



36th Annual

JOSE I. RICARD, MD

FAMILY MEDICINE & SPORTS MEDICINE CONFERENCE



Presented by the Family Medicine Foundation of West Virginia



Drug Diversion Training and Best Practice Prescribing of Controlled Substances

Tracy Hendershot, MD, FAAFP
WV Family Medicine Foundation
Huntington, WV
Thursday November 10th, 2022

Credentials

Dr. Hendershot, MD, DC, FAAFP is a past Paul Ambrose Health Policy Fellow. He trained at Marshall University's Joan C. Edwards School of Medicine with completion of a family practice residency at the same.

He's worked in private practice as a chiropractor from 1996-2004, observing the WV opioid crisis develop from the vantage point of a non-prescribing provider. Since becoming an MD he's worked at the Ebenezer Clinic- a free clinic blocks from Huntington, WV's initial opioid epicenter. He's been CMO of a rural FQHC and Past Chair of the WV PCA CMO committee. He's now employed in the WVU Medicine Health System. At each location he's been handed his share of chronic opioid patients. He manages < 30 chronic opioid patients in the outpatient setting.

Finally, Dr. Hendershot has served as Past President of the WVAFP, currently serving as WVAFP Delegate to the AAFP Congress of Delegates. He also serves as chair of the WVAFP Legislative Committee.

Disclaimers

Dr. Hendershot has **no** conflicts of interest or disclaimers to announce.

The use of brand specific names are not meant as an endorsement,
But to ensure familiarity of the prescriber with the common opioid
products.

I receive no remuneration from any manufacturer.

Objectives

1. Review the climate and trends in WV that contribute to opioid overdose deaths.
2. Review best practice prescribing of controlled substances.
3. Review drug diversion concerns and best practices.
4. Encourage the appropriate use of Naloxone and MAT.

Agenda

The First 30 Hour:

Why We Are here

The current situation

Assessment of Risk

Assessment of Need

BOP

Contracts

The Hour After That:

Prescribing Opioids

Opioid equivalents

Testing:

Urine

Pill counts

The Last Hour:

The New CDC
Guidelines

The WV Opioid
Reduction Act

Why We Are Here...its required

Mandatory Controlled Substance CME for all Licensees

(SB 437 passed 2012)

“Physicians who have prescribed, administered, or dispensed any controlled substance in any jurisdiction in the two year license cycle preceding renewal, are required to complete three hours of Board-approved CME in drug diversion and best practices prescribing of controlled substances **during each reporting period. This is not a one-time only requirement.**

A physician who has **not prescribed any controlled substances whatsoever during the reporting period may seek a waiver** of this requirement by attesting on the renewal application that he or she has not prescribed, administered or dispensed any controlled substances whatsoever since July 1, 2016.”

Why We Are Here...its a state objective

The screenshot shows a web browser window with the URL <https://dhhr.wv.gov/office-of-drug-control-policy/news/Pages/Reports-and-Data.aspx>. The page content includes a sidebar with 'WV.gov' and 'Health Resources' logos, a main header 'Goal 2: Monitor opioid prescriptions and distribution', and a 'Guidelines' section with a link to 'WV Pain Management Guidelines'. A table titled 'Expert Pain Management Panel Members' is displayed, listing 18 members and their organizations. A red circle highlights the table and the 'Guidelines' section. A separate box on the right contains the title 'West Virginia Expert Pain Management Panel Safe & Effective Management of Pain Guidelines 2016' and the page number '1'.

Panel Member	Org
Mark Garofoli, PharmD, MBA (Coordinator)	West Virginia University (WVU)
Timothy Deer, MD (Chairperson)	Centers for Pain Relief
Richard Vaglianti, MD (Vice Chairperson)	WVU Pain Management
Rahul Gupta, MD	West Virginia DHHR, Public Health
Ahmet Ozturk, MD	Marshall University
Denzil Hawkinberry, MD	Community Care Centers
Bradley Hall, MD	WV Medical Professionals Hospital
Matt Cupp, MD	Board Certified
Michael Mills, DO	West Virginia Office of Health
Jimmy Adams, DO	Active Physician
Richard Gross, PhD	WVU Pain Management
Jason Roush, DDS	West Virginia
Stacey Wyatt, RN	St. Francis
Vicki Cunningham, RPh	WV Bureau of Medical
Felice Joseph, RPh	PEIA
Stephen Small, RPh, MS	Rational Drug
Patty Johnston, RPh	Colony Drug & Wellne
Charles Ponte, PharmD, CPE	WVU School
James Jeffries, MS	WV DHHR, Division of Inf
Michael Goff	West Virginia Prescription

West Virginia Expert Pain Management Panel
Safe & Effective Management of Pain Guidelines
2016

1

WVBM and WVBOM Approved Courses:

The Boards of Medicine maintain a list of all three-hour courses that have been approved..

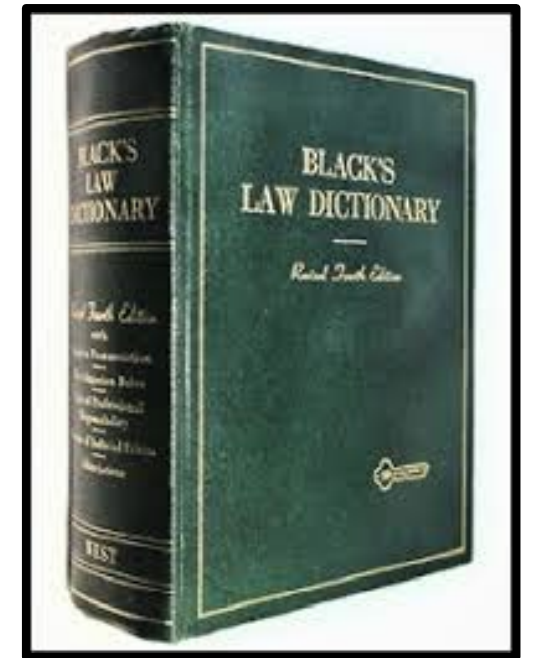
-Thank you for attending **this** lecture

The screenshot displays the website for the West Virginia Board of Osteopathic Medicine. The main navigation bar includes 'Legislative Rules', 'Calendar', 'News', 'About', 'IMLCC', and 'Contact'. A secondary navigation bar features 'Licensing', 'Verification', and 'File Complaint'. The page title is 'Licensure' and the breadcrumb trail is 'Home / Continuing Education / 2022 Renewal CME Courses'. A sidebar on the left lists 'Licensing Options' for 'OSTEOPATHIC PHYSICIANS', including 'Application', 'Emergency Temporary Permit', 'Application Status', 'Renewal', 'Print Licensure Card', 'CSL Application', 'CSL Renewal', 'CSL Print Licensure Card', and 'CME'. The main content area is titled '2022 Renewal CME Courses' and includes a description: 'List of Board-approved CME courses which satisfy the 3-hour Drug Diversion Training and Best Practice Prescribing of Controlled Substances Training requirement for medical doctors whose last names begin with the letters A through L, and who will be renewing on or before June 30, 2022.' Below this is a table with three columns: 'Course Name', 'Sponsor', and 'Location / Date'.

Course Name	Sponsor	Location / Date
Pain & Addiction, Best Practices & Proper Prescribing: Changing a Culture by Changing the Culture of Medicine	WVU School of Medicine and WV Medical Professionals Health Program	ONLINE COURSE # Expires 01/01/2022
From Prescription Drug Abuse to Street Heroin...The Tale of West Virginia's Drug Abuse Epidemic	CAMC Health Education and Research Institute	ONLINE COURSE #
Prescribing Opioids, Providing Naloxone, and Preventing Drug Diversion: The West Virginia Requirement, #91601 or #91602	NetCe	ONLINE COURSE # Expires 03/19/2022

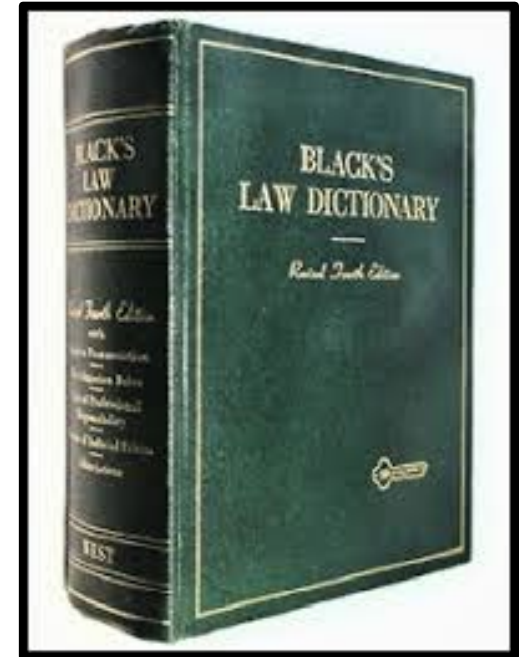
Common Terms:

- **Opiates:** refer to **natural** opioids such as heroin, morphine, and codeine.
- **Opioids:** refers to **all** natural, semisynthetic (hydrocodone, oxycodone, hydromorphone..), and synthetic opioids (excludes methadone, includes tramadol and fentanyl)
- **MAT:** Medication assisted treatment for opioid use disorder when combined with counseling and behavioral therapies.
- **MME:** Morphine milligram equivalents, accounts for different drug types and strengths.



Common Terms:

- **Illicit drugs:** drugs prohibited by law or illicitly manufactured drugs, i.e. fentanyl, ecstasy.
- **Drug Misuse:** The use of drugs in a manner other than prescribed by a doctor.
- **Tolerance:** Reduced response to a drug with repeated use.
- **Dependence:** adaption to a drug that produces symptoms of withdrawal when drug is stopped.
- **Drug addiction:** Preferred term is **Substance Use Disorder**, a problematic pattern of opioid use that causes significant impairment or distress.
 - Unsuccessful efforts to reduce.
 - Use resulting in personal, social, and/or work problems

