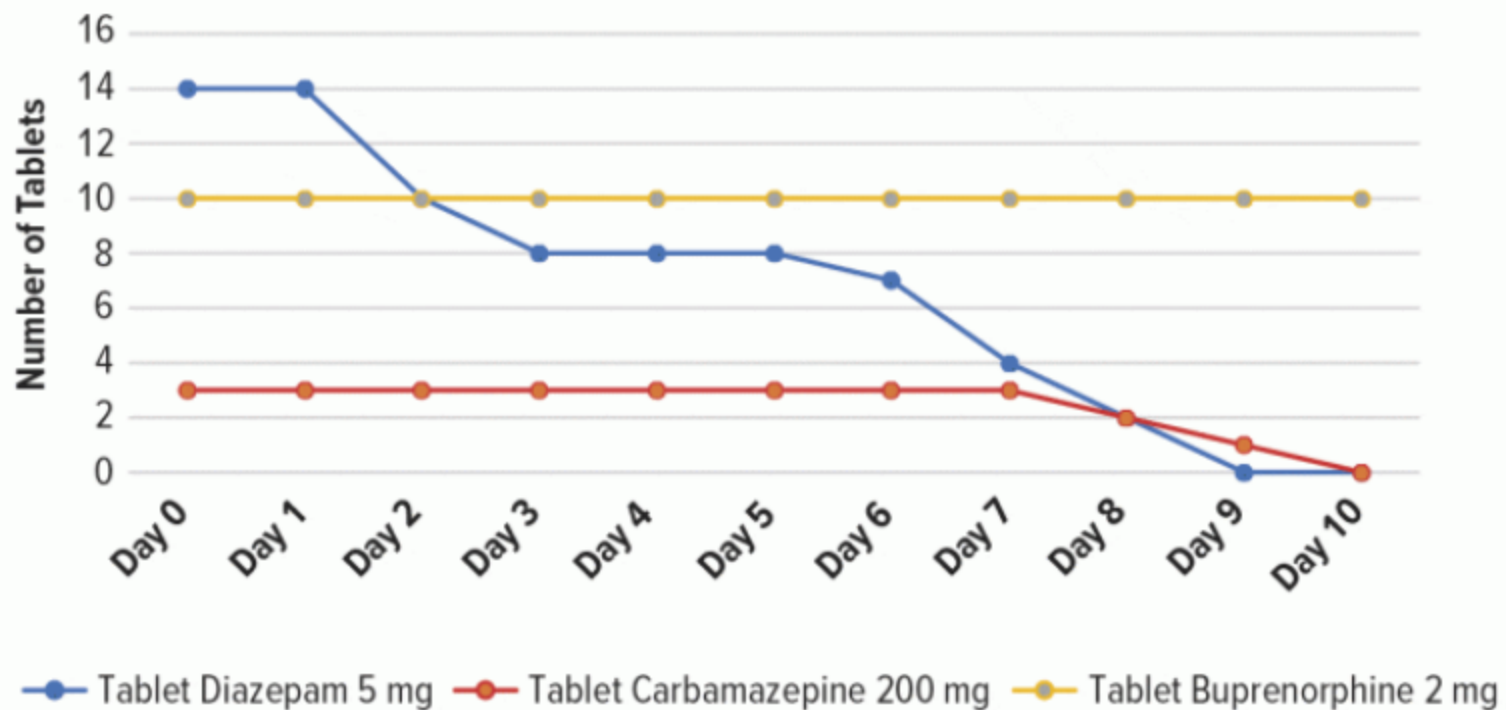
<https://www.oregonpainguidance.org/app/content/uploads/2016/05/Opioid-and-Benzodiazepine-Tapering-flow-sheets.pdf>

# Rapid Detoxification in an Adult With Benzodiazepine Dependence With the Aid of an Antiepileptic

Diveesha Munipati, MBBS; Rahul Mathur, MD, DNB; and Siddharth Sarkar, MD, DNB

**Published:** August 17, 2023

## Graphical Representation of Rapid-Tapering Schedule of Diazepam<sup>a</sup>



<sup>a</sup>Prior to tapering, an antiepileptic was added. Dose of opioid agonist (buprenorphine) was unchanged throughout the schedule.

