



# Fall Update: Managing Pain in 2024

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# Objectives

**To help the provider to improve treating acute and chronic pain in 2024 with better understanding:**

1. How to use multi-modal pain treatment approaches to avoid excessive and prolonged doses of opioids
2. Recently updated definitions for pain
3. Pain signaling pathways and therapeutic targets
4. The new 2022 CDC Guidelines for Using Opioids
5. Principles of assessment and treatment for acute and chronic pain
6. When to consider using buprenorphine for chronic pain
7. Additional considerations

**CONFLICTS: None**

**Updated Pain  
Definitions and Management  
Goals**

## 2020 Revised Definition of Pain

### Pain

An unpleasant sensory and emotional experience associated with, or resembling that associated with, actual or potential tissue damage.

### Notes

- Pain is always a personal experience that is influenced to varying degrees by biological, psychological, and social factors.
- Pain and nociception are different phenomena. Pain cannot be inferred solely from activity in sensory neurons.
- Through their life experiences, individuals learn the concept of pain.
- A person's report of an experience as pain should be respected.
- Although pain usually serves an adaptive role, it may have adverse effects on function and social and psychological well-being.
- Verbal description is only one of several behaviors to express pain; inability to communicate does not negate the possibility that a human or a nonhuman animal experiences pain.

## Pain Terms and Definitions

- **Allodynia:** Pain due to a stimulus that does not normally provoke pain.
- **Hyperalgesia:** Increased pain from a stimulus that normally provokes pain.
- **Nociceptive pain:** Pain that arises from actual or threatened damage to non-neural tissue and is due to the activation of nociceptors.
- **Central sensitization:** Increased responsiveness of nociceptive neurons in the central nervous system to their normal or subthreshold afferent input.
- **Neuropathic pain:** Pain caused by a lesion or disease of the somatosensory nervous system.
- **Nociplastic pain:** Pain that arises from altered nociception despite no clear evidence of actual or threatened tissue damage causing the activation of peripheral nociceptors or evidence for disease or lesion of the somatosensory system causing the pain.

*IASP Terminology. International Association for the Study of Pain. Available at: <https://www.iasp-pain.org/Education/Content.aspx?ItemNumber=1698> (Accessed on December 1, 2019).*

## GOAL FOR ADEQUATE PAIN CONTROL

The goal for pain control should not be zero pain, but rather a tolerable level of pain that allows physical and emotional function. Often this means balancing analgesia with achieving functional goals, while avoiding preventable complications.

## PRINCIPLES

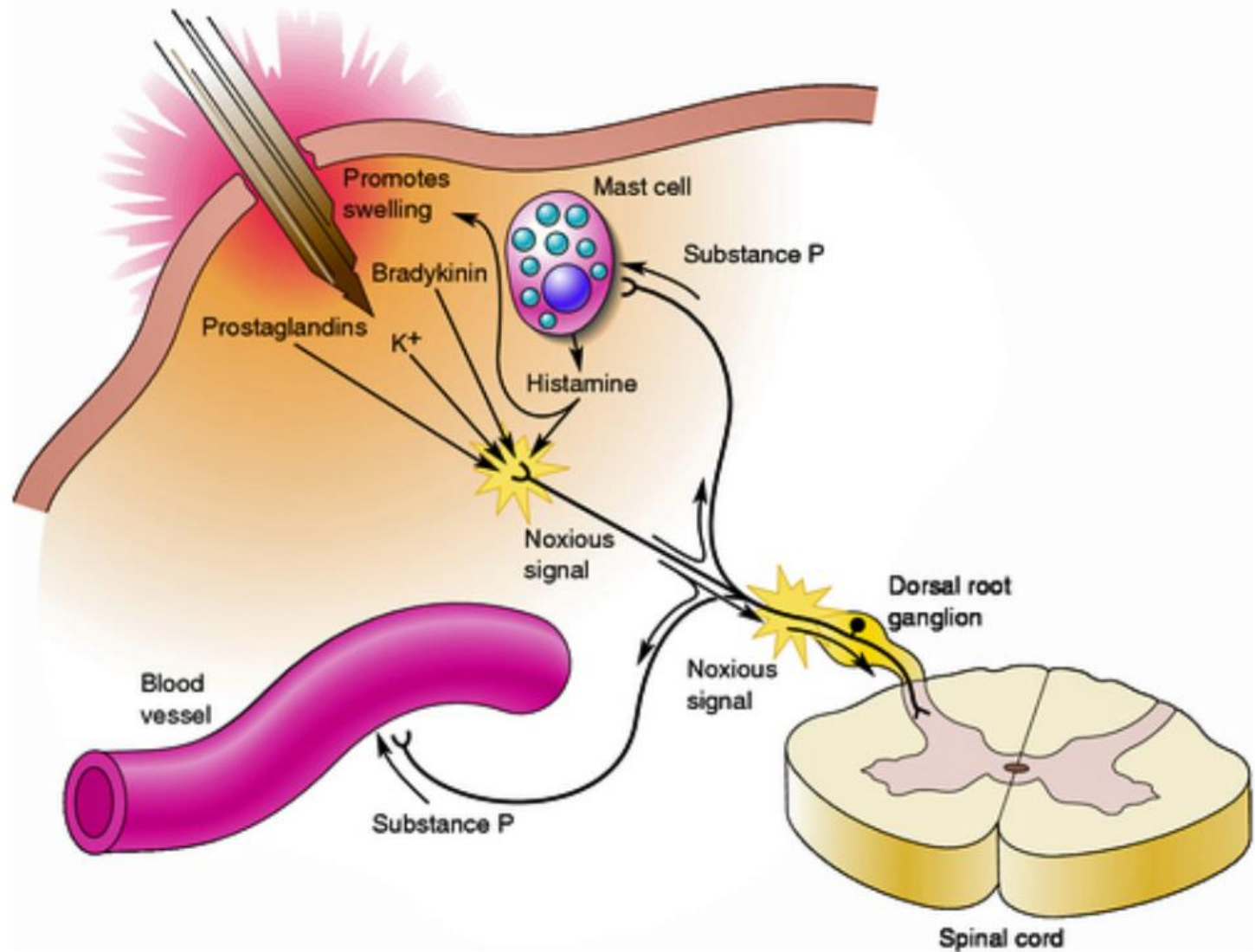
1. Create an individualized plan for pain management based on the expected degree of pain and patient factors that may affect the plan
2. Offer multimodal analgesia, adding opioids only as necessary
3. Provide patient education
4. Adjust the pain management plan based on adequacy of pain relief and the occurrence of adverse events

## TACTICAL APPROACH

1. **Use multimodal analgesia** — Use a multimodal approach to analgesia for acute pain, with nonpharmacologic techniques, regional anesthesia techniques as appropriate, nonopioid analgesics, and opioids only as necessary. Multimodal analgesia involves the use of two or more agents that employ different mechanisms for pain management, thereby reducing overreliance on and adverse effects from a single class of agents, most importantly opioids.
2. **Use opioids safely** — An overarching principle of acute pain management is to avoid excessive or prolonged use of opioids. Opioids are associated with short term side effects (eg, respiratory depression, sedation, nausea and vomiting, pruritus, urinary retention, constipation) and long term adverse effects (eg, tolerance, dependence, opioid induced hyperalgesia, withdrawal upon conclusion of therapy, opioid use disorder, and overdose).