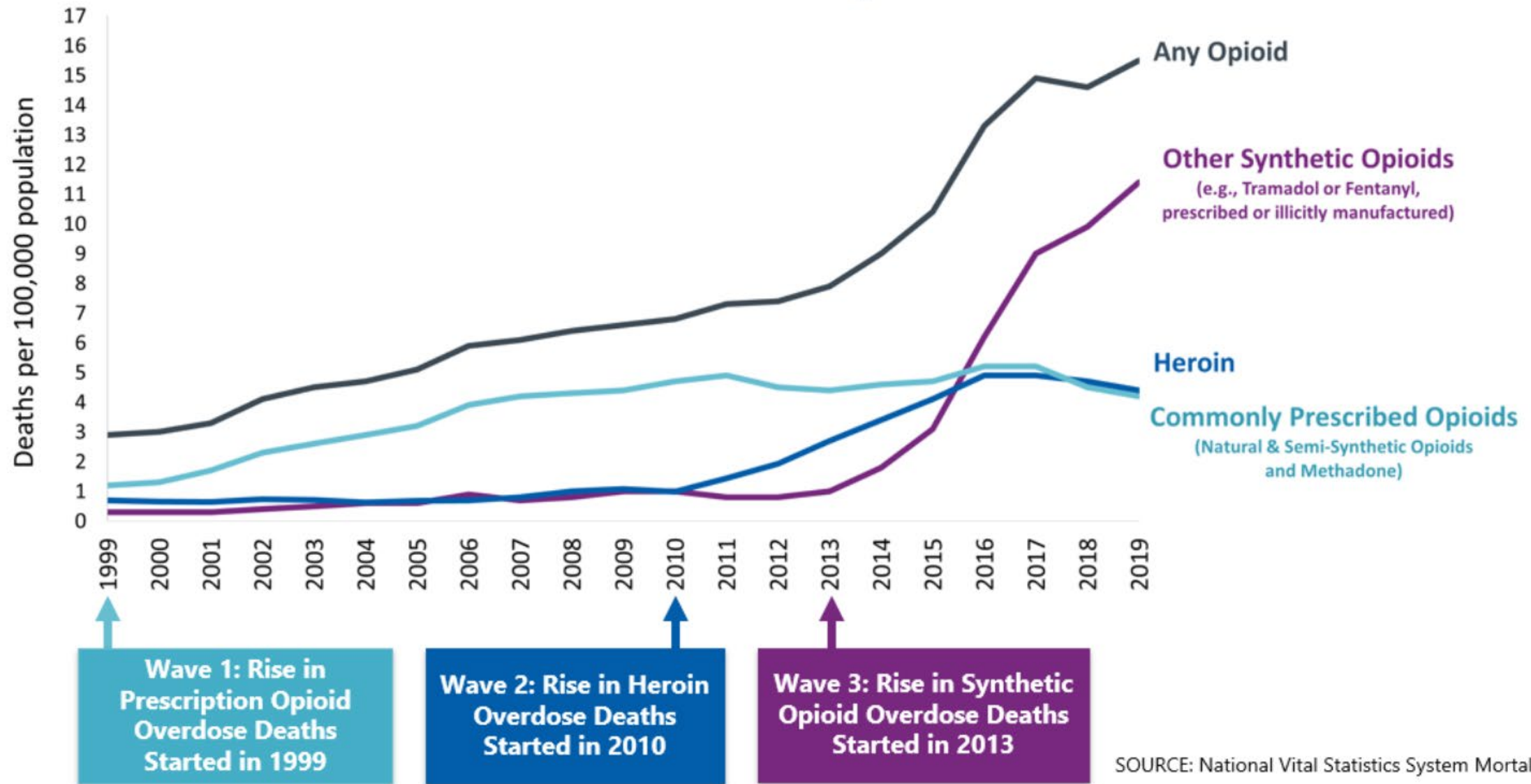


The Opioid Crisis

Nearly **841,000 people have died since 1999** from a drug overdose.
In 2019, **70,630** drug overdose deaths occurred in the United States.
The age-adjusted rate of overdose deaths **increased by over 4%** from
2018 (20.7 per 100,000) to
2019 (21.6 per 100,000).



Three Waves of the Rise in Opioid Overdose Deaths



SOURCE: National Vital Statistics System Mortality File.

Current National Trends

The COVID-19 pandemic worsened the Opioid Crisis.

While there was a **4.6% drop** from 2017 (21.7 per 100,000) to 2018 (20.7 per 100,000).

Recent provisional data available from the CDC indicated that approximately 81,230 drug overdose deaths occurred in the United States in the 12-months ending in May 2020.

This represents a worsening of the drug overdose epidemic in the United States and is the largest number of drug overdoses for a 12-month period ever recorded.

Synthetic opioids (other than methadone)—remain the main driver of drug overdose deaths.

67.0% of opioid-involved overdose deaths involve synthetic opioids.

Current National Trends

In 2020, WV lead the nation with **the highest rates of drug overdose deaths.**

West Virginia (81.4 per 100,000 or 1330 persons),

District Of Columbia (58.1 per 100,000 or 424 persons),

Kentucky (49.2 per 100,000 or 2083 persons),

Delaware (47.3 per 100,000 or 444 persons),

Ohio (47.2 per 100,000 or 5204 persons),

Tennessee (45.6 per 100,000 or 3034 persons), and

Maryland (44.6 per 100,000 or 2771 persons).

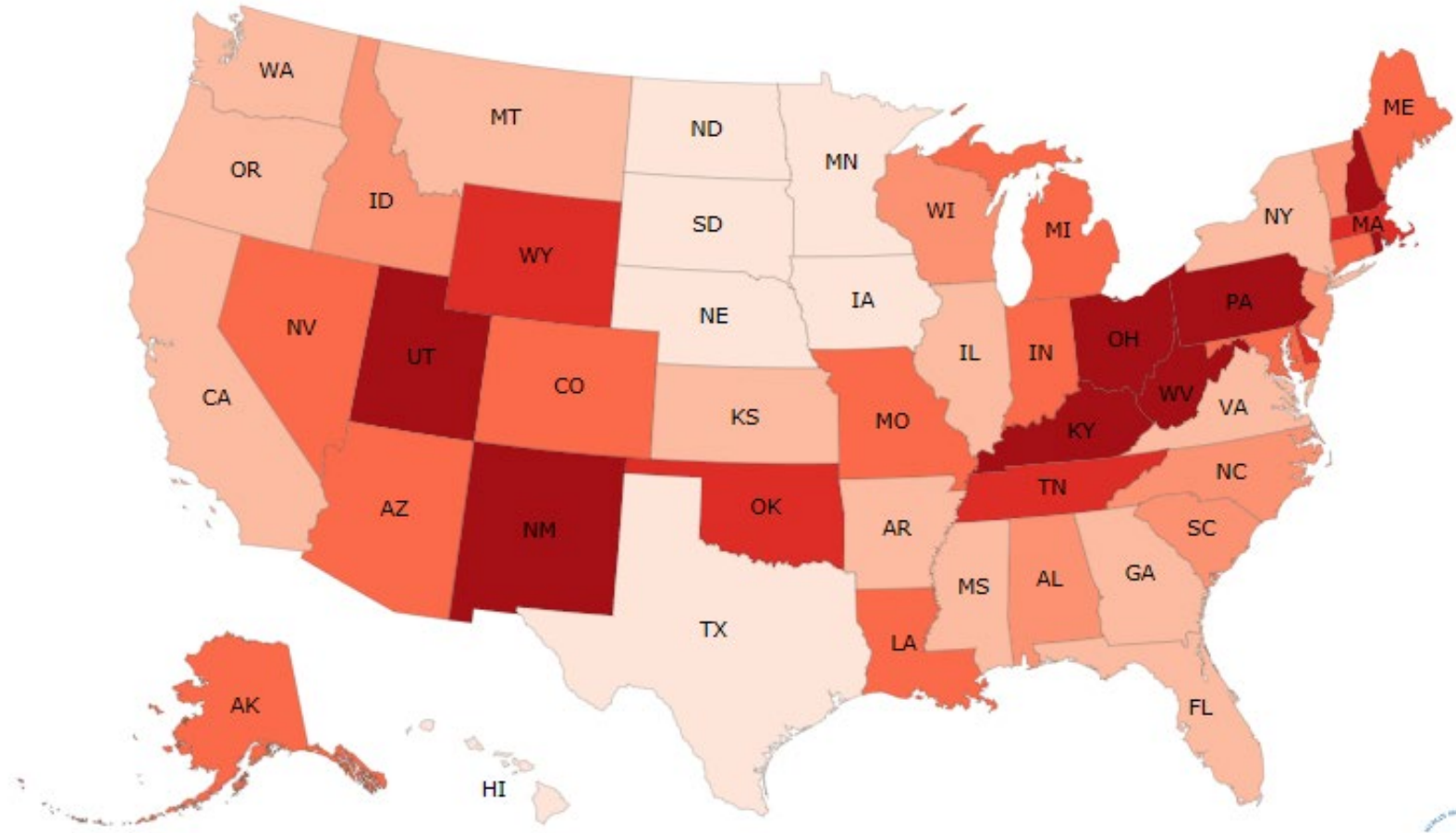
California and Florida lead actual losses with 8908 and 7231 respectively