# Adjunctive Medications

Medication	Effect of Medication	Study
Hydroxyzine	Patients taking 25-50 mg had a decrease in anxiety during a benzodiazepine taper compared to placebo.	Lemoine et al., 1997
Carbamazepine	When given 200-800 mg/day during and after a benzodiazepine taper, it reduced withdrawal symptoms and promoted abstinence compared to placebo.	Schweizer et al., 1991
Trazodone	A significantly higher percentage of patients taking trazodone during a benzodiazepine taper were abstinent from benzodiazepines at 5 weeks post-taper compared to patients taking placebo, but there was no difference at 12 weeks post-taper.	Rickels et al., 1999
Sodium valproate	A significantly higher percentage of patients taking sodium valproate during a benzodiazepine taper were abstinent from benzodiazepines at 5 weeks post-taper compared to patients taking placebo, but there was no difference at 12 weeks post-taper.	Rickels et al., 1999
Imipramine	Pretreatment and use of imipramine during benzodiazepine taper increased taper success rate; a significantly higher percentage of patients taking imipramine were abstinent from benzodiazepines at 12 weeks post-taper compared to those taking placebo.	Rickels et al., 2000

# Adjunctive Medications

Medication	Effect of Medication	Study
Pregabalin	Patients treated with pregabalin (150-600 mg/day) had significantly lower withdrawal symptoms compared to placebo, both during taper and 6 weeks after. Group treated with pregabalin had lower anxiety during taper.	Hadley et al. (2012)
Buspirone	Subjects given buspirone during BZD withdrawal had lower levels of anxiety than subjects given placebo.	Morton & Lader (1995) Udelman & Udelman (1990)
Gabapentin	In MMT patients taking doses up to 1200 mg TID, there were no significant differences between gabapentin and placebo on amount of BZD use per day (both groups reduced use), days abstinent per week, and CIWA-B scale.	Mariani et al. (2016)
Flumazenil	Randomized, placebo-controlled study found subjects given flumazenil infusion plus oxazepam significantly reduced withdrawal symptoms and cravings compared to oxazepam and placebo. Subjects given flumazenil infusion had lower relapse rates up to 30 days later.	Gerra et al. (2002)
Melatonin	Cross-over study, compared melatonin to placebo in MMT patients using BZD. Sleep quality improved with cessation of BZD, regardless of group. In each group, ~30% stopped using BZD. <a href="https://www.aoaam.org/resources/Documents/2018%20Convention%20Slides/Saturday%20-%2010-6-2018%20-%2011am%20-%20Management%20of%20Withdrawal%20-%20Alcohol%20Benzodiazepines%20Opioids%20-%20Kmicc.pdf">https://www.aoaam.org/resources/Documents/2018%20Convention%20Slides/Saturday%20-%2010-6-2018%20-%2011am%20-%20Management%20of%20Withdrawal%20-%20Alcohol%20Benzodiazepines%20Opioids%20-%20Kmicc.pdf</a>	Peles et al. (2007)





® Rule

## Schedules of Controlled Substances: Temporary Placement of Etizolam, Flualprazolam, Clonazolam, Flubromazolam, and Diclazepam in Schedule I

A Rule by the Drug Enforcement Administration on 07/26/2023

Paper that list some equivalency tables for Designer Benzos

<https://academic.oup.com/jat/article/47/1/1/6549532>

June 2022

### Bromazolam Prevalence Surging Across the United States Driven In Part by Increasing Detections Alongside Fentanyl



[Pharmaceuticals \(Basel\)](#). 2021 Jun; 14(6): 560.

PMCID: PMC8230725

Published online 2021 Jun 11. doi: [10.3390/ph14060560](https://doi.org/10.3390/ph14060560)

PMID: [34208284](https://pubmed.ncbi.nlm.nih.gov/34208284/)