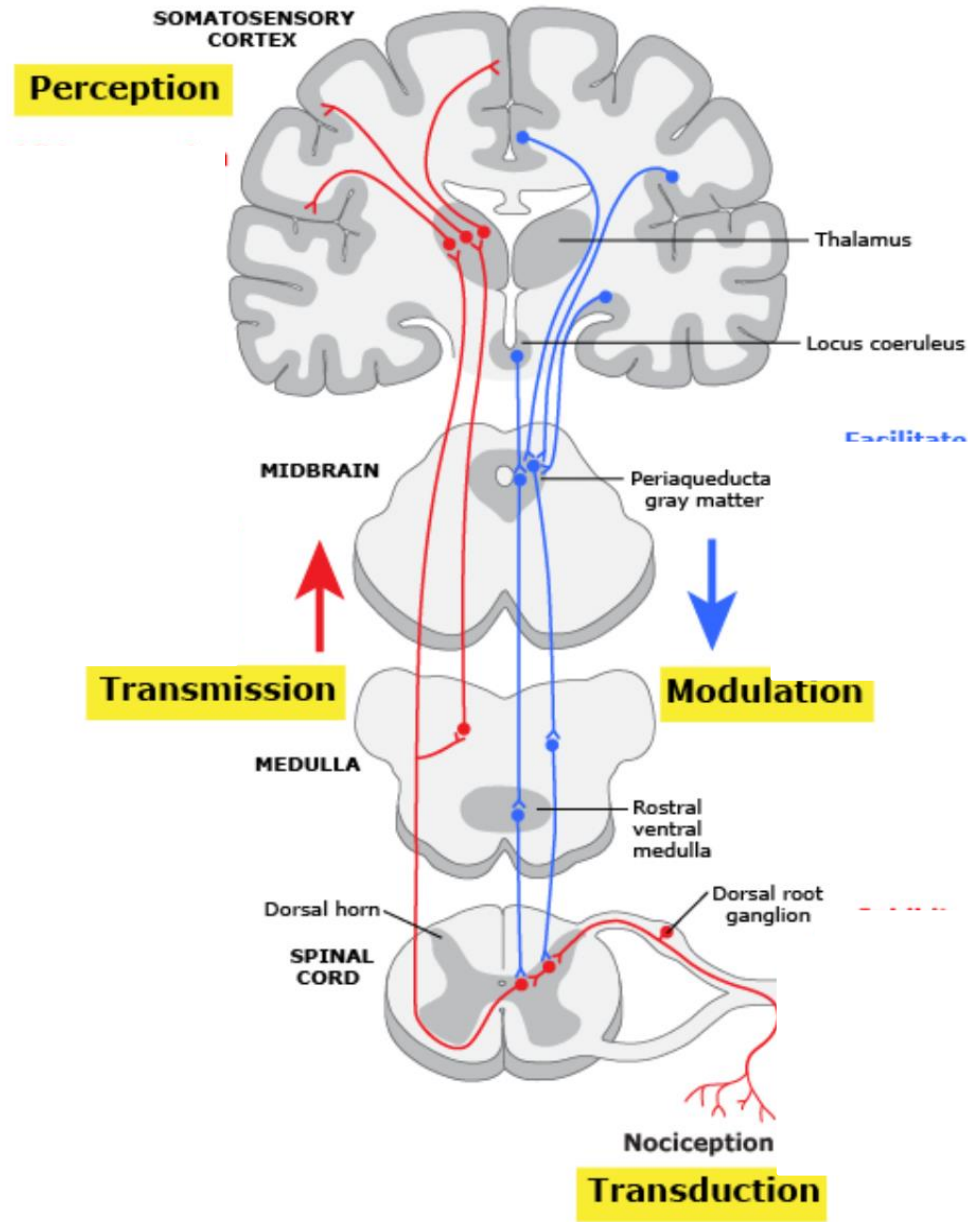
# Pain Signaling Pathways

## Pain Signaling Pathways





# Pain Signaling Pathways



# **Key Principles of Pain Evaluation**

## History and Physical

- Identify possible pain etiology
- Identify comorbidities that may affect treatment options
- Examine for allodynia and/or sensory changes in painful body part
- For patients who use opioids or patients with risk factors for opioid misuse or use disorder:
  - Check PDMP
  - Screen for opioid risk with ORT, SOAPP, COMM, or similar

PDMP: prescription drug monitoring programs; ORT: Opioid Risk Tool; SOAPP: Screener and Opioid Assessment for Patients with Pain; COMM: Current Opioid Misuse Screen

### Body diagram

- Useful for all patients

- For patients with multisite pain, screen for chronic widespread pain disorders with Widespread Pain Index and Symptom Severity Score

### Pain history

#### OLDCARTS

- **O**nset ("When did your pain start?")

- **L**ocation ("Where does it hurt?")

- **D**uration ("How long does your pain last?")

- **C**haracter ("How does your pain feel?", ie, aching, burning, shooting, tingling)

- **A**lleviating/**A**ggravating ("What makes your pain better/worse?") and **A**tribution ("What do you think is the cause?")

- **R**adiation ("Does this pain spread anywhere else?")


- **T**emporal pattern ("Does your pain vary over the course of a day?")

- **S**ymptoms associated ("How does your pain impact your physical function, your mood, your sleep?")

## Visual Pain Scales

**A**

**Visual analog scale**  
Place a mark on the line below to indicate how bad your pain feels.

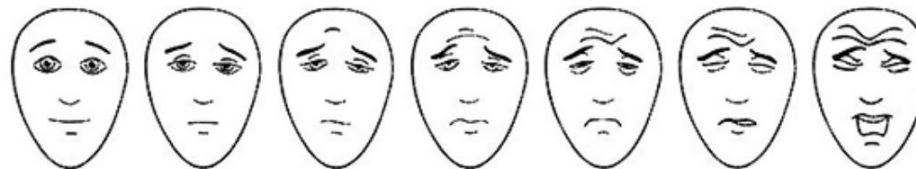
No pain  Worst pain imaginable

**B**

**Numeric rating scale**  
What does your pain feel like?

0 1 2 3 4 5 6 7 8 9 10

None Mild Moderate Very bad Unbearable



Schematic representation of the faces pain scale, rated from 0 to 6 left to right.



## Mood assessment

### PHQ-4

Over the past 2 weeks, have you been bothered by these problems?	Not at all	Several days	More days than not	Nearly every day
•Feeling nervous, anxious, or on edge	0	1	2	3
•Not being able to stop or control worrying	0	1	2	3
•Feeling down, depressed, or hopeless	0	1	2	3
•Little interest or pleasure in doing things	0	1	2	3

•**Scoring:**Add total score

•**For score >5**, screen for anxiety, depression, and post-traumatic stress, with GAD-7, PHQ-9, and PTSD-5

## Sleep assessment

### Sleep initiation and maintenance

•Does pain interfere with falling asleep?

•Does pain interfere with staying asleep?

### Screen for obstructive sleep apnea (OSA) – STOP-Bang<sup>[2,3]</sup>

Yes	No	Snore – Do you snore loudly (loud enough to be heard through closed doors, or your bed partner elbows you for snoring at night)?
Yes	No	Tired – Do you often feel tired, fatigued, or sleepy during the day?
Yes	No	Observed – Has anyone observed you stop breathing or choking/gasping during sleep?
Yes	No	Pressure – Do you have or are you being treated for high blood pressure?
Yes	No	Body mass index >35 kg/m <sup>2</sup> ?
Yes	No	Age older than 50 years?
Yes	No	Neck size large (male: ≥17 inches, female: ≥16 inches)?
Yes	No	Gender = male?

•**Scoring:**Low risk of OSA: Yes to 0 to 2 questions

•Intermediate risk of OSA: Yes to 3 to 4 questions

•High risk of OSA: Yes to ≥5 questions

1.Reproduced from: Kroenke K, Spitzer RL, Williams JB, Löwe B. An ultra-brief screening scale for anxiety and depression: The PHQ-4. Psychosomatics 2009; 50:613. Table used with the permission of Elsevier Inc. All rights reserved.

2.Chung F, Subramanyam R, Liao P, et al. High STOP-Bang score indicates a high probability of obstructive sleep apnoea. Br J Anaesth 2012; 108:768.



# Confirm Appropriateness for Prescribing Opioids (DIRE Score)

Name: \_\_\_\_\_ DOB \_\_\_/\_\_\_/\_\_\_

## DIRE Score: Patient Selection for Chronic Opioid Analgesia

For each factor, rate the patient's score from 1-3 based on the explanations in the right-hand column

SCORE	FACTOR	EXPLANATION
	<b>DIAGNOSIS</b>	1 = Benign chronic condition with minimal objective findings or no definite medical diagnosis. Examples: fibromyalgia, migraine headaches, non-specific back pain. 2 = Slowly progressive condition concordant with moderate pain, or fixed condition with moderate objective findings. Examples: failed back surgery syndrome, back pain with moderate degenerative changes, neuropathic pain. 3 = Advanced condition concordant with severe pain with objective findings. Examples: severe ischemic vascular disease, advanced neuropathy, severe spinal stenosis.
	INTRACTABILITY	1 = Few therapies have been tried and the patient takes a passive role in his/her pain management process. 2 = Most customary treatments have been tried but the patient is not fully engaged in the pain management process, or barriers prevent (insurance, transportation, medical illness). 3 = Patient fully engaged in a spectrum of appropriate treatments but with inadequate response.
<b>RISK</b>		<b>(R = Total of P+C+R+S below)</b>
	<del>Psychological</del>	1 = Serious personality dysfunction or mental illness interfering with care. Example: personality disorder, severe affective disorder, significant personality issues. 2 = Personality or mental health interferes moderately. Example: depression or anxiety disorder. 3 = Good communication with clinic. No significant personality dysfunction or mental illness.
	Chemical Health	1 = Active or very recent use of illicit drugs, excessive alcohol, or prescription drug abuse. 2 = Chemical copier (uses medications to cope with stress) or history of chemical dependence (CD) in remission. 3 = No CD history. Not drug-focused or chemically reliant.
	Reliability	1 = History of numerous problems: medication misuse, missed appointments, rarely follows through. 2 = Occasional difficulties with compliance, but generally reliable. 3 = Highly reliable patient with meds, appointments & treatment.
	Social Support	1 = Life in chaos. Little family support and few close relationships. Loss of most normal life roles. 2 = Reduction in some relationships and life roles. 3 = Supportive family/close relationships. Involved in work or school and no social isolation.
	<b>EFFICACY SCORE</b>	1 = Poor function or minimal pain relief despite moderate to high doses. 2 = Moderate benefit with function improved in several ways (or insufficient info - hasn't tried opioid yet or very low doses or too short of a trial). 3 = Good improvement in pain and function and quality of life with stable doses over time.