Number and age-adjusted rates of drug overdose deaths by state, US 2014

West Virginia

35.5 per 100,000

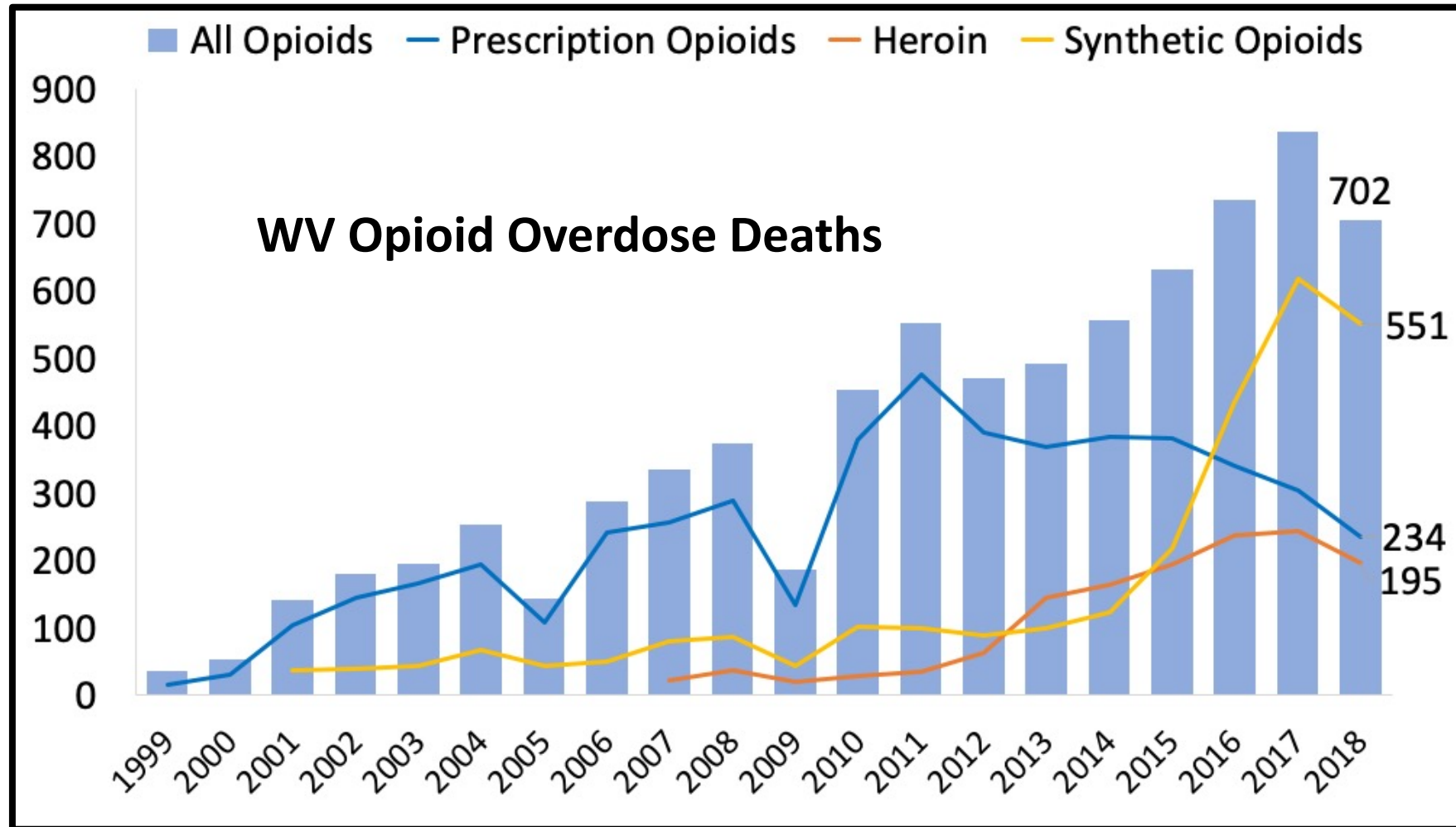
627 total deaths



- CT
- DC
- DE
- MD
- NH
- NJ
- RI
- VT

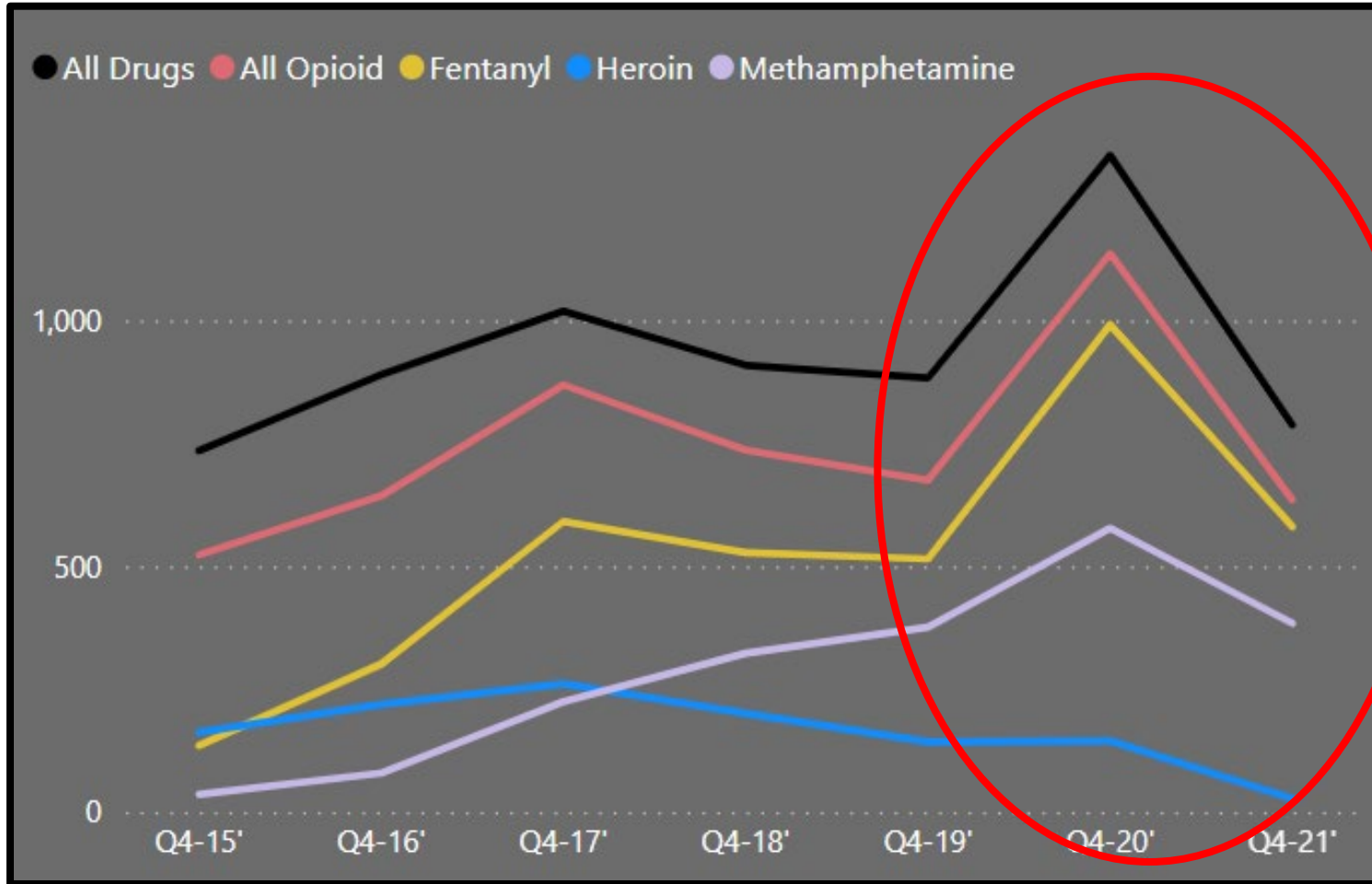


Pre-COVID WV Trends



..WV follows national picture.

Recent WV Overdose Trends

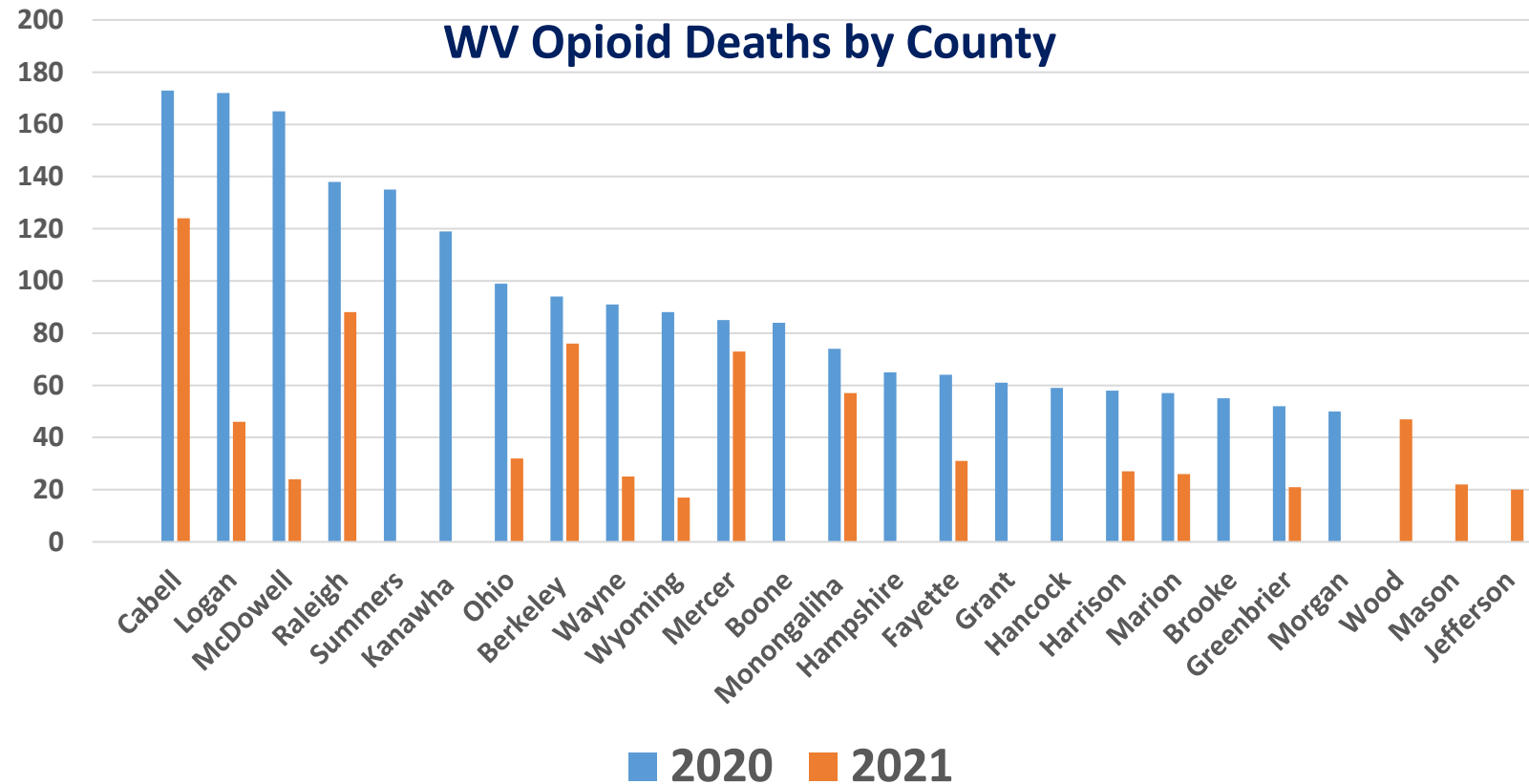


CDC data has lagged; and,

State data may not be directly comparable.