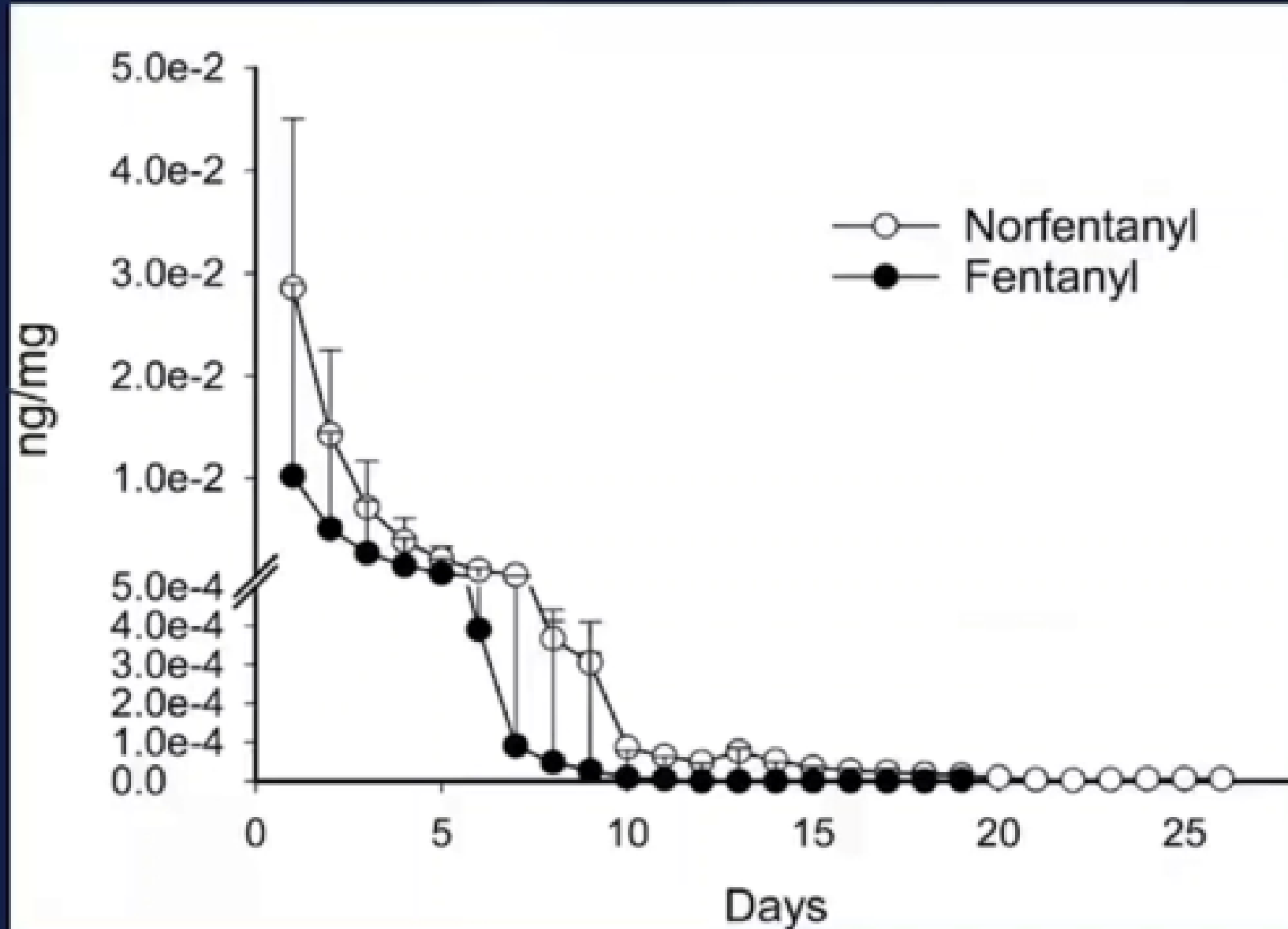
### Designer Benzodiazepines: A Review of Toxicology and Public Health Risks

[Pietro Brunetti](#),<sup>1</sup> [Raffaele Giorgetti](#),<sup>1</sup> [Adriano Tagliabracci](#),<sup>1</sup> [Marilyn A. Huestis](#),<sup>2,\*</sup> and [Francesco Paolo Busardò](#)<sup>1</sup>