\_\_\_\_\_ **Total score= D + I + R + E**

**Score 7-13:** Not a suitable candidate for long-term opioid analgesia

**Score 14-21:** May be a good candidate for long-term opioid analgesia

### NOTES

A DIRE Score of :513 indicates that the patient may not be suited to long-term opioid pain management. Used with permission by Miles J. Belgrade, MD |



# Assess Risk for Overdose (Using RIOSORD)

Risk Index for Overdose or Serious Opioid-Induced Respiratory Depression (RIOSORD)		
Question	Points for Positive Response	Actual Response
<b>In the past 6 mo, has the patient had a health care visit (outpatient, inpatient, or emergency department) involving any of the following health conditions</b>		
Substance use disorder (abuse or dependence), including alcohol, amphetamines, antidepressants, cannabis, cocaine, hallucinogens, opioids, and sedatives	25	
Bipolar disorder or schizophrenia	10	
Stroke or other cerebrovascular disease	9	
Kidney disease with clinically significant renal impairment	8	
Heart failure	7	
Nonmalignant pancreatic disease (e.g., acute or chronic pancreatitis)	7	
Chronic pulmonary disease (e.g., emphysema, chronic bronchitis, asthma, pneumoconiosis, asbestosis)	5	
Recurrent headache (e.g., migraine)	5	
<b>Does the patient use any of the following substances?</b>		
Fentanyl	13	
Morphine	11	
Methadone	10	
Hydromorphone	7	
<b>Does the patient use an extended-release or long-acting formulation of any prescription opioid?</b>		
Prescription benzodiazepine (e.g., diazepam, alprazolam)	9	
Prescription antidepressant (e.g., fluoxetine, citalopram, venlafaxine, amitriptyline)	8	
<b>Is the patient's current maximum prescribed daily morphine-equivalent dose <math>\geq 100</math> mg for all opioids used on a regular basis?</b>		
	7	
<b>Total possible score</b>	<b>146</b>	

Risk Classes and Predicted Probability of Serious Opioid-Induced Respiratory Depression during the Next 6 Months.			
Risk Class	RIOSORD Score	Average Predicted Probability (Percent)	Actual Observed Incidence (Percent)
1	<5	1.9	2.1
2	5–7	4.8	5.4
3	8–9	6.8	6.3
4	10–17	15.1	14.2
5	18–25	29.8	32.2
6	26–41	55.1	58.8
7	$\geq 42$	83.4	82.4

# Check the Michigan Automated Prescription Service (MAPS) to Assess for Potential Misuse, Abuse, Diversion, or Overdose Risk

Menu Admin Patient Alerts Henry Smith

RxSearch > Patient Request > Justin Cooper

STATE DEPARTMENT OF HEALTH  
Powered by NarxCare™

**Justin Cooper, 37M**

Narx Report Resources

Date: 06/15/2017 Download PDF Download CSV

Justin Cooper

Risk Indicators

NARX SCORES			OVERDOSE RISK SCORE	ADDITIONAL RISK INDICATORS (2)
Narcotic	Sedative	Stimulant	<b>650</b> (Range 0-999)	<ul style="list-style-type: none"> <li>Active MME &gt; Threshold</li> <li>Patient has Benzodiazepine/ Narcotic overlap</li> </ul>
<b>672</b>	<b>512</b>	<b>190</b>		
Explain these scores			Explain this score	Explain these indicators

Graphs

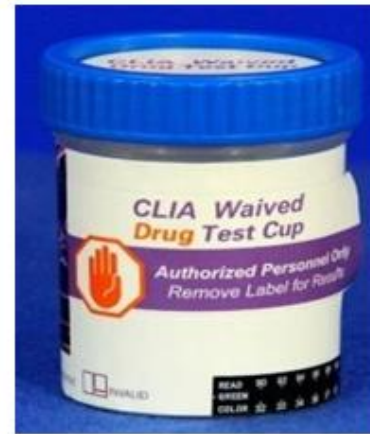
RX GRAPH  Narcotic  Sedative  Stimulant

All Prescribers

Prescribers	10	9	8	7	6	5
10. King, James						
9. Hawkins, Norma						
8. Jenknis, Gerald						
7. Ramos, Jesse						
6. Jackson, Janice						
5. Medina, Martha						

# Check for Anticipated and Unanticipated Medications and Other Substances

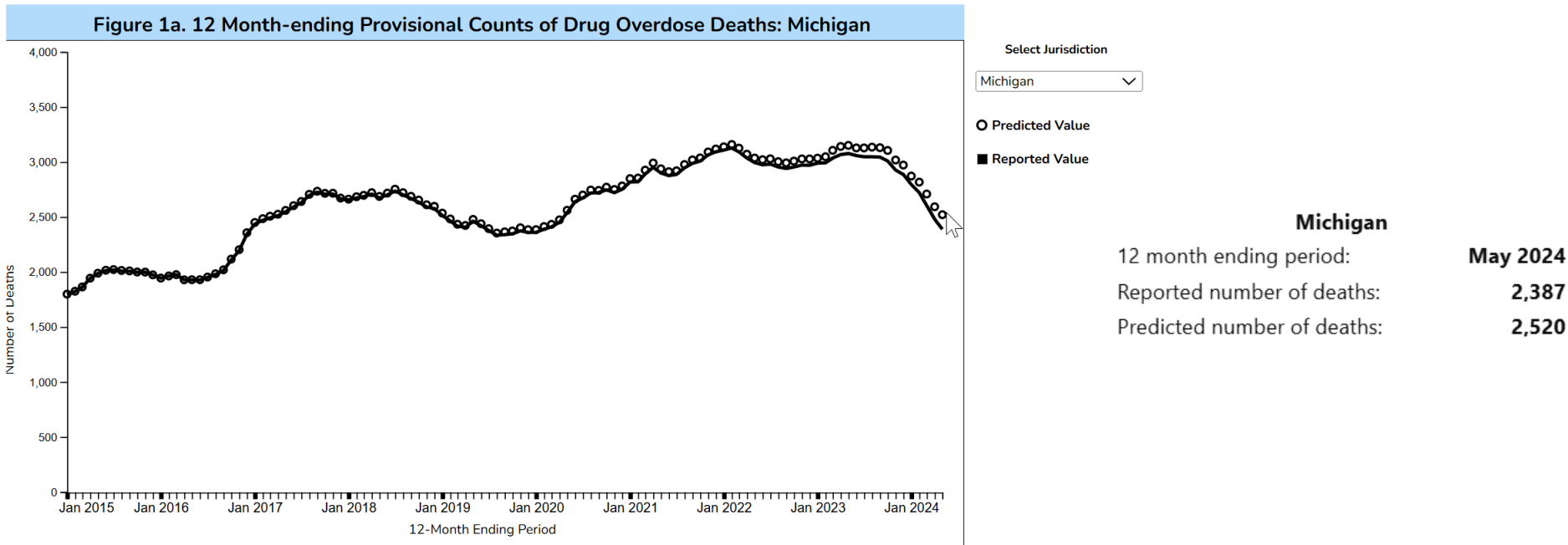
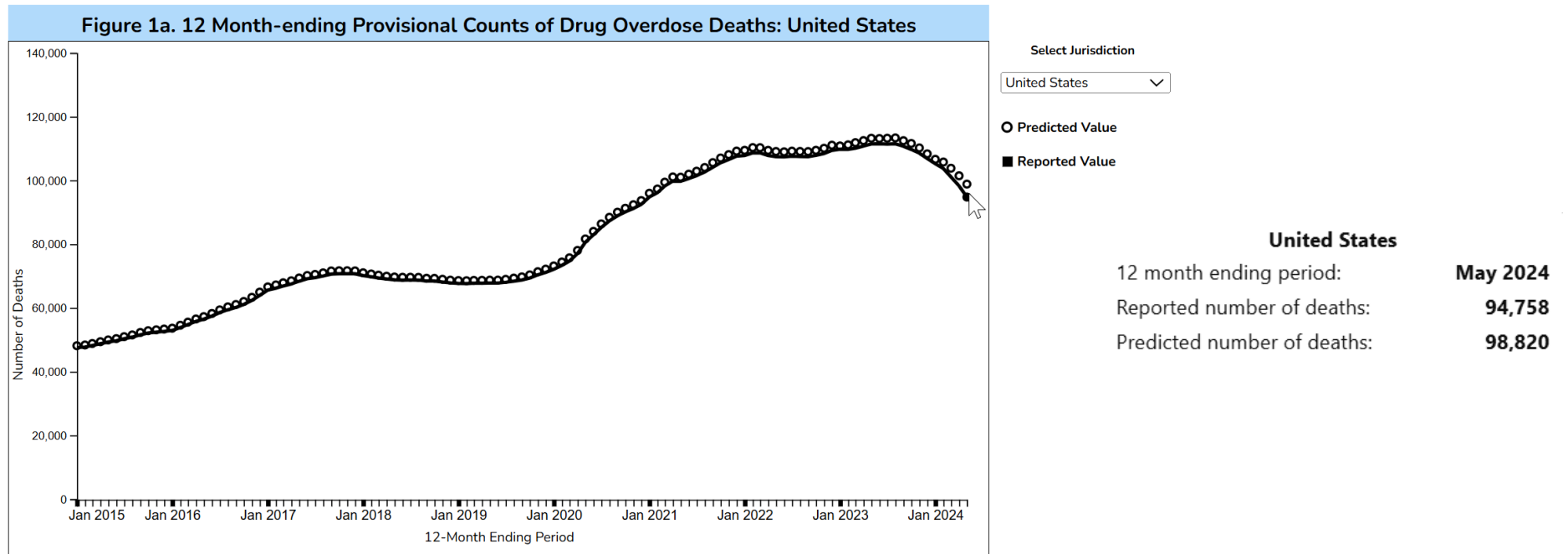
- If you don't check, you will have no idea.
- Qualitative - in Office
  - Test either positive or negative
  - Immunoassay
- Quantitative (in Lab)
  - Test measures concentration of drug
  - GC/MS or LC/MS
  - Can check for all psychoactive substances prescribed and unprescribed
  - Matrix - Urine, Saliva, Blood, Hair, Exhaled Air (breathalyzer), etc.