Regardless, **The COVID crisis resulted in an increase in opioid overdosing.**

WV Opioid County Deaths

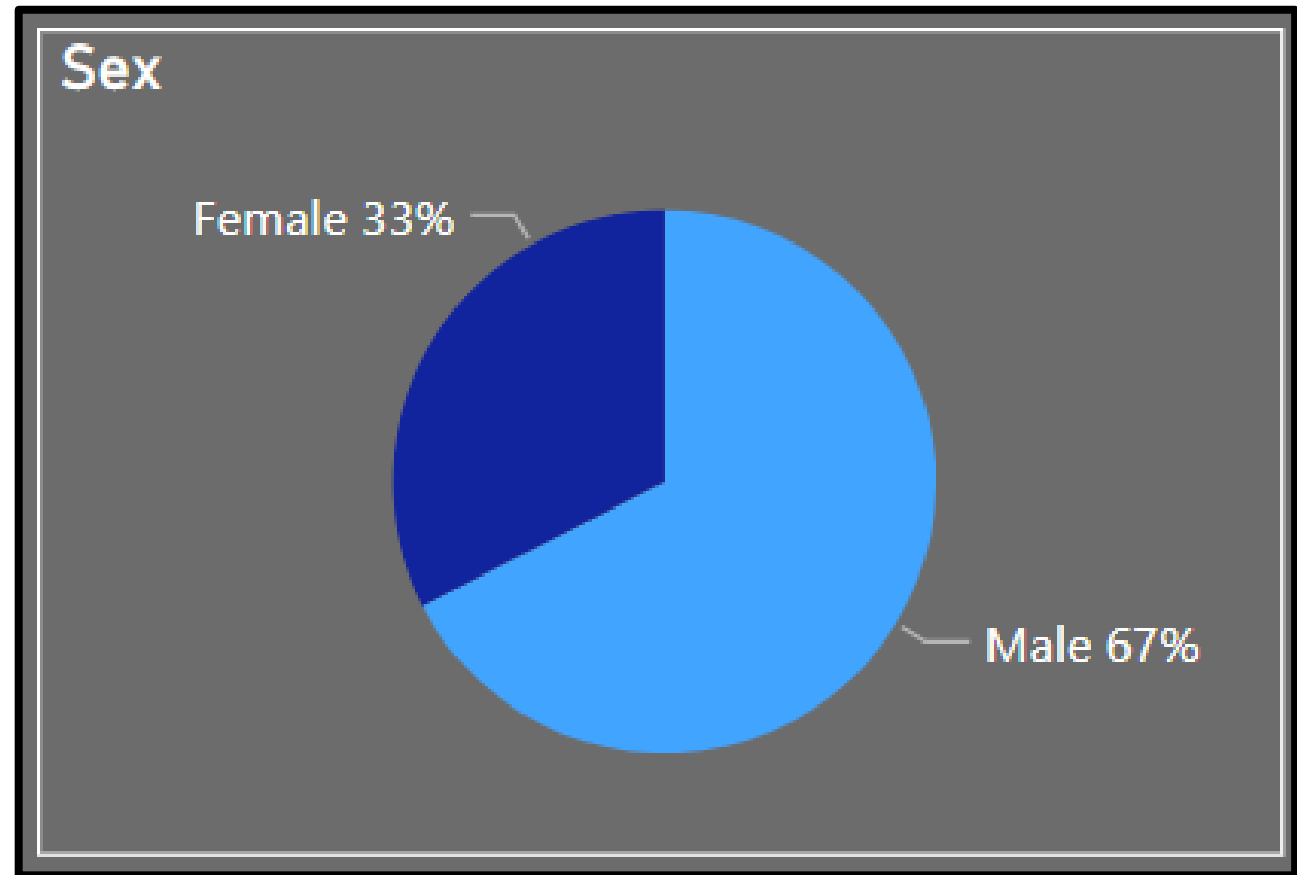
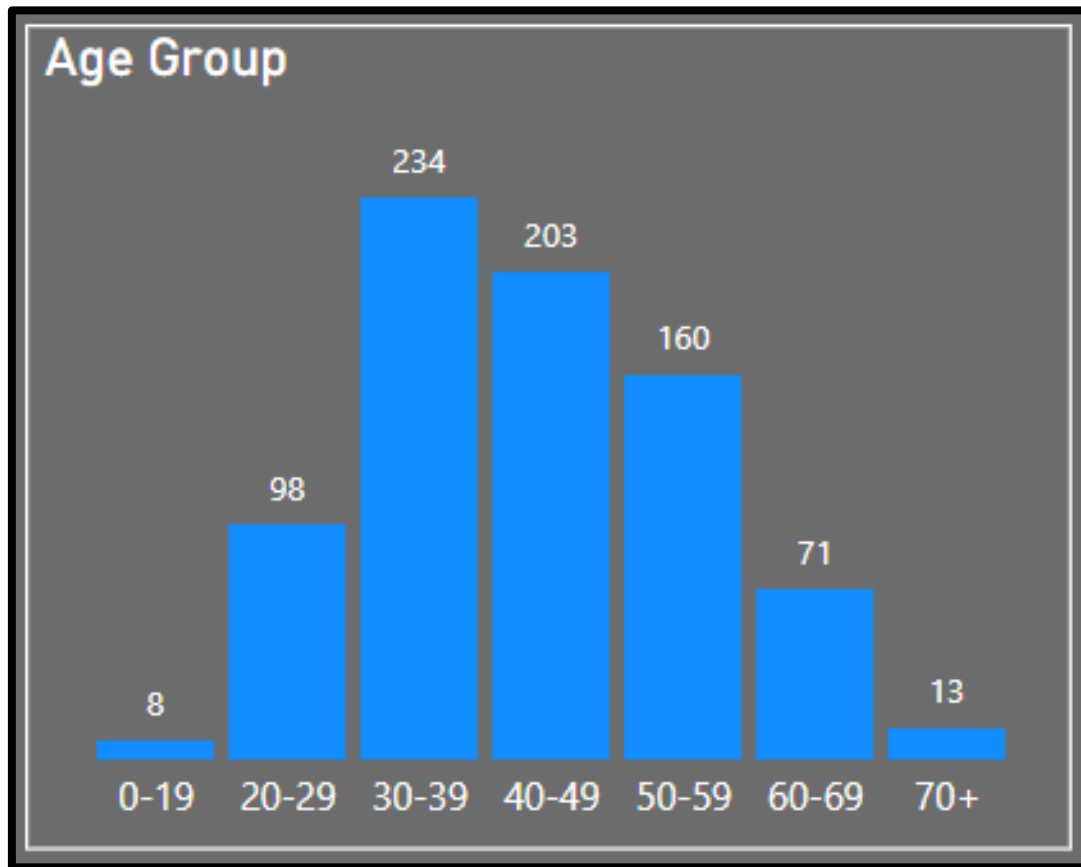


Not CDC Data.

2021 data is incomplete as pending deaths have not been confirmed.

WV Demographics 2021

WV Fatal Overdoses by age and sex



Fentanyl is Leading Opioid Deaths

It continues to be **illegal opioid** deaths, **not prescription opioids**, that drive the current national epidemic.

- 72.9% of overdose deaths are from synthetic opioids.
- Deaths involving psychostimulants such as **cocaine and methamphetamine are increasing** with and without synthetic opioid involvement.

Drug overdose deaths have shifted geographically.

- From 2018 to 2019, the largest increase in death rates involving synthetic opioids occurred in the West (67.9%).
- The largest increase in death rates involving psychostimulants occurred in the Northeast (43.8%).

Current National Trends

The overall national opioid prescribing rate has declined from 2012 to 2020.

In 2012 the national rate was **81.3** prescriptions per 100 persons.
In 2020 the rate had fallen to **43.3** prescriptions per 100 persons.

However,
the prescribing rates continued to remain high in a
few counties across the country.

In 3.6% of U.S. counties the rate is still 100:100

(this is a drop from 11% in 2018)

WV Trends?

In 2020, West Virginia providers wrote:

53.7 opioid prescriptions for every 100 persons,

(**The 13th highest rate in the U.S.** that year)

Positive news?

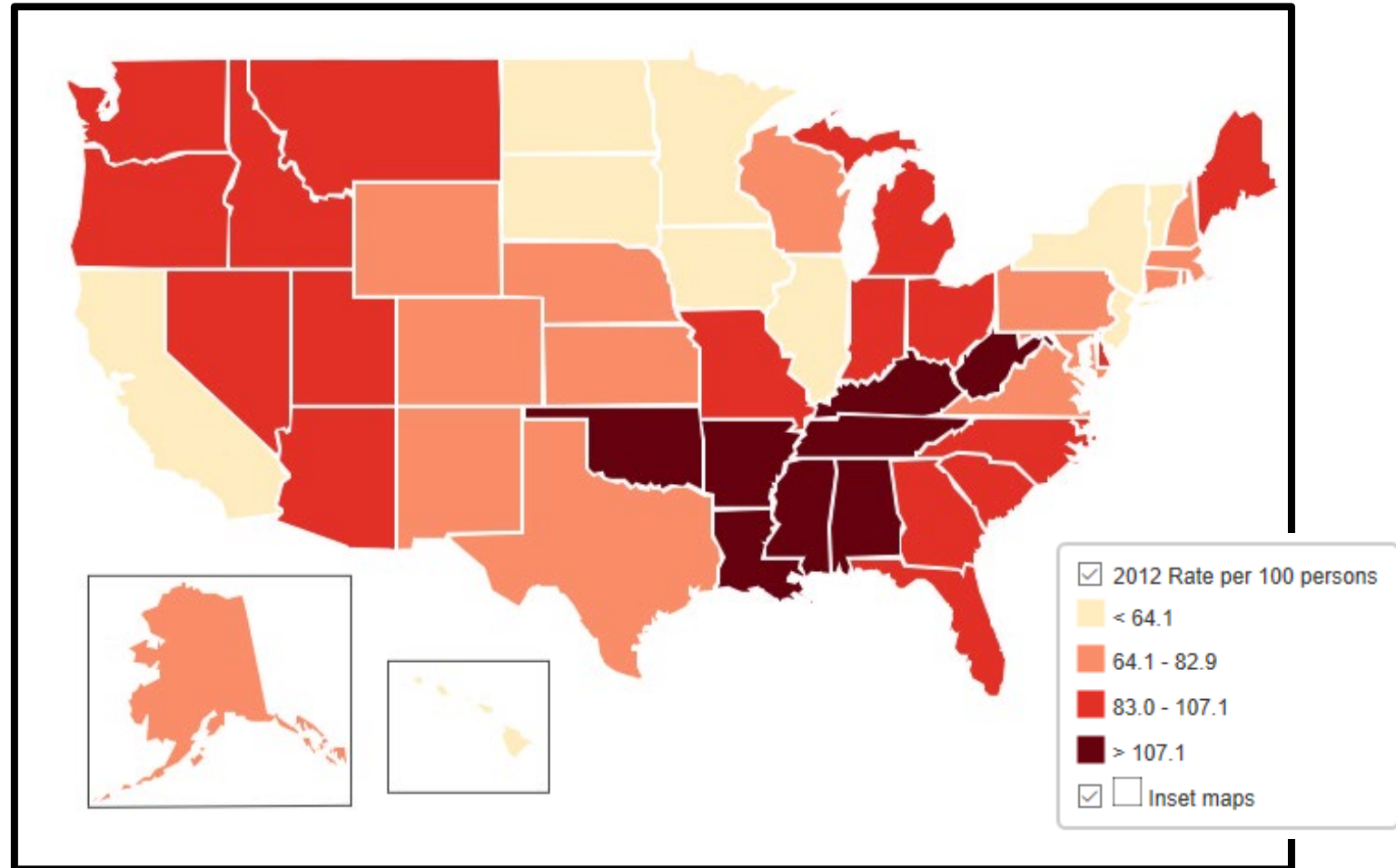
**This was the lowest WV rate since data became available in
2006**



U.S. State Prescribing Rates, 2009

West Virginia

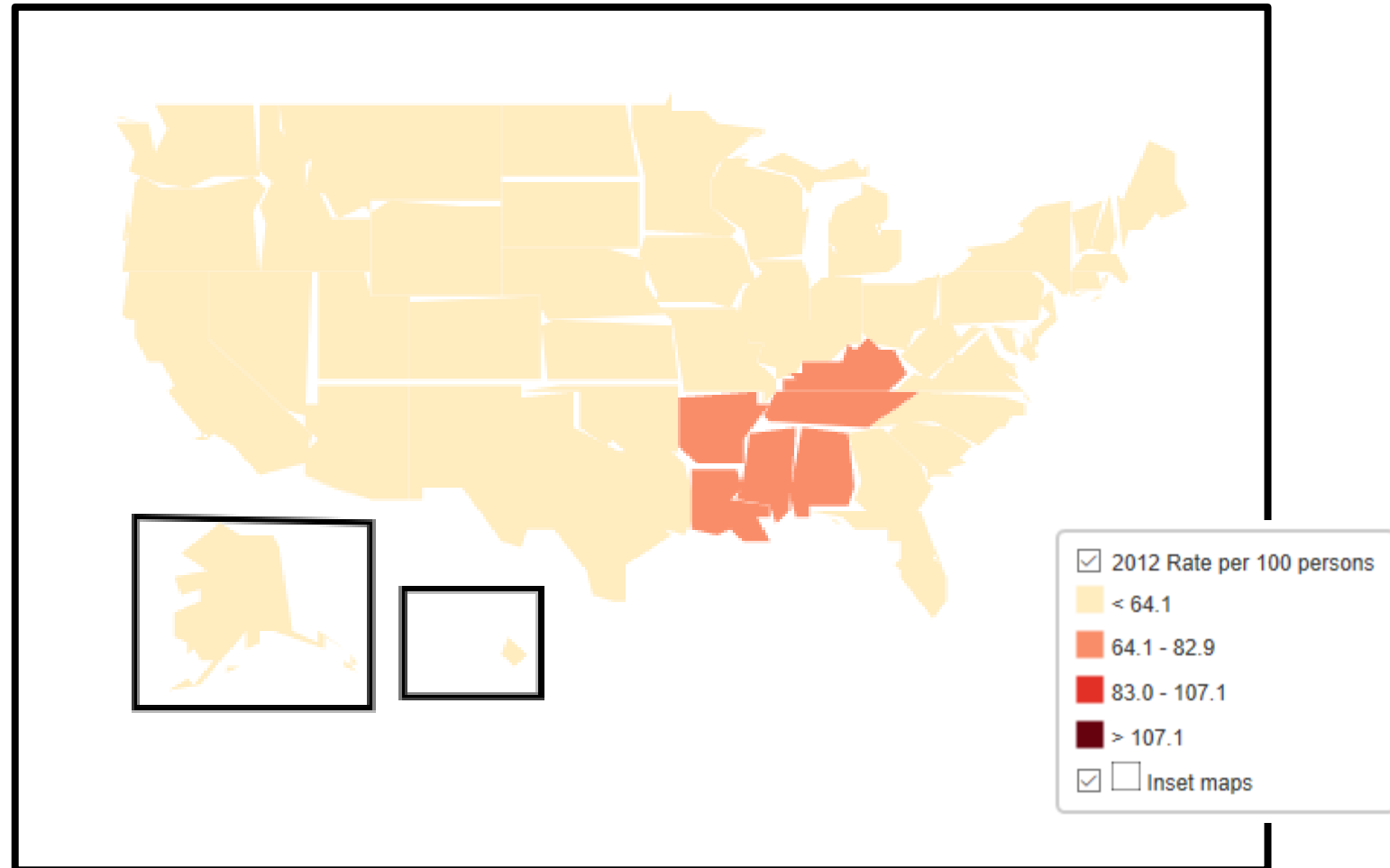
146.9



U.S. State Prescribing Rates, 2020

West Virginia

53.7



WV County Rx Data

By 2009 individual WV county physicians began to respond to the crisis.