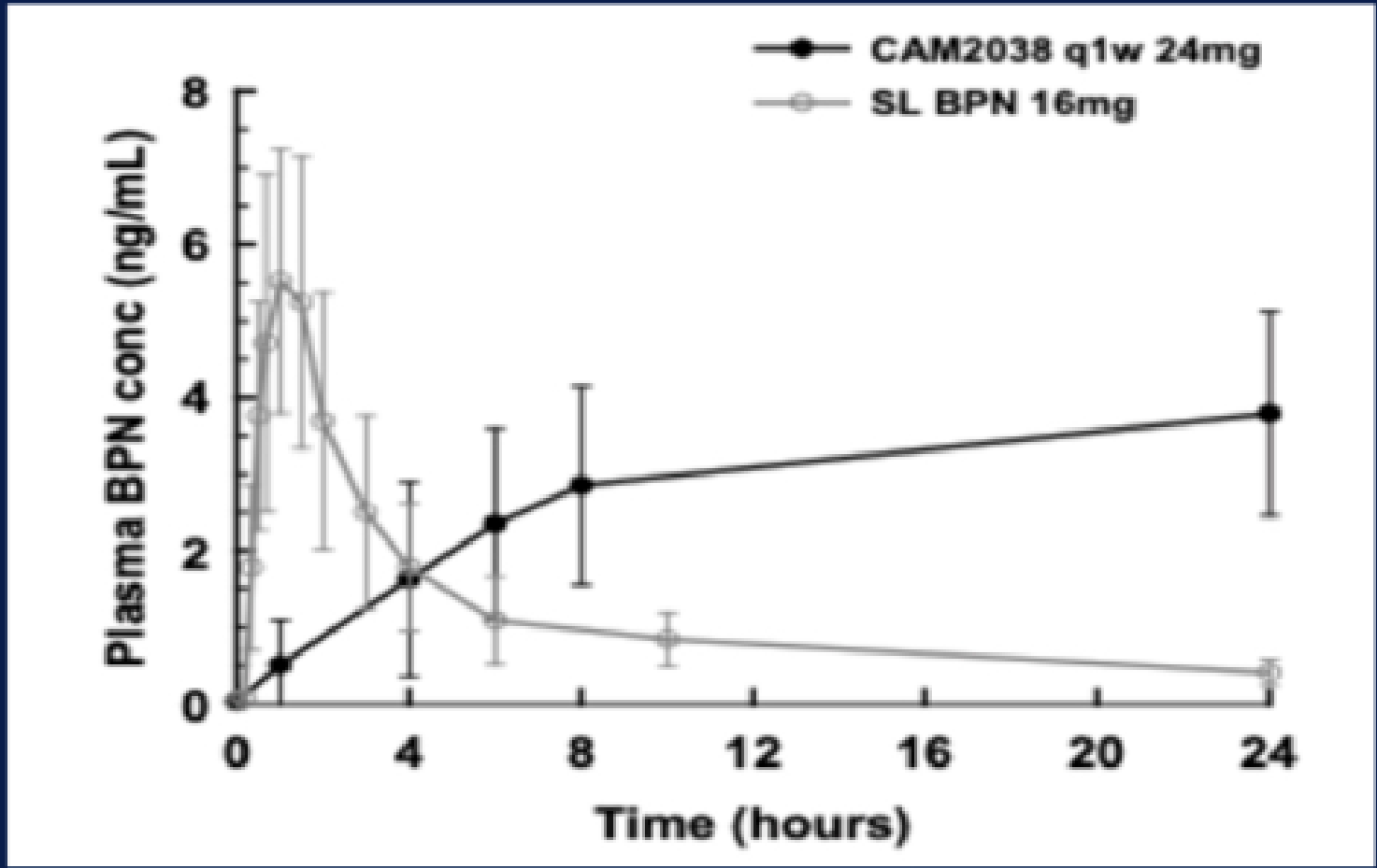
# Outpatient Opioid Withdrawal Management for Patients Initiating Buprenorphine



# FENTANYL NORFENTANYL ELIMINATION IN URINE



# ED-Initiated Buprenorphine VALIDATION Network Trial ED INNOVATION



**XR BUP given in  
ED  
1200 patients  
9 cases of PW  
(0.76%)**

Pharmacokinetics of XR- & SL- Buprenorphine

<https://jamanetwork.com/journals/jamanetworkopen/fullarticle/2802898>

# High dose buprenorphine initiation

California ED Bridge Program  
<https://cabridge.org/tools/resources/>



## Buprenorphine Self-Start

Guidance for patients starting buprenorphine outside of hospitals or clinics

- 1 Plan to take a day off and have a place to rest.
- 2 Stop using and wait until you feel very sick from withdrawals (at least 12 hours is best, if using fentanyl it may take a few days).
- 3 Dose one or two 8mg tablets or strips UNDER your tongue (total dose of 8-16mg).
- 4 Repeat dose (another 8mg-16mg) in an hour to feel well.
- 5 The next day, take 16-32mg (2-4 tablets or films) at one time.

### If you have started bup before:

- If it went well, that's great! Just do that again.
- If it was difficult, talk with your care team to figure out what happened and find ways to make it better this time. You may need a different dosing plan than what is listed here.