**Consider What Else the Patient May  
Be Misusing - The Overdose  
Epidemic is Turning the Corner,  
But It Is Far From Over**

# 12 Month-Ending Provisional Number and Percent Change of Drug Overdose Deaths

Based on data available for analysis on: October 6, 2024



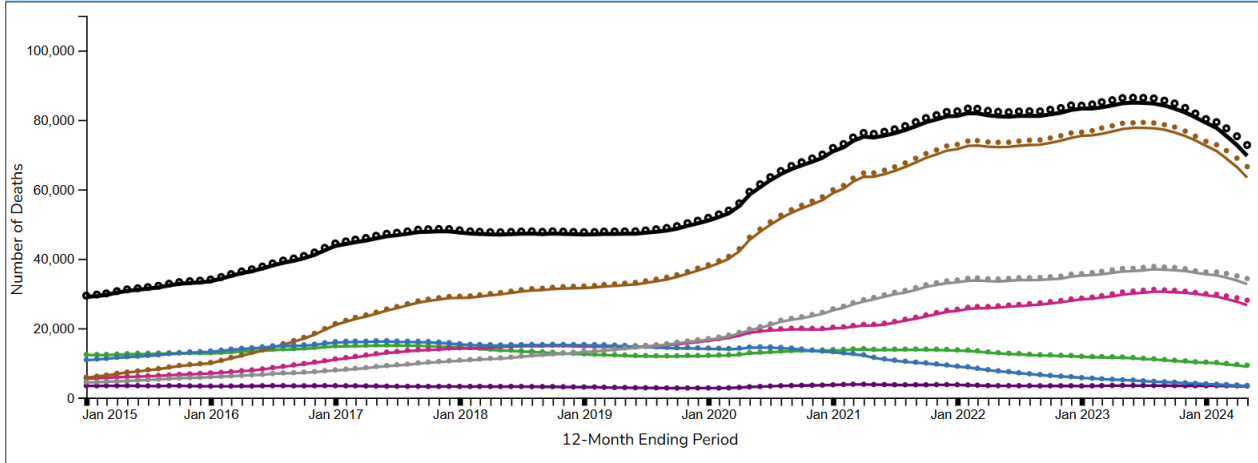
# 12 Month-Ending Provisional Number of Drug Overdose Deaths by Drug or Drug Class

Based on data available for analysis on: October 6, 2024

After opening the **drug class dropdown**, click the top of the dropdown menu again to make the checkboxes disappear.

Select Jurisdiction:  Select specific drugs or drug classes:

**Figure 2. 12 Month-Ending Provisional Number of Drug Overdose Deaths by Drug or Drug Class: United States**



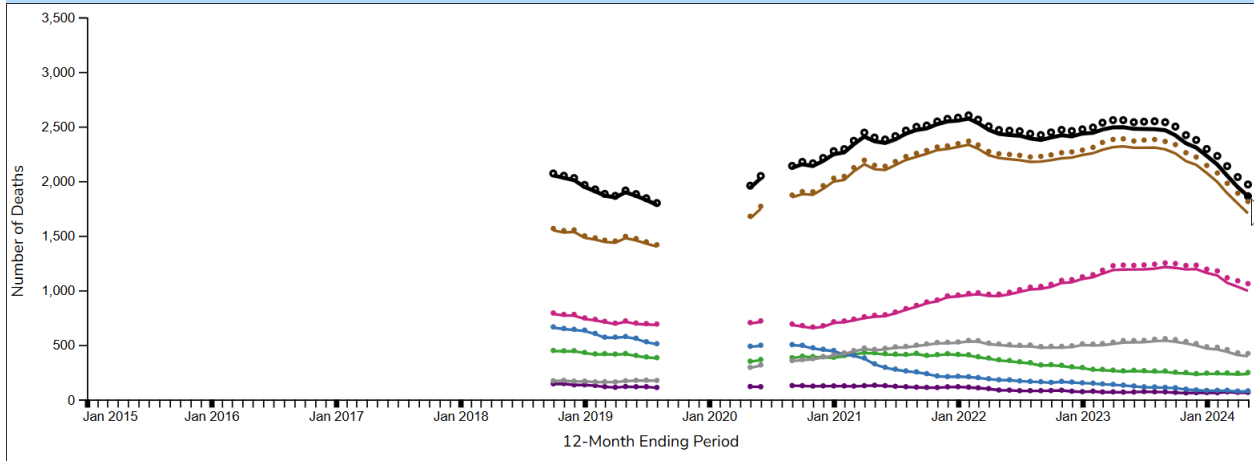
**United States, May 2024,  
Opioids (T40.0-T40.4, T40.6)**

Reported number of deaths:	<b>69,649</b>
Predicted number of deaths:	<b>72,755</b>

**Legend for Drug or Drug Class**

- Cocaine (T40.5)
- Heroin (T40.1)
- Methadone (T40.3)
- Natural & semi-synthetic opioids (T40.2)
- Opioids (T40.0-T40.4, T40.6)
- Psychostimulants with abuse potential (T43.6)
- Synthetic opioids, excl. methadone (T40.4)
- Reported Value
- Predicted Value

**Figure 2. 12 Month-Ending Provisional Number of Drug Overdose Deaths by Drug or Drug Class: Michigan**



**Michigan, May 2024,  
Opioids (T40.0-T40.4, T40.6)**

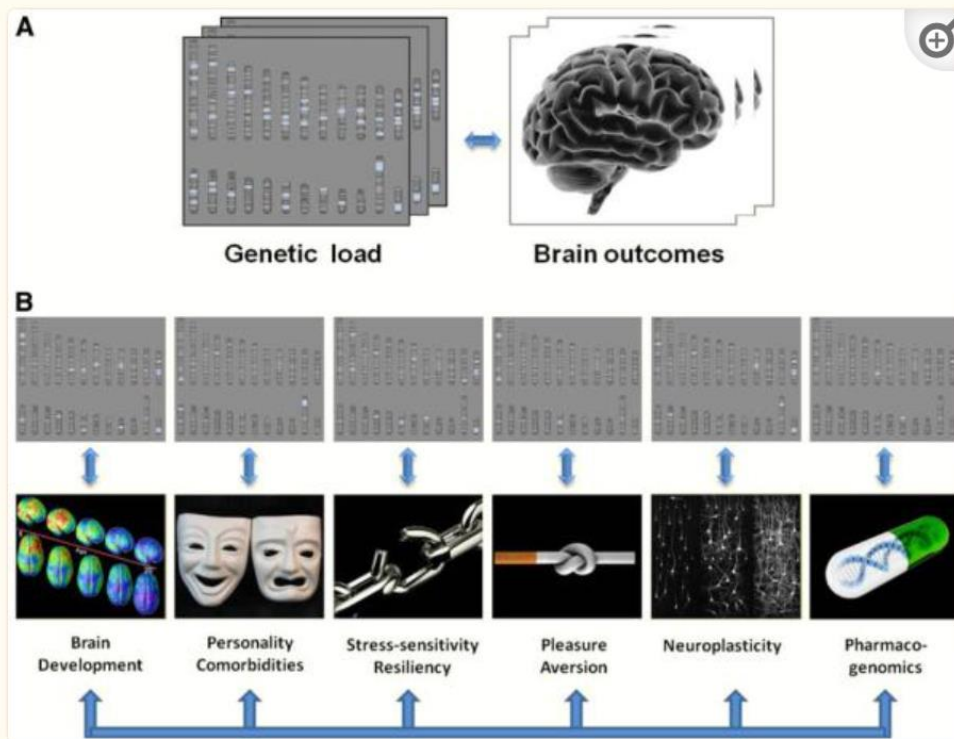
Reported number of deaths:	<b>1,862</b>
Predicted number of deaths:	<b>1,969</b>

# Epigenetic Effects of Psychoactive Substances – Prolonged Exposure Can Lead to Changes in Brain Cell DNA and Behavior

At least 3 different epigenetic mechanisms in chromatin have been identified: 1) DNA methylation, 2) histone modification, and 3) non-coding RNA (ncRNA)-associated gene silencing.