At that time, **Mingo county** lead the state in Rx rates-With **nearly triple** the rate of other high prescribing counties.

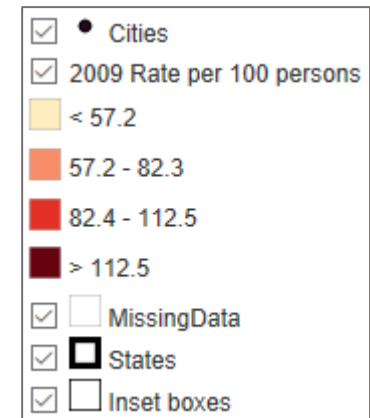
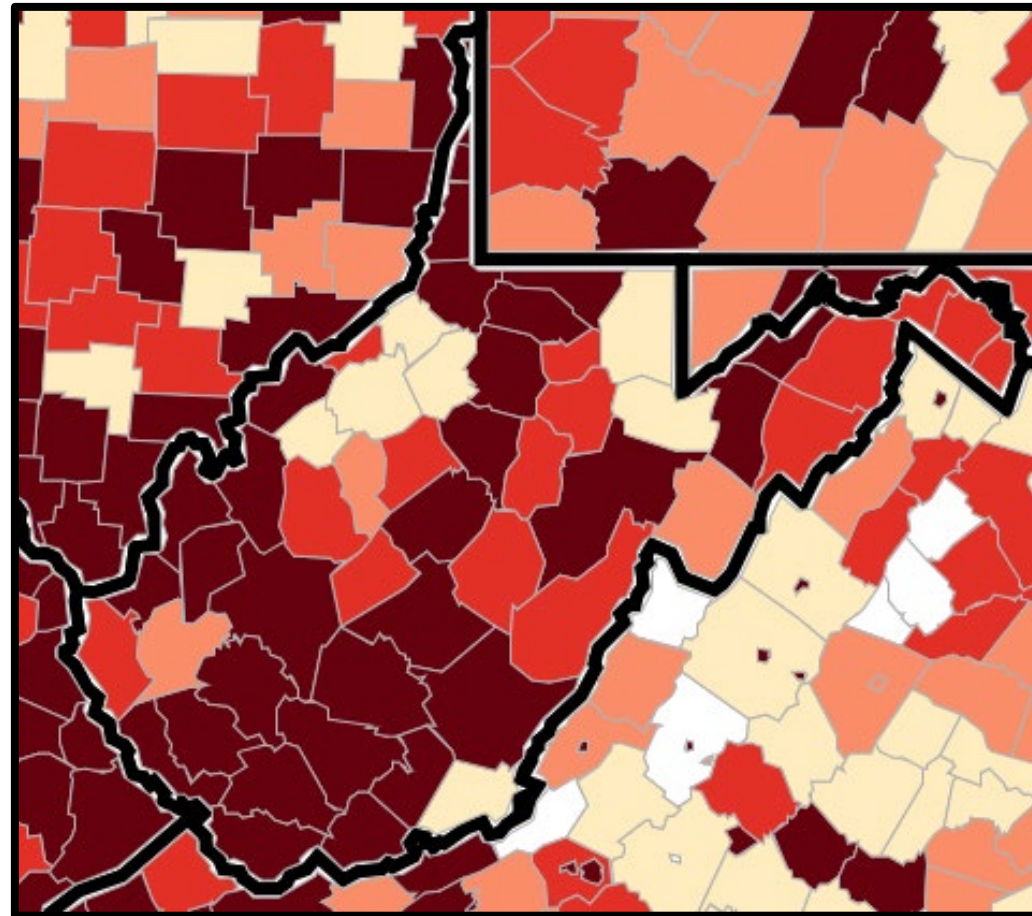
This abruptly stopped by 2010 and **Logan county** alone drove the WV outlier status.

This because it has a larger population than other outliers and because its rate is triple the other outliers. (36,000 pop and 15th in WV)

WV County Prescribing Rates, 2009

Mingo, WV

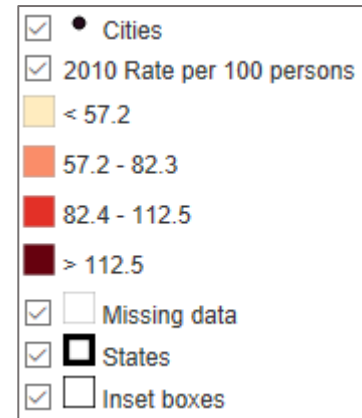
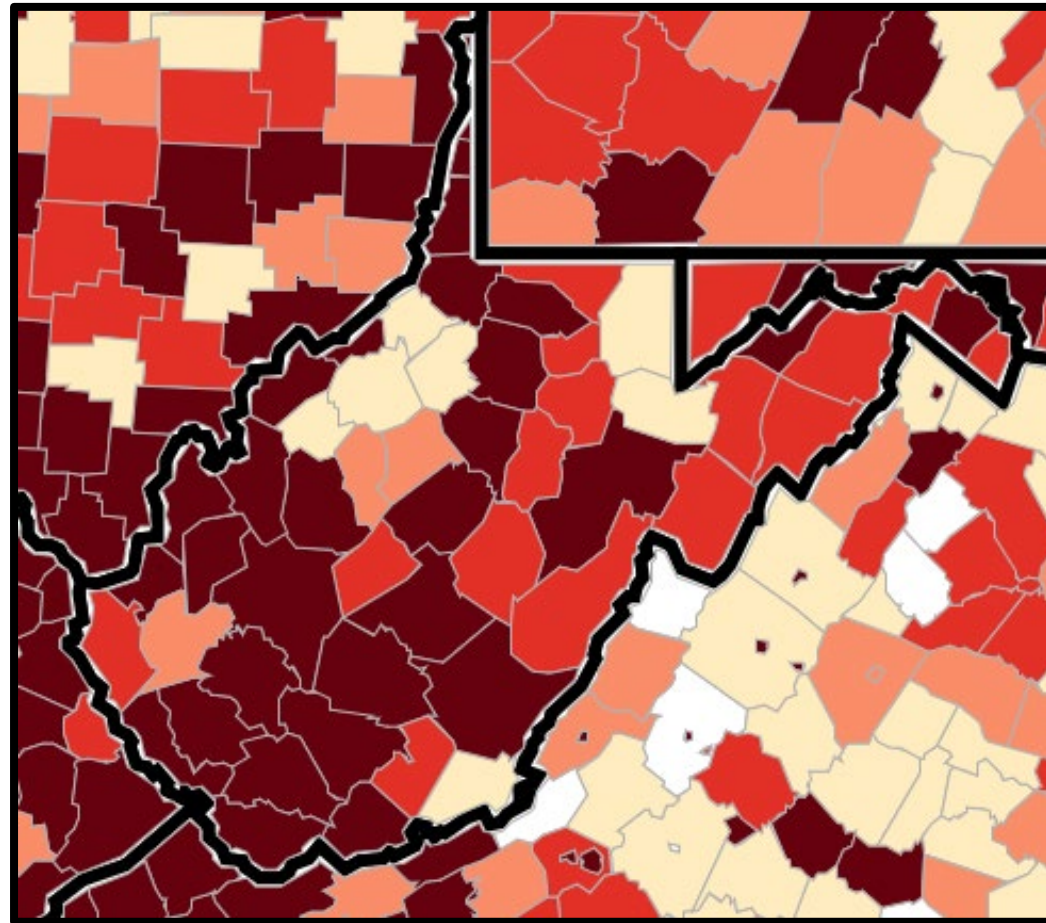
437.2



WV County Prescribing Rates, 2010

Logan, WV

287.4



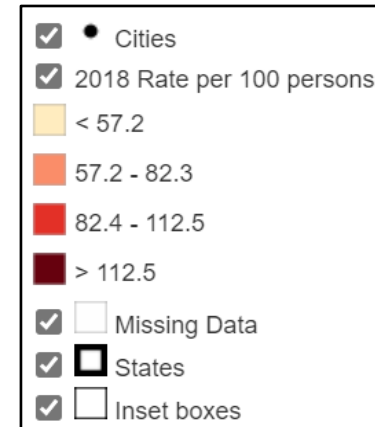
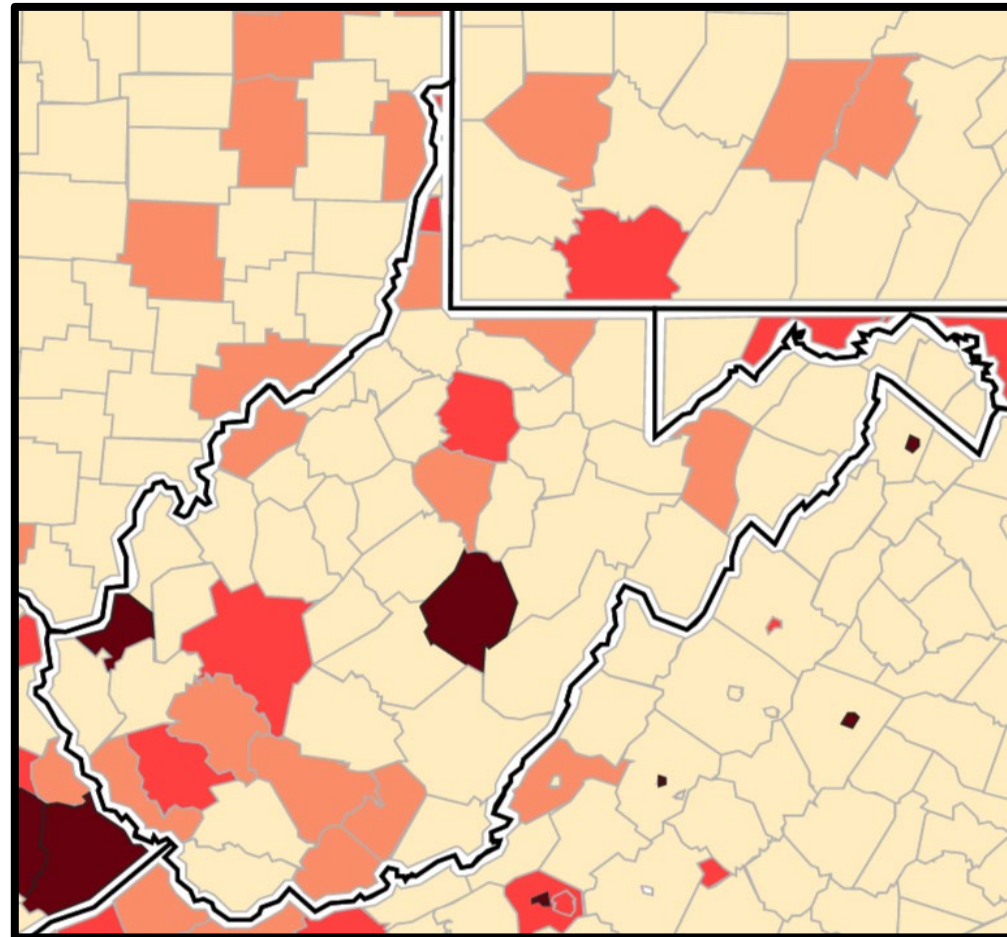
WV County Prescribing Rates, 2019