### If you have never started bup before:

- Gather your support team and if possible take a "day off."
- You are going to want space to rest. Don't drive.
- Using cocaine, meth, alcohol or pills makes starting bup harder, and mixing in alcohol or benzos can be dangerous.



Place dose under your tongue (sublingual).

### If you have a light habit: (For example, 5 "Norco 10's" a day)

- Consider a low dose: start with 4mg and stop at 8mg total.
- **WARNING:** Withdrawal will continue if you don't take enough bup.

### If you have a heavy habit: (For example, injecting 2g heroin a day or smoking 1g fentanyl a day)

- Consider a high dose: start with a first dose of 16mg.
- For most people, the effects of bup max out at around 24-32mg.
- **WARNING:** Too much bup can make you feel sick and sleepy.

# Low-dose Overlapping SLBUP Start

*How to*

	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7
<b>Buprenorphine dose</b>	0.5mg daily	0.5mg BID	1mg BID	2mg BID	4mg BID	4mg TID	8mg BID
<b>Film size</b>	2mg	2mg	2mg	2mg	2mg	2mg	8mg
<b>Morning dose</b>							
<b>Afternoon Dose</b>							
<b>Night dose</b>							
<b>Full agonist</b>	Continue	Continue	Continue	Continue	Continue	Continue	STOP



# Long-term buprenorphine treatment for kratom use disorder: A case series

Viktoriya R. Broyan , MS, Jessica K. Brar , BS, Tristen Allgaier, Student & Jeffrey T. Allgaier , MD, FACEP, FASAM

Pages 763-766 | Published online: 03 Feb 2022

[Cureus](#). 2023 Jun; 15(6): e41146.

PMCID: PMC10386870

Published online 2023 Jun 29. doi: [10.7759/cureus.41146](https://doi.org/10.7759/cureus.41146)

PMID: [37519540](https://pubmed.ncbi.nlm.nih.gov/37519540/)

## Successful Management of Kratom Use Disorder With Buprenorphine and Naloxone

Monitoring Editor: Alexander Muacevic and John R Adler

[Martin Arhin](#),<sup>✉1</sup> [Julian Mobley](#),<sup>1</sup> [Hamad Hamad](#),<sup>2</sup> and [Paul Remick](#)<sup>2</sup>

> [J Addict Med](#). 2021 Apr 1;15(2):167-172. doi: [10.1097/ADM.0000000000000721](https://doi.org/10.1097/ADM.0000000000000721).

## Treatment of Kratom Withdrawal and Dependence With Buprenorphine/Naloxone: A Case Series and Systematic Literature Review

[Stephanie T Weiss](#) <sup>1</sup>, [Heather E Douglas](#)