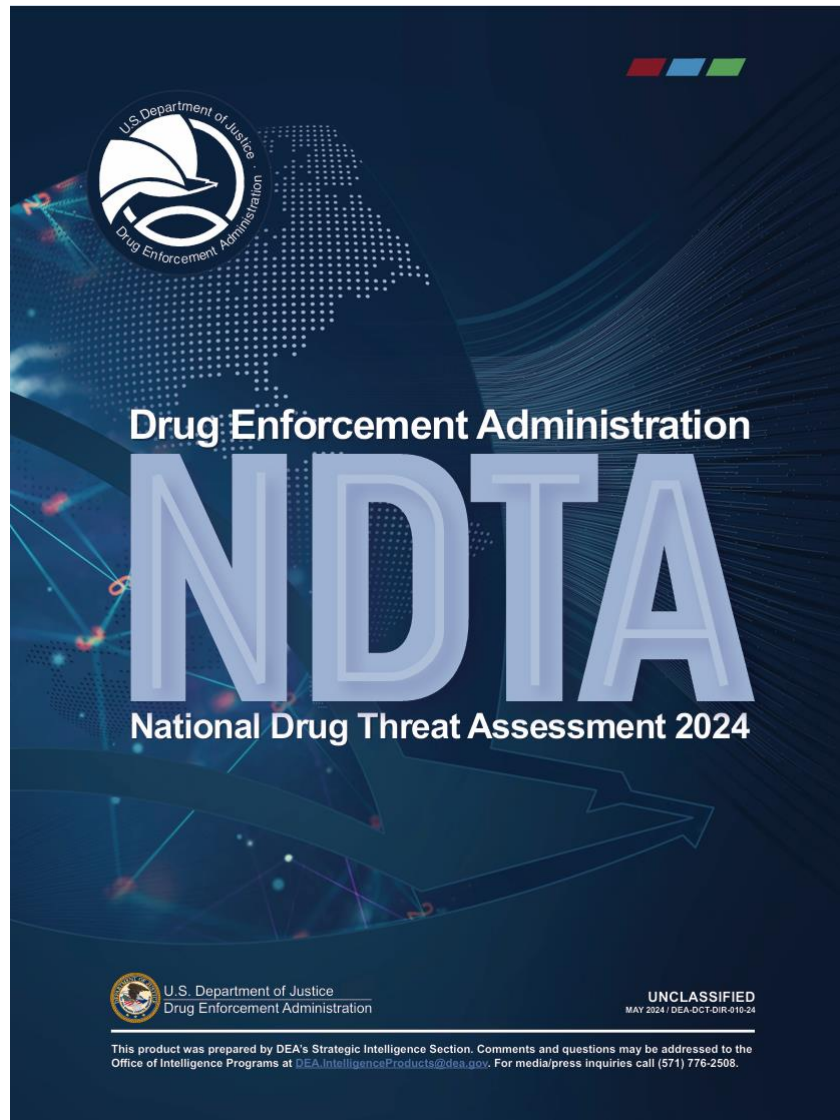
Substances that Demonstrate Epigenetic Effects

1. **Opioids**
2. Benzodiazepines
3. Amphetamines
4. Cocaine
5. Methamphetamine
6. Antipsychotics
7. Antidepressants
8. Cannabinoids
9. Kratom
10. Nicotine
11. Others

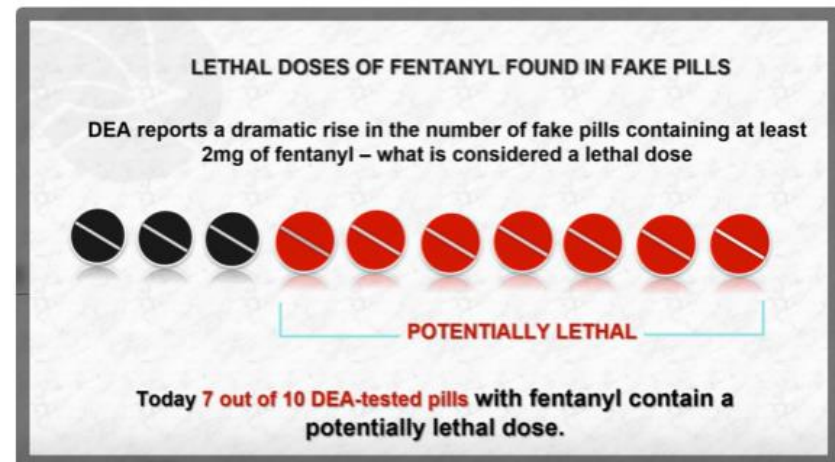




# DEA: Synthetic Drugs Driving the Overdose Landscape



*“The shift from plant-based drugs, like heroin and cocaine, to synthetic, chemical-based drugs, like fentanyl and methamphetamine, has resulted in the most dangerous and deadly drug crisis the United States has ever faced,”*

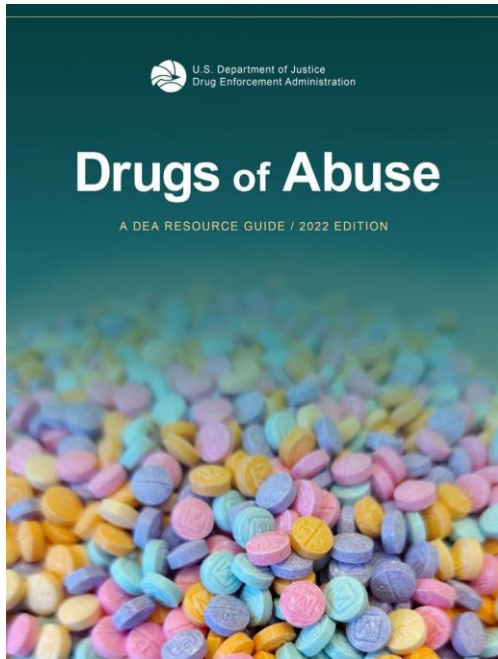


Xylazine is making the deadliest drug threat our country has ever faced, fentanyl, even deadlier.  
DEA Public Safety Alert March 2023

Thirty-one percent of the drug-related deaths in the United States include psychostimulants – mostly methamphetamine.



# DEA: Multiple Drugs of Abuse on the Street



Counterfeit Oxycodone Tablet Containing Fentanyl



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Xylazine

Nitazines

# Polysubstance Use is Common

- Polysubstance use has markedly increased throughout the entire global pandemic (2020 – 2023)
- According to the National Survey On Drug Use and Health from 2019, people who use one substance often use another.
- Many individuals with one substance use disorder are at risk of having a concurrent substance use disorder.
- Among people with a cocaine use disorder, nearly 60% have a co-occurring alcohol use disorder and over 20% have a marijuana use disorder, and among people with an opioid use disorder, more than 25% have at least two other substance use disorders.
- Finally, people with mental health disorders have been found to have higher rates of substance use and substance use disorders versus the general population.
- Having a mental disorder can increase the risk for developing multiple substance use disorders.

## Concurrent Substance Use

Substance	Marijuana	Opioid	Cocaine	Methamphetamine	Alcohol
<b>Heavy Alcohol Use</b>	45%	9.2%	11%	1.7%	
<b>Heavy Marijuana Use</b>		16%	16.3%	3.3%	16.2%
<b>Opioid</b>	53%		15.6%	8.7%	14.7%
<b>Methamphetamine</b>	68.1%	43.7%	32.2%		13.4%

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AMERICAN  
PSYCHIATRIC  
ASSOCIATION 