Logan, WV

93.7

Webster, WV

124.8



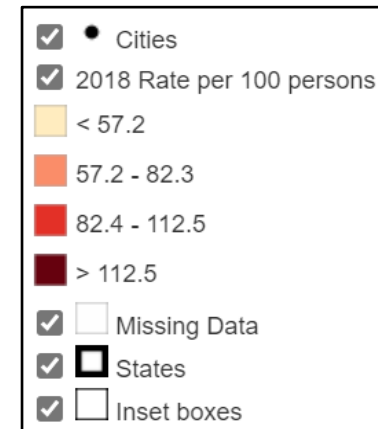
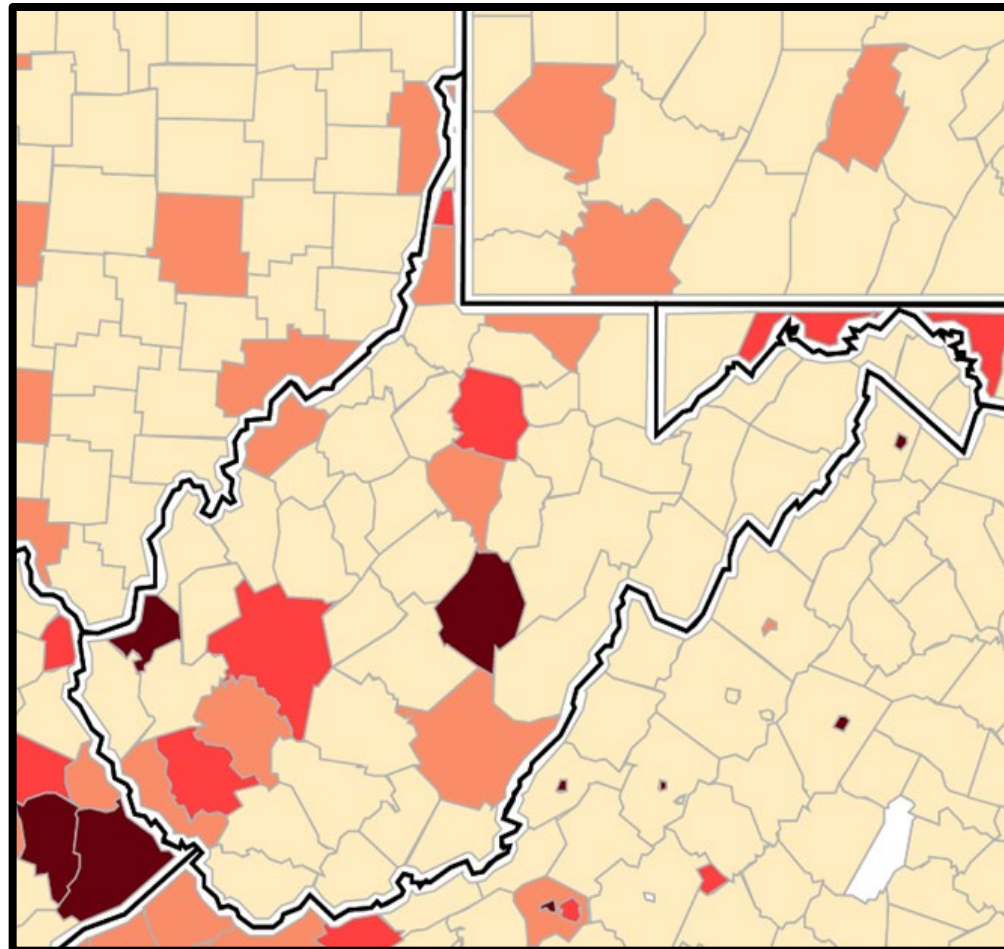
WV County Prescribing Rates, 2020

Logan, WV

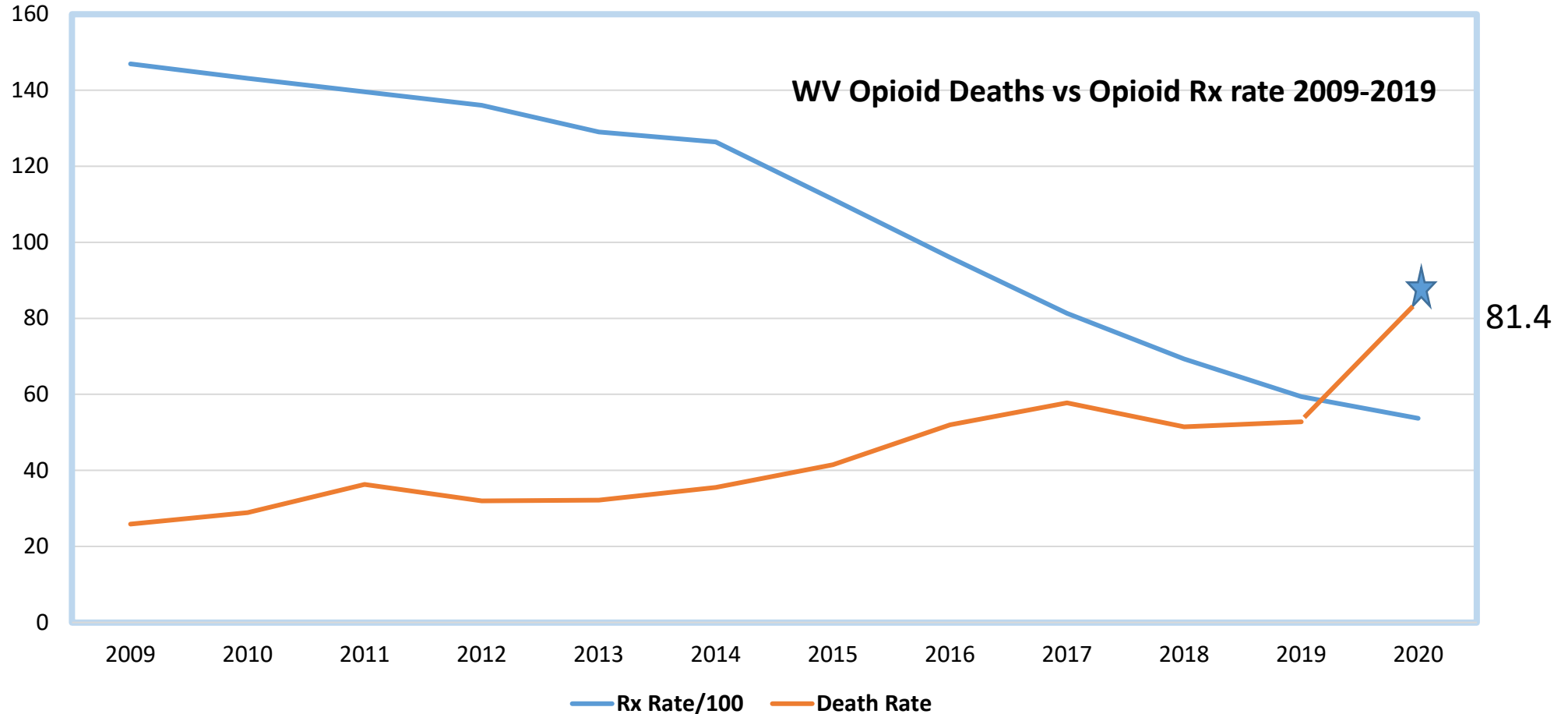
85.7

Webster, WV

118

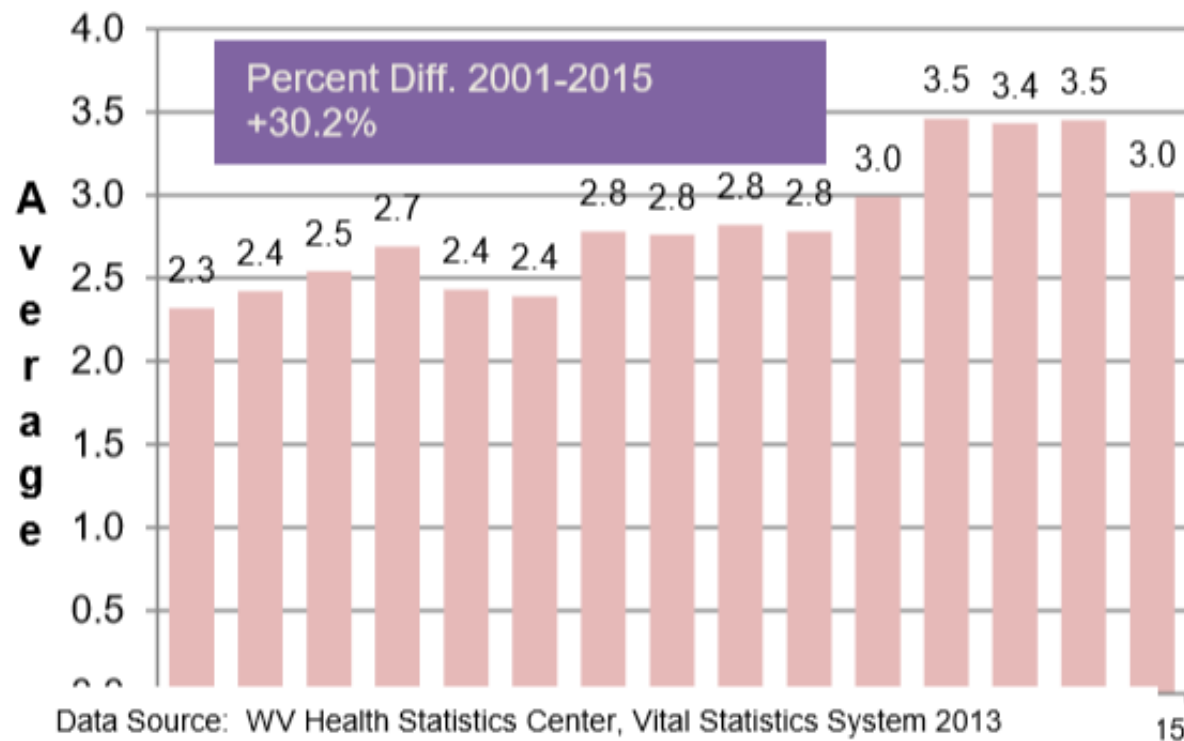


WV Opioid Reduction Not Tied to Less Deaths



Illegal and Rx Opioids Often Mixed:

Figure 2: Average Number of Drugs Involved Per Fatal Overdose West Virginia Occurrences, 2001-2015



This is a primary rationale for appropriate, ongoing, prescribing education as opposed to ongoing education regarding Addiction Treatment.