# Management of Co-morbid Stimulant Use Disorder





# ASAM and AAAP Announce New Clinical Practice Guideline to Address Rising Stimulant Use Disorders

Nov 7, 2023

Download

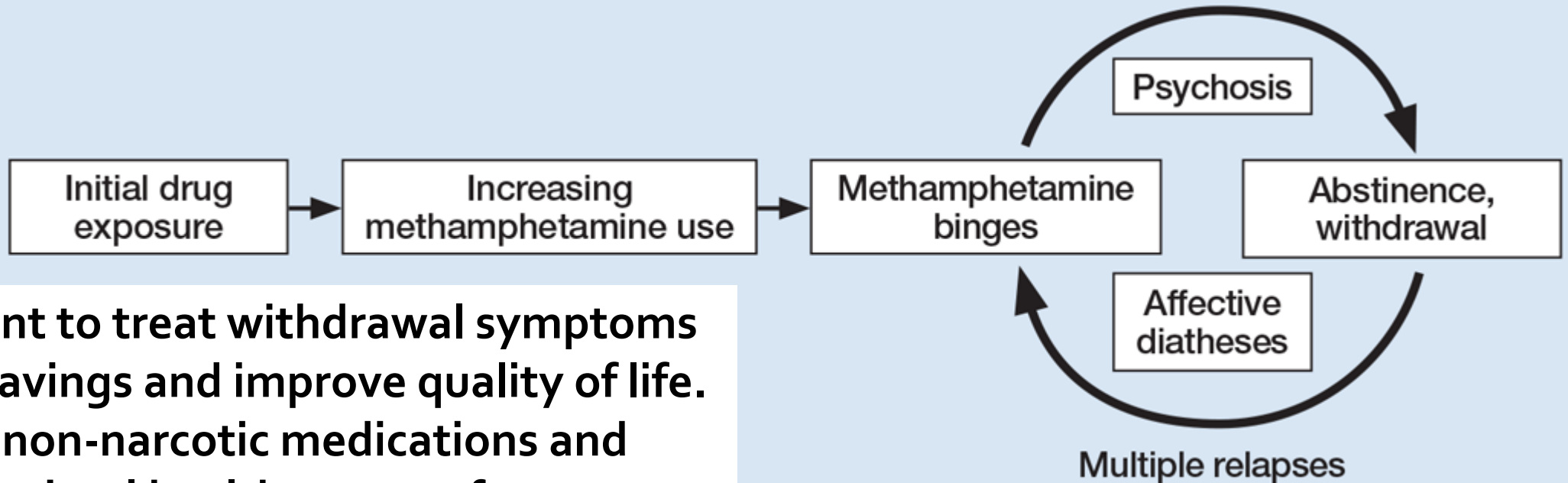
Contingency  
Management is  
#1

*Board certified  
providers may  
consider Rx  
stimulants in select  
patients with close  
monitoring*



[https://downloads.asam.org/sitefinity-production-blobs/docs/default-source/quality-science/stud\\_guideline\\_document\\_final.pdf?sfvrsn=71094b38\\_1](https://downloads.asam.org/sitefinity-production-blobs/docs/default-source/quality-science/stud_guideline_document_final.pdf?sfvrsn=71094b38_1)

# Supporting patients in the cycle of stimulant use



**It's important to treat withdrawal symptoms to reduce cravings and improve quality of life. Provide non-narcotic medications and behavioral health support for:**

- Depression**
- Anxiety**
- Insomnia**
- Pain**
- Thought disorders**

# Pharmacological Management of Psychosis

☀ *“Recent research suggests that **olanzapine or quetiapine may be preferred for the management of methamphetamine-induced psychosis**; however, the evidence is considered low quality... When managing psychosis prior to confirming the etiology of stimulant intoxication or withdrawal, clinicians should conduct an evaluation with a focus on identifying potential causes of the patient’s psychosis other than stimulant intoxication. Clinicians should focus treatment of psychosis on management of the underlying causes of the patient’s psychotic symptoms and monitor for medication side effects with usual care”*

# ASK ABOUT ANXIETY AND DEPRESSION

**Mirtazapine** helps insomnia and anorexia

**Bupropion** helps with energy and appetite suppression

Less evidence for SSRIs

Avoid Benzos → try **buspirone** instead

Consider treating physiological WD sx's that mimic anxiety with  
prn **clonidine, hydroxyzine, propranolol**

# Ask About Sleep

- ☀️ Make a differential diagnosis to determine whether a client's sleep problems likely stem from protracted withdrawal or are the result of other causes.
- ☀️ Educate clients about good sleep habits: adopting a regular sleep routine (going to bed and getting up at the same times), exercising early in the day, minimizing caffeine intake, eating well, and avoiding late afternoon naps.
- ☀️ Utilize non-narcotic meds, consider co-treating comorbid pain/mood disorders: eg **TCA**s, **mirtazapine**, **trazodone**, **doxepin**, **quetiapine** (all off label), low-dose melatonin
- ☀️ Test for sleep apnea (opioids=central vs obstructive)
- ☀️ CBTI (free VA app)



# Harm Reduction



# Harm Reduction Based Low Threshold Care

- Don't discharge patients for ongoing drug use
- Flexible walk-in/same day/tele-med appointments
- Peer support (via text)
- Treatment of co-morbid medical/MH issues
- Contingency Management
- Contraception
- Hep C treatment/ PREP for active users
- Naloxone kits
- Injection and smoking supplies
- Fentanyl/xylazine test strips
- Low threshold MOUD access

Education and Training

Harm Reduction Kit

*Rural Alaska Harm  
Reduction Initiative*

[www.iknowmine.org/harmreduction](http://www.iknowmine.org/harmreduction)

Narcan® (Naloxone) Kit

Safe Medication  
Storage and Disposal

# ANTHC Harm Reduction Training and Supplies

<https://www.iknowmine.org/topic/harm-reduction/>

[https://www.iknowmine.org/wp-content/uploads/2021/01/ANTHC\\_Harm-Reduction\\_Toolkit.pdf](https://www.iknowmine.org/wp-content/uploads/2021/01/ANTHC_Harm-Reduction_Toolkit.pdf)