# PSYCHIATRIC NEWS

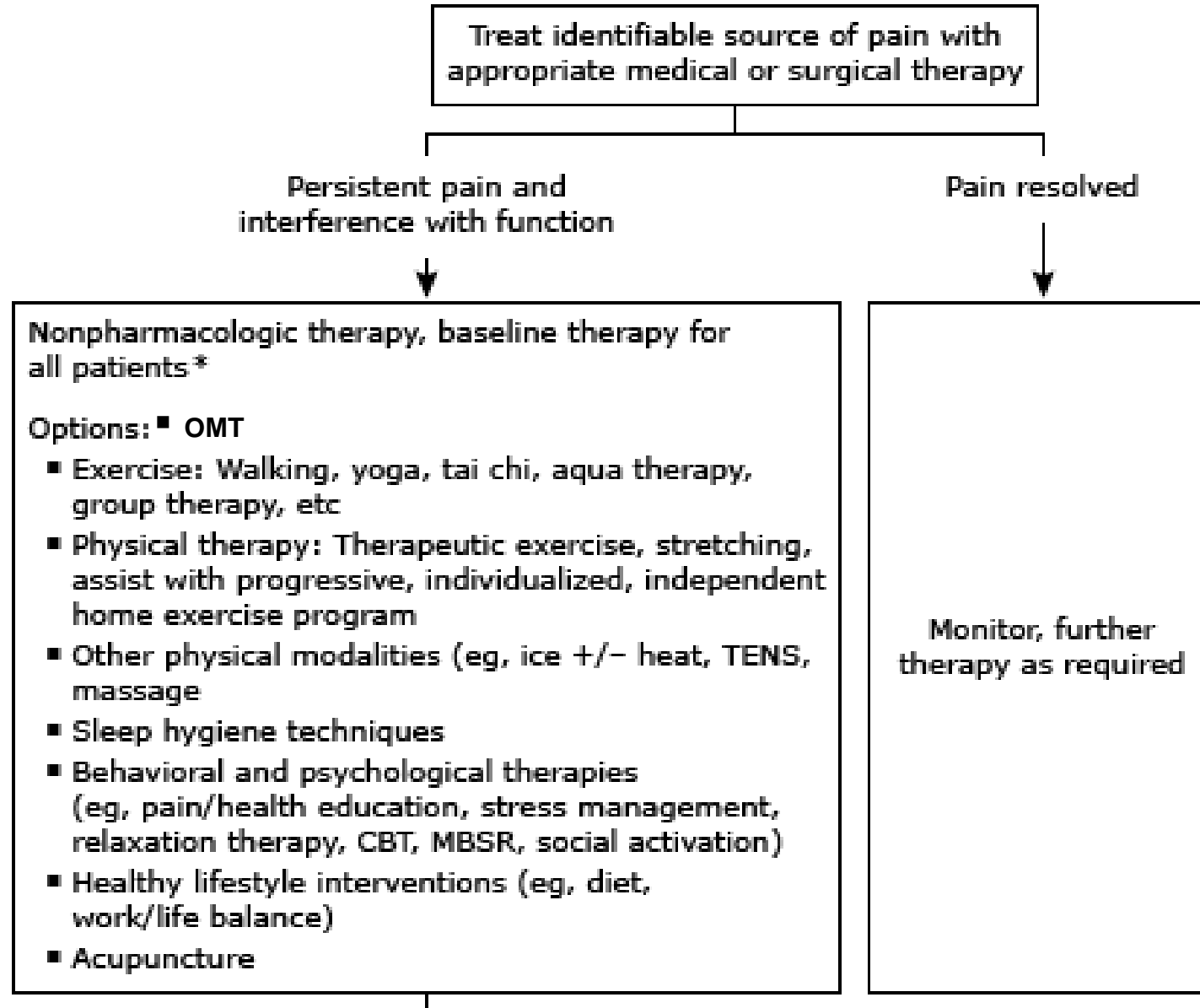
PSYCHNEWS.ORG

ISSN 0033-2704



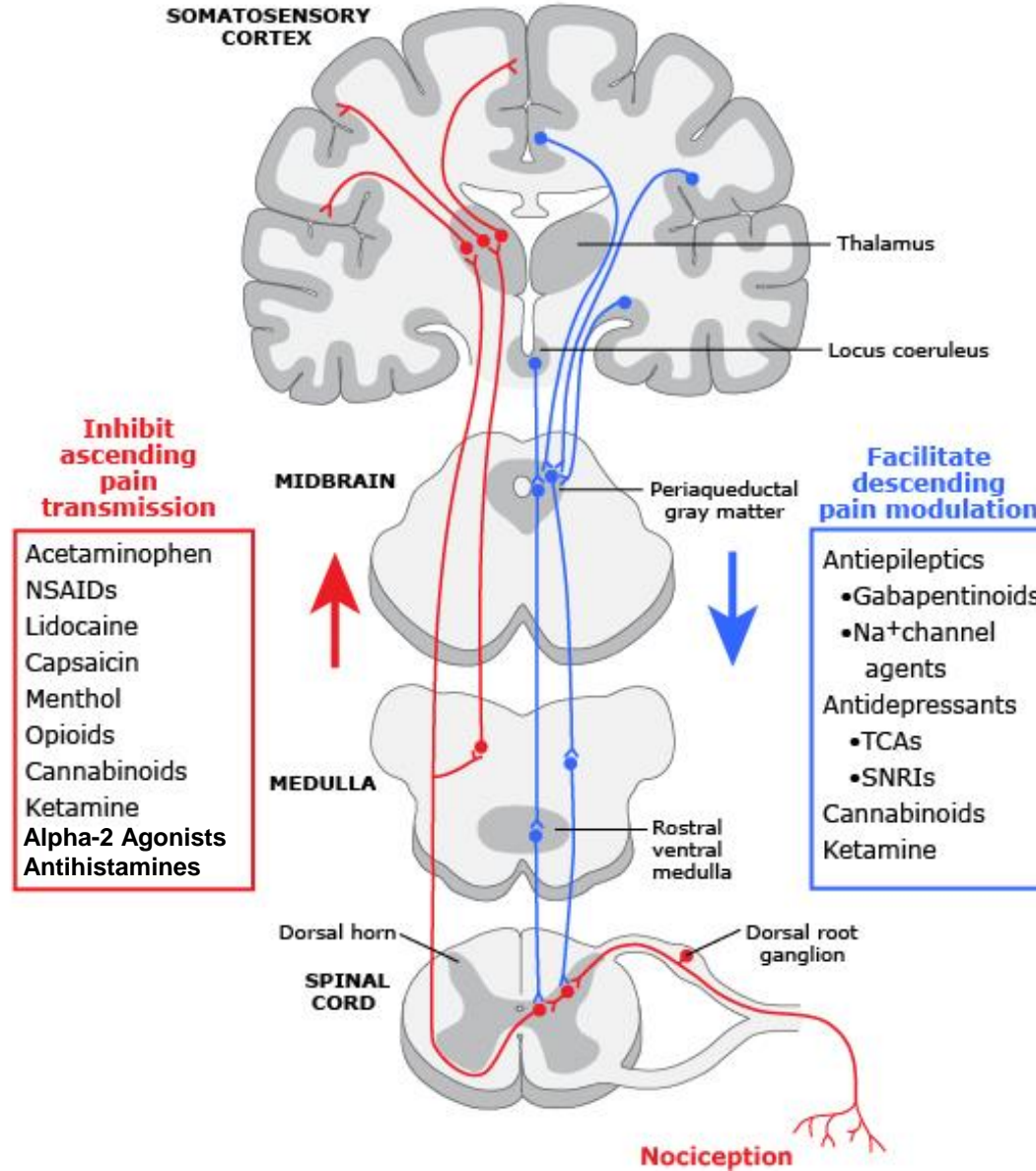
URL: [Special Report: The Art of Treating Complex Substance Use Disorders | Psychiatric News](#)