A new, good, reason for ongoing training is less physician experience with chronic opioid management.

Break



Starting a Patient on Opioids



Clinical Practice Guideline
Opioid Prescribing for Chronic Pain
Affirmation of Value, **April 2016**

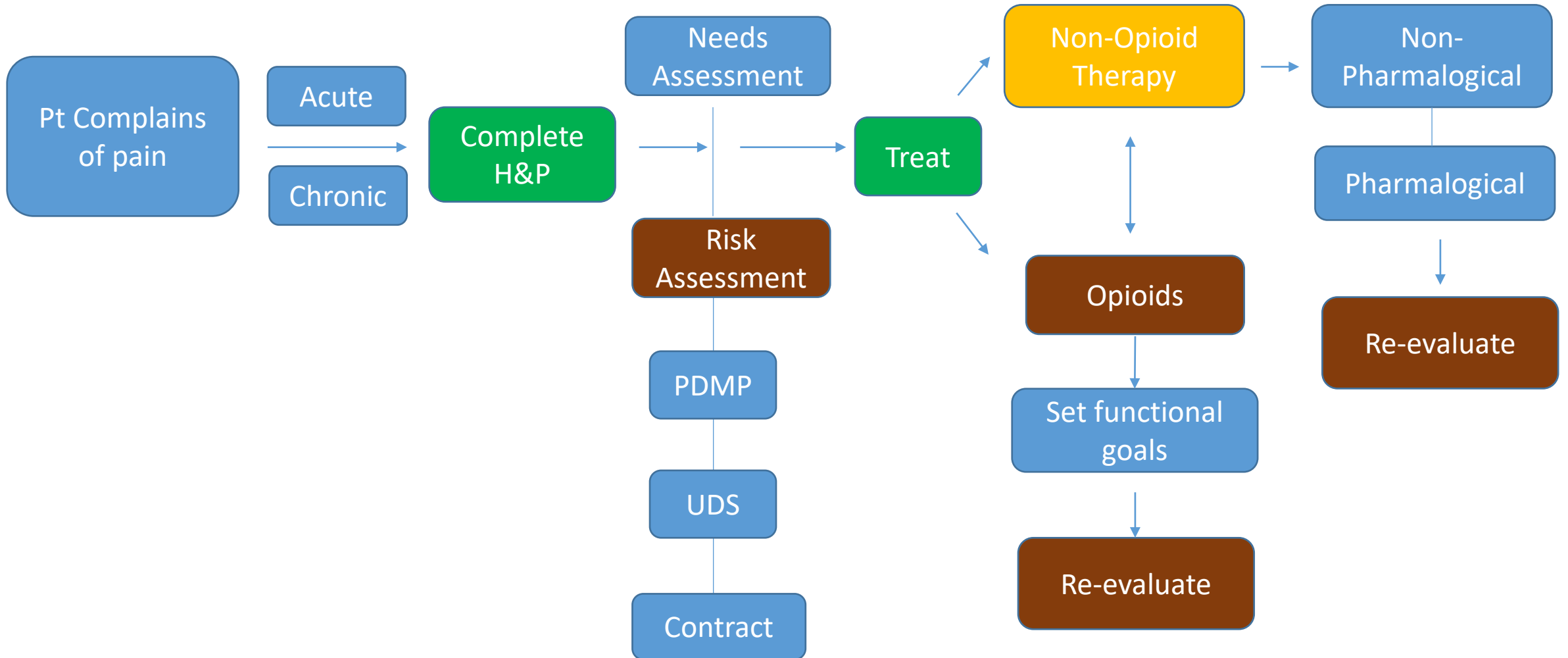
CDC Guideline for Prescribing Opioids for Chronic Pain — United States, **2016** Recommendations and Reports / March 18, 2016 / 65(1);1–49

Dowell D, Ragan KR, Jones CM, Baldwin GT, Chou R. CDC Clinical Practice Guideline for Prescribing Opioids for Pain — United States, 2022. MMWR Recomm Rep 2022;71(No. RR-3):1–95. DOI: <http://dx.doi.org/10.15585/mmwr.rr7103a1>.



Responsible, safe, and effective prescription of opioids for Chronic Non-Cancer Pain: ASIPP Guidelines. Pain physician 2017; 2017; 20;S3-S92. Year updated **2017**.

Opioid Prescribing Flow:



Assessment of Risk-ORT

Indicates the **probability of opioid-related aberrant behaviors**

In a study, 158 consecutive new patients treated in a pain clinic took the **Opioid Risk Tool** (ORT). It measured valid risk factors associated with substance abuse. All patients were monitored for aberrant behaviors for 12 months after their initial visits.

The ORT displayed **excellent discrimination** for both the male ($c = 0.82$) and the female ($c = 0.85$) prognostic models.

Assessment of Risk-ORT

Those Scored as..

- **Low Risk: scores of 0-3**, 17 out of 18 (94.4%) did **not** display an aberrant behavior.
- **Moderate risk: Score of 4-7**
- **High Risk: Score of > or = to 8**, 40 out of 44 (90.9%) **did** display an aberrant behavior.

Mark each box that applies	Female	Male
Family history of substance abuse		
Alcohol	1	3
Illegal drugs	2	3
Rx drugs	4	4
Personal history of substance abuse		
Alcohol	3	3
Illegal drugs	4	4
Rx drugs	5	5
Age between 16—45 years	1	1
History of preadolescent sexual abuse	3	0
Psychological disease		
ADD, OCD, bipolar, schizophrenia	2	2
Depression	1	1
Scoring totals		

Assessment of Risk-SOAPP

The **Screening and Opioid Assessment for Patients with Pain (SOAPP)**[®] helps determine **how much monitoring** a patient on long-term opioid therapy might require. It comes as a short and a standard form.

SOAPP endeavors to **minimize the chances of missing high-risk patients**. This means that patients who are truly at low risk may still get a score above the cutoff. The SOAPP is less good at identifying who is not at-risk.

The SOAPP is scored as $>$ than 4 or less than 4.

Assessment of Risk-SOAPP

The tool asks interesting risk factors:

How often do you have **mood Swings**?

How often do you **smoke a cigarette** within an hour after you wake up?

How often have you had **legal problems** or been arrested?

SOAPP Version	SOAPP Cutoff Score	Sensitivity	Specificity	Positive Predictive Value	Negative Predictive Value	Positive Likelihood Ratio	Negative Likelihood Ratio
Short Form	Score 4 or above	.86	.67	.69	.85	2.59	.20
Standard	Score 7 or above	.91	.69	.71	.90	2.94	.13

SOAPP® Version 1.0 - SF

Name: _____ Date: _____

The following are some questions given to all patients at the Pain Management Center who are on or being considered for opioids for their pain. Please answer each question as honestly as possible. This information is for our records and will remain confidential. Your answers alone will not determine your treatment. Thank you.

Please answer the questions below using the following scale:

0 = Never, 1 = Seldom, 2 = Sometimes, 3 = Often, 4 = Very Often

1. How often do you have mood swings? 0 1 2 3 4
2. How often do you smoke a cigarette within an hour after you wake up? 0 1 2 3 4
3. How often have you taken medication other than the way that it was prescribed? 0 1 2 3 4
4. How often have you used illegal drugs (for example, marijuana, cocaine, etc.) in the past five years? 0 1 2 3 4
5. How often, in your lifetime, have you had legal problems or been arrested? 0 1 2 3 4

Please include any additional information you wish about the above answers. Thank you.

Keeping Patients on Opioids-The PEG

Checking for Functional Improvement

What number, from 0 – 10 best:

Q1: Describes your Pain in the past week?

Q2: Describes how, during the past week, pain has interfered with your Enjoyment of life?

Q3: Describes how, during the past week, pain has interfered with your General activity?

PEG score = 30% improvement from baseline is clinically meaningful.

BOP

Evaluating for Misuse

Prescription Drug Monitoring Programs (PDMPs) collect data from pharmacies on dispensed controlled substance prescriptions and make those data available to authorized users through a secure, electronic database. Currently, all states have operating PDMPs. **More than 20 have joined a shared database.**

Some EMRs can pull PDMP data.
Making a query more convenient.



Contracts

In order to receive controlled medications, patients often sign a:

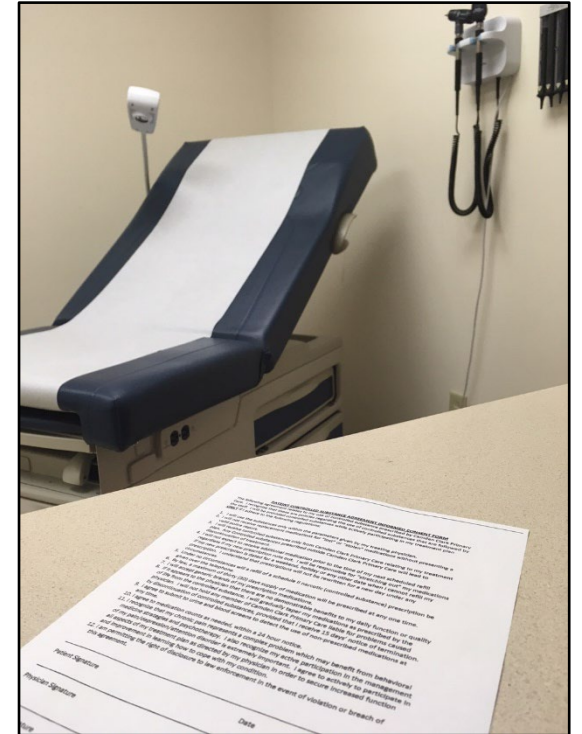
“Pain Contract,” or

“Treatment Consent Form,” or

“Narcotic Contract, ” or

“Opioid Treatment Agreement” (OTA),

There is **some criticism that OSAs are ethically suspect**, if not unethical, and should be used with extreme care.



Contracts

Use an agreement that defines the terms and expectations of therapy

It should outline:

- Appropriate intervals for follow-up,
- Refill policies,
- Participation in any indicated multimodal management plan (e.g., physical therapy, psychological treatment),
- Use of only one prescriber and one pharmacy for all controlled medications, and prohibition of illicit substance use or prescription diversion.
- Should be part of an ongoing treatment plan for all patients receiving chronic opioid therapy, thereby avoiding reliance on physician judgment, suspicion, or bias.