## WHAT IS NEXT Distro?

An online and mail-based **harm reduction service** designed to reduce opioid overdose death, prevent injection-related disease transmission, and improve the lives of people who use drugs. <https://nextdistro.org>



# An example of a prescription for syringes

*Diabetic syringes*

***29g, 1/2in “longs” or 31g, 5/16in “shorts”***

*(ask patient which they prefer)*

***1/2 or 1 cc***

*(ask patient which they prefer, 1/2 cc is more common)*

*Dispense #\_\_ boxes of 100 syringes*

*Refill PRN X 1year*

**Allows for online ordering**

# Patient Cases



Groups will work together to apply their clinical knowledge and unique experience to develop a proposed treatment plan for their group's case.

# Community Resources

- ☀ In a roadless village: Community health aid, Behavioral health aid, village clinic with emergency medication, Medivac required for hospital care, Rx get mailed to patient or clinic.
- ☀ On road system: Peer supports, Primary care clinic, critical access hospital, pharmacy (all may require transportation).
- ☀ Not available locally: inpatient withdrawal management facilities, OTP, homeless shelter, residential SUD facility.

# Your plan should address:

- Medical **management options**, including standard of care recommendations vs level of care patient is willing to accept and outlining a **risk-benefit conversation**
- Creative **utilization of available supports** at home and in the community (virtual recovery supports, telemedicine, family/friends, peer support, village CHAP/behavioral health aides, clergy, tribal elders and traditional healers, social services)
- **Harm reduction options**
- **Safety and follow-up planning**

#1. A 25 yo single female with PTSD at 16 WGA 1<sup>st</sup> pregnancy, referred to your addiction specialty clinic from a local primary care practice to take over her SLBUP prescribing. She has been intermittently taking SLBUP but has moved between 3 different practices in the past 2 months due to chaotic life circumstances, currently couch surfing. She frequently no-shows for visits and has many gaps in medication continuity. She reported to her PCP last week that she has been struggling to take her SLBUP daily and has continued to inject fentanyl most days, as well as using about 6mg/day of non-prescribed alprazolam and methamphetamine daily. She has had 2 admissions to WD management but left AMA on 1<sup>st</sup> day. The nearest OTP is 200 miles away and she refuses residential treatment. She is interested in stopping all drugs but unsure how she can possibly manage her anxiety without benzos. She lives 10 miles outside of town, staying with mom (strained relationship) and has no sober friends.

#2. 69 yo male veteran, PMH CAD, MI x2 (stent 6 mos ago), Leukemia CLL (off meds), Chronic Hep C and had been prescribed methadone 50mg/day for chronic pain. He has been injecting fentanyl daily for the past year due to worsening pain, along with methamphetamine to treat fatigue. His methadone prescriber discharged him from care after he tested positive for stimulants. He presents requesting assistance with withdrawing off methadone. He's interested in buprenorphine but scared about PW and worried it won't be strong enough to treat his pain and wonders what will happen if he injects fentanyl while on BUP. He says he cannot stop using meth, he needs it to function due to severe fatigue. He refuses residential treatment, he has to care for home/dog. Nearest neighbors are also PWUD. His truck is broken down and he lives 10 miles from the clinic and 20 miles from the pharmacy. He has not had labs done in over 2 years.

#3. 24 yo AK Native fisherman with h/o anxiety and chronic back pain, who lives in a roadless village (seen by telemed) requests withdrawal management for kratom (drinks 8 cups of tea/day) and bromazolam (4-6 mg/day), both ordered online, so that he can leave in 2 weeks for his 2-month commercial fishing season. When he has run out of kratom he experiences significant withdrawal and has smoked heroin a few times which helped symptoms. He has experienced a seizure once when he ran out of benzos in the past, and sometimes drinks alcohol to help with insomnia when running low on bromazolam. He will be on boat with a sober crew with no access to medical care or pharmacy for at least 2 months. He is an established patient at the local village clinic staffed by a CHAP.