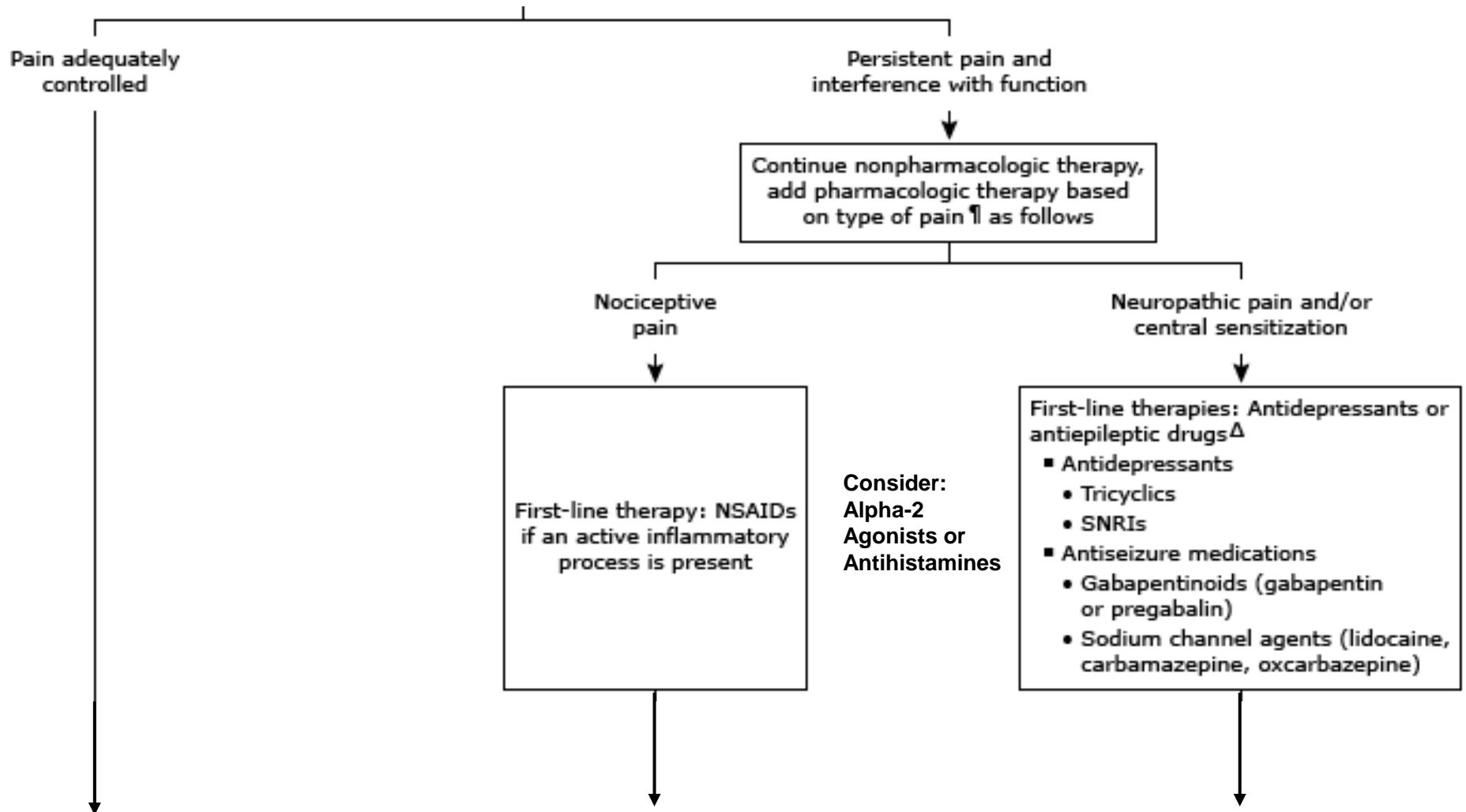
# Therapeutic Approaches

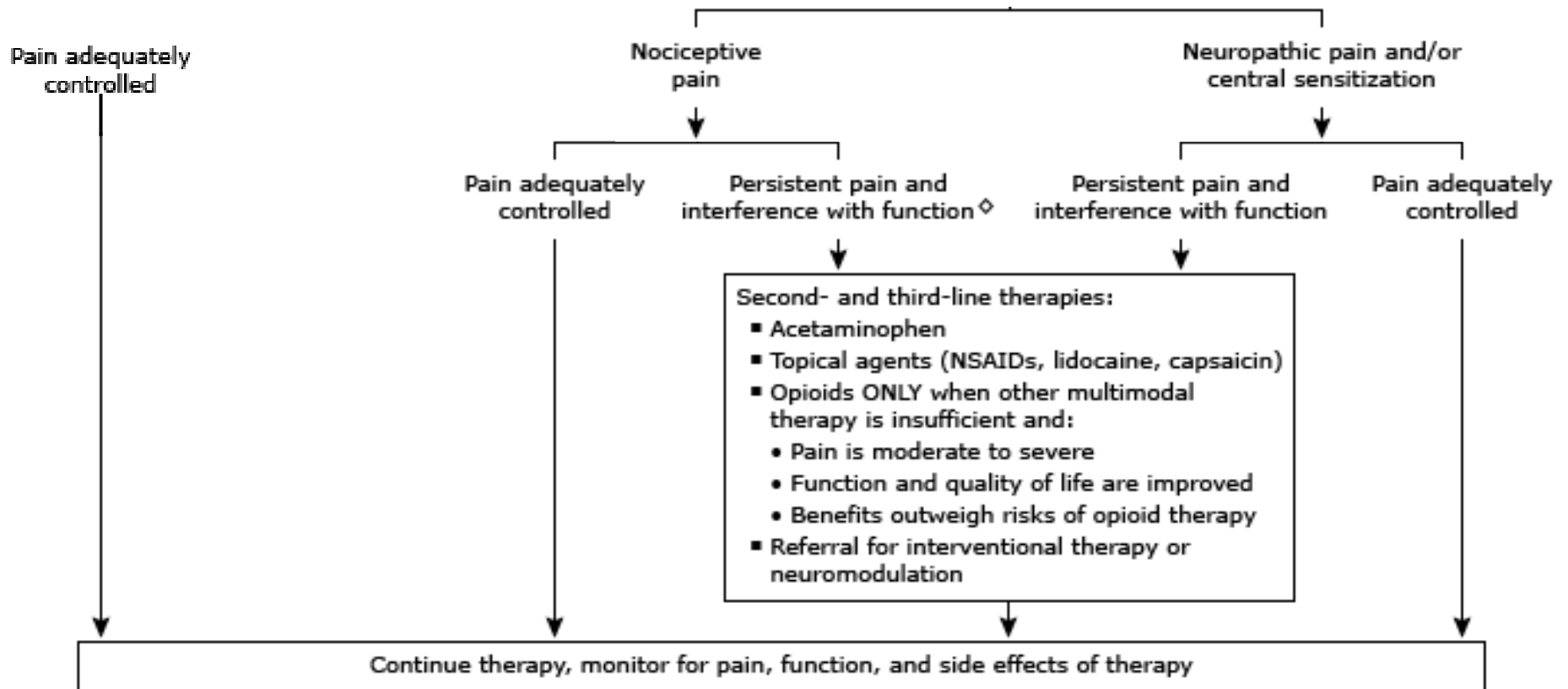




## Targeting Therapeutic Pathways









# **So What About Using Opioids - The 2022 CDC Guideline For Prescribing Opioids**

# CDC Clinic Practice Guideline for Prescribing Opioids – United States, 2022

## Summary

### This Guidelines is

- **A clinical tool to improve communication between clinicians and patients and empower them to make informed, person-centered decisions related to pain care together**
- **Intended for primary care clinicians and other clinicians providing pain care for outpatients aged ≥18 years old with:**
  - acute pain (duration <1 month);
  - subacute pain (duration of 1-3 months); or
  - chronic pain (duration of >3 months)
- **Intended to be flexible to enable person-centered decision-making, taking into account an individual’s expected health outcomes and well-being.**

### This clinical practice guideline is not

- **A replacement for clinical judgment or individualized, person-centered care**
- **Intended to be applied as inflexible standards of care across patients, and/or patient populations by healthcare professionals, health systems, pharmacies, third-party payers, or governmental jurisdictions or to lead to the rapid tapering or discontinuation of opioids for patients**
- **A law, regulation, and/or policy that dictates clinical practice or a substitute for FDA-approved labeling**
- **Applicable to the following types of pain treatment:**
  - sickle cell disease-related pain
  - cancer pain
  - palliative care
  - end-of-life care

URL - [CDC Clinical Practice Guideline for Prescribing Opioids for Pain — United States, 2022](#)  
(Accessed 5/20/23)

URL - [Prescribing Opioids for Pain — The New CDC Clinical Practice Guideline | NEJM](#) (Accessed 5/20/23)

# CDC Clinic Practice Guideline for Prescribing Opioids – United States, 2022

## Summary

### 1. For all patients with acute, subacute, or chronic pain

- Initiate the lowest dose to achieve expected results
- For opioid naïve patients, start with immediate-release opioids instead of extended-release/long-acting (ER/LA) opioids
- Use extreme caution when prescribing opioids, benzodiazepines and other sedating substances concurrently
  - consider whether benefits outweigh risks
  - Taper cautiously to a less risky dose or discontinue
- Check the state prescription drug monitoring program (PDMP) also known as the Michigan Automated Prescription Service (MAPS), to determine whether the patient is receiving opioid dosages or combinations that put the patient at high risk for overdose
  - When initiating therapy
  - Periodically when continuing
- Consider toxicology testing to assess for prescribed medications as well as other prescribed and non-prescribed controlled substances
- Offer naloxone and other overdose mitigation strategies when risk factors for opioid overdose are present

### 2. For acute pain, consider initiating opioid therapy only if benefits are anticipated to outweigh risks to the patient

- Nonopioid therapies are effective for many common types of acute pain
- Prescribe no greater quantity than needed for the expected duration of pain severe enough to require opioids

# CDC Clinic Practice Guideline for Prescribing Opioids – United States, 2022

## Summary

- 3. For subacute and chronic pain, consider initiating opioid therapy only if expected benefits for pain and function are anticipated to outweigh risks to the patient**
  - Work with patients to establish treatment goals for pain and function
  - Nonopioid therapies are preferred
  - Discuss the known risks and realistic benefits of opioid therapy
  - Consider how opioid therapy will be discontinued when benefits do not outweigh risks
  - If opioids are continued
    - use caution when prescribing opioids at any dosage
    - avoid increasing dosage above levels likely to yield diminishing returns in benefits relative to risks to patients
    - re-evaluate benefits and risks after starting opioid therapy or when escalating dose
    - Initially at 1 to 4 weeks
    - Then every 3 months (or more frequently as indicated)
- 4. Carefully weigh benefits and risks for patients already receiving higher opioid dosages**
  - Do not abruptly discontinue opioid therapy unless there are indications of a life-threatening issue, such as warning signs of impending overdose (e.g., confusion, sedation, or slurred speech)
  - Exercise care when reducing or continuing opioid dosage
  - Work closely with patients to optimize other therapies
  - Gradually taper to lower dosages if risks outweigh benefits of continued opioid therapy
  - Taper and discontinue opioids if warranted based on the individual clinical circumstances of the patient
  - Consider transitioning to buprenorphine if opioids cannot be sufficiently tapered or discontinued
- 5. Offer Medications for Opioid Use Disorder (MOUD) to patients without pain and exhibiting opioid use disorder (OUD)**