#4. 54 yo AK Native female (who lives in a roadless village, seen by telemed) with schizophrenia, RA, AUD with compensated cirrhosis, and opioid dependence. She lost her opioid prescriber 3 years ago when the itinerant provider lost their DEA, then transitioned to smoking fentanyl to treat her pain. She ran out of her mood stabilizers 6 mos ago, triggering increased fentanyl use to help with sleep and hallucinations. She wants to stop using fentanyl. She drinks 1/5 of vodka daily and had seizures in the past with abstinence. She often experiences blackouts from drinking and injuries from falls, last year medi-evoked to ANMC for hip fracture. She wants to cut down on her alcohol consumption, but she is apprehensive about stopping. She presents with her husband today, also seeking treatment, who revived her from an overdose last week. He is also dependent on fentanyl but does not drink. She relies on him to obtain alcohol for her and describes him as supportive. She refuses to go to the ANMC hospital for inpatient care (her mother died there) but she does utilize the village clinic staffed by a CHAP.



# Final Takeaways/Summary

- ☀️ Current society guidelines for withdrawal management do not always address the management of patients with complex needs in resource limited communities.
- ☀️ Some patients can benefit from a low-threshold, harm reduction approach to withdrawal management when a higher level of care is not accessible.
- ☀️ Providers should discuss risks and benefits of various withdrawal management options while considering creative usage of available supports to reduce harms and maximize quality of life for patients with severe, complex SUDs.

# References

1. The ASAM Clinical Practice Guideline on Alcohol Withdrawal Management 2022 <https://www.asam.org/quality-care/clinical-guidelines/alcohol-withdrawal-management-guideline>
2. The ASAM National Practice Guideline for the Treatment of Opioid Use Disorder 2020 <https://www.asam.org/quality-care/clinical-guidelines/national-practice-guideline>
3. The ASAM/AAAP Clinical Practice Guideline on the Management of Stimulant Use Disorder 2023 <https://www.asam.org/quality-care/clinical-guidelines/stimulant-use-disorders>
4. Tiglao SM, et al, Alcohol Withdrawal Syndrome: Outpatient Management. Am Fam Physician. 2021 Sep <https://pubmed.ncbi.nlm.nih.gov/34523874/>
5. Smith-Bernardin, S.M., et al. Scoping review of managed alcohol programs. Harm Reduct J 19, 82 (2022). <https://doi.org/10.1186/s12954-022-00646-0>

# References

6. Munipati D, et al. Rapid detoxification in an adult with benzodiazepine dependence with the aid of an antiepileptic. *Prim Care Companion CNS Disord.* 2023;25(4):22cr03457.

<https://doi.org/10.4088/PCC.22cr03457>

7. Mireya Pérez Orts, et al, The Evolution Toward Designer Benzodiazepines in Drug-Facilitated Sexual Assault Cases, *Journal of Analytical Toxicology*, Volume 47, Issue 1, January 2023,

<https://doi.org/10.1093/jat/bkac017>

8. Cohen, Shawn M. MD; et al, Low Dose Initiation of Buprenorphine: A Narrative Review and Practical Approach. *Journal of Addiction Medicine* 16(4):p 399-406, 7/8 2022. DOI: 10.1097/ADM.0000000000000945

9. D’Onofrio G, et al. Incidence of Precipitated Withdrawal During a Multisite Emergency Department–Initiated Buprenorphine Clinical Trial in the Era of Fentanyl. *JAMA Netw Open.* 2023;6(3):e236108.

<https://jamanetwork.com/journals/jamanetworkopen/fullarticle/2802898>

# Resources

1. Bridge to Treatment <https://bridgetotreatment.org/tools/resources/>
2. Curbsiders' Addiction Medicine, Get in the Spirit of Ambulatory Alcohol Withdrawal, July 2022 <https://thecurbsiders.com/addiction-medicine-podcast/2-get-in-the-spirit-of-ambulatory-alcohol-withdrawal>
3. Oregon Pain Guidance Opioid and Benzodiazepine Taper Flowsheets <https://www.oregonpainguidance.org/app/content/uploads/2016/05/Opioid-and-Benzodiazepine-Tapering-flow-sheets.pdf>
4. The Ashton Manual; Benzodiazepines: How They Work and How to Withdraw <https://www.benzoinfo.com/ashtonmanual/#contents>
5. Alaska Native Tribal Health Consortium Harm Reduction Toolkit <https://www.iknowmine.org/topic/harm-reduction/>
6. NEXT Distro: online and mail-based harm reduction services <https://nextdistro.org/>

Sarah Spencer DO, FASAM  
Addiction Medicine Specialist & Consultant

Ninilchik Traditional Council Community Clinic  
Ninilchik, Alaska  
Cell 907-299-7460

[sarahspencerak@gmail.com](mailto:sarahspencerak@gmail.com)