## **6. Other Considerations**

# Consider Using Buprenorphine for Pain Management

**Effective analgesia**

**Relative ceiling on respiratory depression**

Fatal overdose limited but possible with other non-opioid respiratory depressants

**Less dysphoria, sedation, constipation**

**Limited tolerance**

**Limited abuse potential and withdrawal**

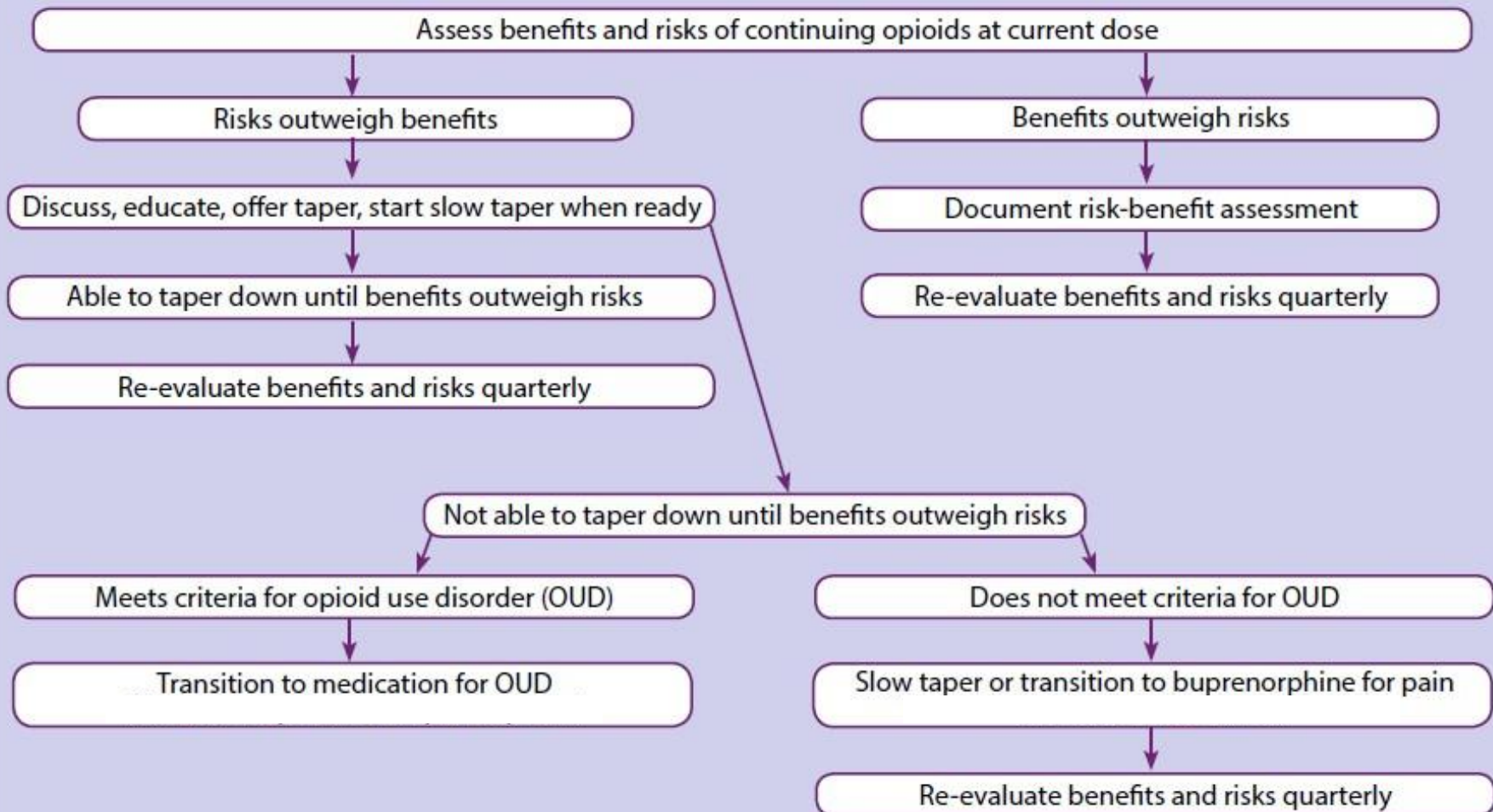
**Reduced endocrinopathies**

**Convenient dosing schedule**

**It's better than the rest.**

# When Using Opioids Long-term – Consider a Taper or Switch to Buprenorphine If and When Possible, Then Go Slow

## Opioid Tapering Flowchart



Adapted from Oregon Pain Guidance. Tapering – Guidance & Tools. Available at <https://www.oregonpainguidance.org/guideline/tapering/>.



# Don't Ignore or Abandon Inherited "Legacy" Pain Patients Already on Opioids

## Inherited Patients Taking Opioids for Chronic Pain — Considerations for Primary Care

Phillip O. Coffin, M.D., and Antje M. Barreveld, M.D.

### Steps in Caring for Patients with Chronic Pain Who Have Received Long-Term Opioid Therapy from a Previous Clinician.

- 1. Review the case with the former clinician if possible.** Try to develop a treatment plan that slowly adjusts to your style of management while avoiding a radical divergence from the previous plan of care.
- 2. Consider providing a therapeutic bridge for the patient until a plan of care is determined, given the risks associated with stopping opioid therapy.** Abruptly tapering or stopping opioid therapy can be dangerous for multiple reasons. Opioids may be crucial for the patient's condition (e.g., sickle-cell disease), and the patient may be at risk for other harms when opioids are tapered or discontinued (see figure).
- 3. Develop a patient-centered care plan.** If a taper is needed, empower the patient to make decisions, including which medications to taper first and how fast. Successful tapers may take years.
- 4. Assess the patient for opioid use disorder and start discussing medication options right away.** Patients may find it challenging to accept an opioid use disorder diagnosis; give them time.
- 5. Document opioid stewardship and the rationale for the treatment plan.** Investigations into opioid prescribing are often based on insufficient documentation.



# Orient Patients to Dispose of Expired or Unused Medications to Avoid Pilfering and Accidental Overdose

**Drug Disposal Options**  
Do you have medicine you want to get rid of?

Do you have a drug take-back option readily available?  
Check the **DEA website**, as well as your local drugstore and police station for possible options.

**NO** **YES**

Is it on the **FDA flush list**?

**NO** **YES**

Follow the FDA instructions for disposing of medicine in the household trash.

Immediately flush your medicine in the toilet. Scratch out all personal info on the bottle and recycle/throw it away.

Take your medicine to a drug take-back location.  
Do this promptly for FDA flush list drugs!

**FDA U.S. FOOD & DRUG ADMINISTRATION**

A single opportunity is all it takes. **Remove the risk** of unused opioids.

[www.fda.gov/DrugDisposal](http://www.fda.gov/DrugDisposal)

**FDA Remove the RISK**

**FDA Remove the RISK**

They're both relying on you to **remove the risk** of unused opioid medicines.

[www.fda.gov/DrugDisposal](http://www.fda.gov/DrugDisposal)

**FDA Remove the RISK**

# Offer Narcan (Naloxone) or Opvee (Nalmefene) to Reverse Overdose

1. Opvee contains nalmefene as its active ingredient, while Narcan contains naloxone.
2. Opvee and Narcan are both effective opioid overdose reversal medications.
3. Narcan can wear off before the opioid does.
4. Opvee's half-life is about 11 hours and Narcan's half-life is about 2 hours.
5. This means Opvee stays in the body longer — a potential advantage for reversing an overdose from synthetic opioids like fentanyl that can last longer in the body.
6. Opvee may require less doses than Narcan for the same amount of fentanyl ingested by the person overdosing.
7. Opvee restored breathing about 2 times better than Narcan within 5 minutes of administering the nasal spray.



# Summary



**This program was intended to optimize clinical practice by helping the clinician better understand:**

1. How to use multi-modal pain treatment approaches to avoid excessive and prolonged doses of opioids
2. Recently updated definitions for pain
3. Pain signaling pathways and therapeutic targets
4. The new 2022 CDC Guidelines for Using Opioids
5. Principles of assessment and treatment for acute and chronic pain
6. When to consider using buprenorphine for chronic pain
7. Additional considerations for pain management



Thank You!