Breaks



Agenda

The First 30 Hour:

Why We Are here

The current situation

Assessment of Risk

Assessment of Need

BOP

Contracts

The Hour After That:

Prescribing Opioids

Opioid equivalents

Testing:

Urine

Pill counts

The Last Hour:

The New CDC
Guidelines

The WV Opioid
Reduction Act

Medication Review:

Initial Dosing

Controlled Substances Act, (1970)

WV Code: Chapter 60A

Recent change

Schedule	Definition	Examples	WV Rx Authority
1	Drugs with potentially severe physiological and psychological dependence. Considered to have no acceptable medicinal qualities.	Heroin, Marijuana , MDMA (ecstasy), Lysergic acid diethylamide (LSD), GHB (date rape drug), Mescaline, Cathinone (used in bath salts), Peyote, Psilocybin and psilocin (mushrooms)	(For Details check w/ Licensing Boards)
2	Commonly abused and causing dependency. They have a legitimate medical use. Must be prescribed to a patient by their provider and received from a pharmacist	Cocaine, OxyContin , Dilaudid, Methadone, Fentanyl, Vicodin , Methamphetamine, Dexedrine, Adderall , Ritalin	
		Ketamine, Tylenol with codeine , Buprenorphine (Suboxone, Subutex), Anabolic steroid, Testosterone,	MD/DO. PA: Prescribe. NP Prescribe, Dispense, Administer
4	Less chance of addiction or abuse. Providers allowed to include 5 refills in a 6 month period without additional consultation.	Xanax , Valium, Klonopin, Ambien , Sonata, Lunesta, Phenobarbital (long acting barbiturate), Modafinil (Stimulant-like drug), lomotil	MD/DO. PA: Prescribe. NP Prescribe, Dispense, Administer
5	Lowest likelihood for abuse, don't require a prescription and may be refilled.	Robitussin AC , Pyrovalerone (for chronic fatigue/appetite suppressant), Some anticonvulsants (Lyrica, gabapentin)	MD/DO. PA: Prescribe. NP Prescribe, Dispense, Administer

The Physician does not Rx Marijuana in WV, the Marijuana dispensary hires an employee to recommend dosing for the patient.

MD/DO
PA/APRN

Initial Dosing

Expect Adverse Effects:

Most Common - Constipation

Greatest Mortality - **Respiratory Depression**

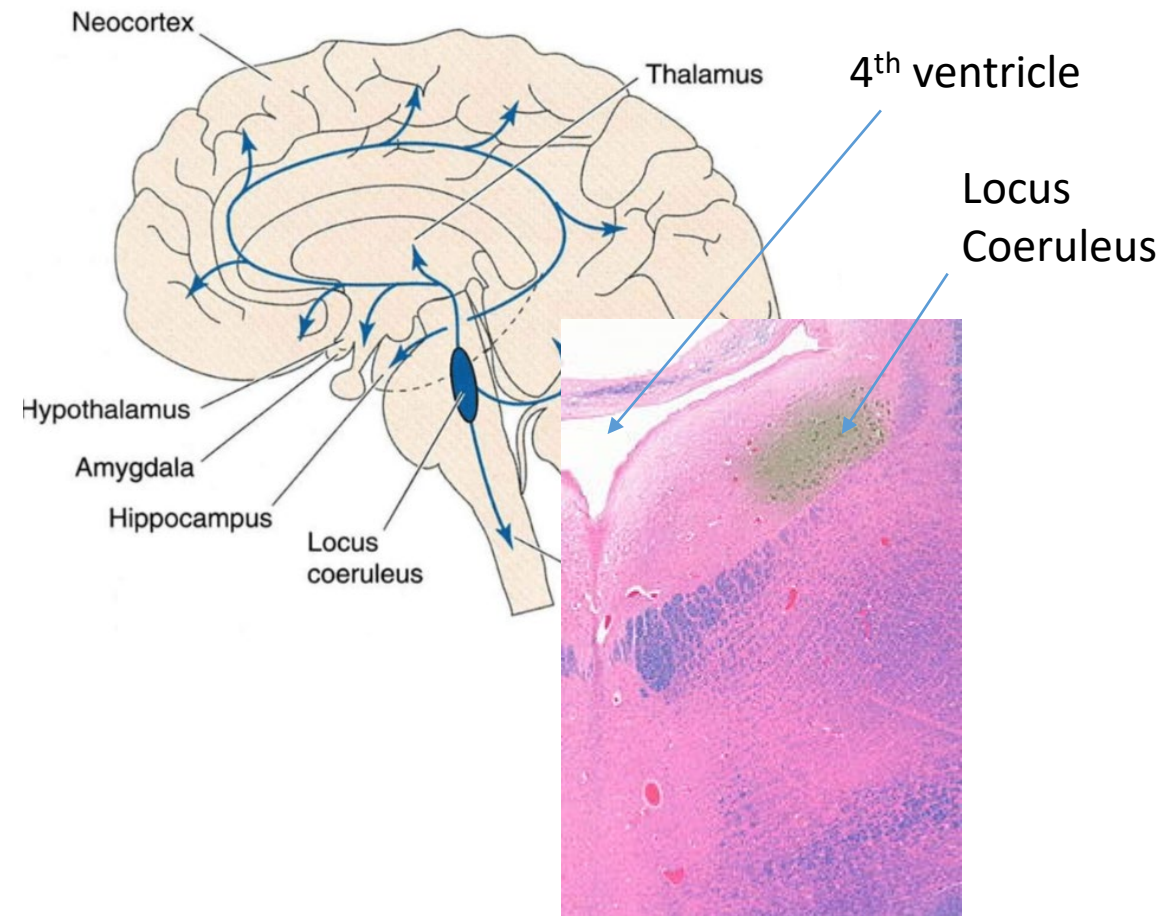
Other serious Mortality - Dependence, Hyperalgesia, Addiction

Other concerns: Sedation, Falls, Hypogonadism, Sweating, Miosis, Urinary Retention.

Initial Dosing

Why Respiratory Depression?

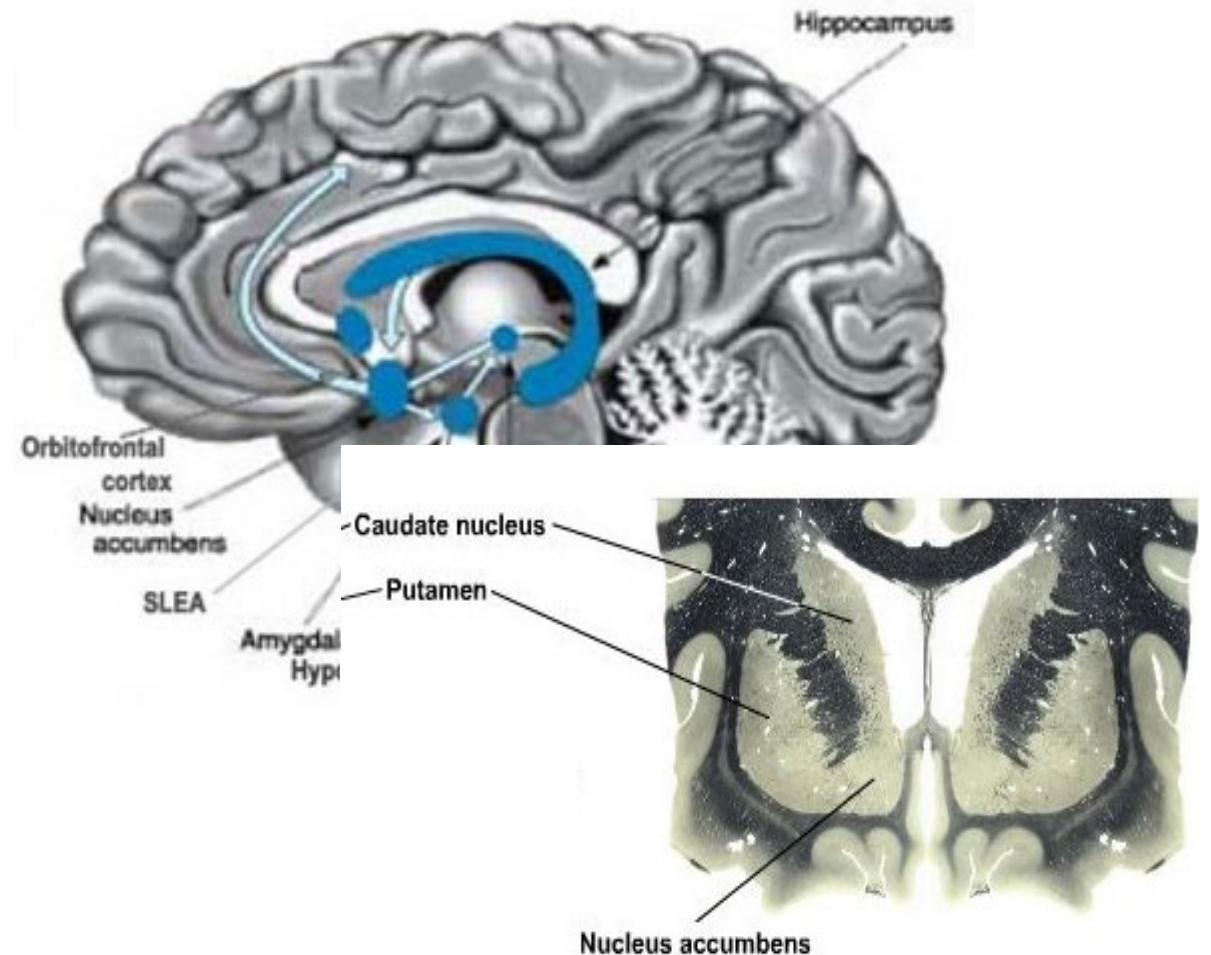
When opioid receptors in the **locus coeruleus** (brainstem) are activated they **prevent the release of noradrenaline** – decreasing alertness and blood pressure, increasing drowsiness, slow respiration (breathing), and induce analgesic (pain relieving) effects.



Initial Dosing

Why Respiratory Depression?

Opioid receptor activation in the **nucleus accumbens** releases **dopamine**, stimulating feelings of pleasure.



Initial Dosing

Expect Adverse Effects:

Identify Risk of Respiratory Depression

- Elderly, Debilitated
- Co-administered with other Resp Depressants
- Opioid Naive patients
- **Contraindicated** in those with Respiratory Suppression risks (COPD?)



<http://oxygenadvantage.co.za/2017/02/15/simu-late-high-altitude-training-acclimatization/>

Initial Dosing

Expect Adverse Effects:

Greatest Mortality-Respiratory Depression

- Discuss risk with patient
- Explain how hypercapnia exacerbates the sedating effect of opioids
- Urge supervision during initial use
- Encourage immediate use of 911 if respiratory depression suspected
- Offer Opioid Antagonists-**Naloxone**

Initial Dosing

Sustained-release preparations..

Offer more consistent drug levels.

Use caution when a patient demands only short-acting medication.

High potential for abuse or diversion:

fentanyl (Duragesic) patches

oxycodone (Roxicodone)

methadone (but easily monitored)

Lower risk, and has less abuse potential:

sublingual or transdermal **buprenorphine**

Initial Dosing

Black Box Warnings:

All opioid agonist-

- Addiction, abuse, misuse.
- Respiratory Depression.
- Accidental Ingestion.
- Neonatal withdrawal syndrome.
- CYP450 3A4 interactions.
- Risk of Concomitant use w/ Benzo./ CNS Depressants.

Acetaminophen warning-for acute liver failure.

MOA: Opioid Receptor Agonist.

Initial Dosing

Safety and Monitoring

Renal Dosing: Cr at baseline, then if severe renal disease or >65YOA check periodically.

Hepatic Dosing: if acetaminophen check LFTs, esp if severe hepatic disease.

Pregnancy/Lactation: may result in androgen deficiency-limited studies. Weigh risk/benefit if prolonged use. Risk of NAS. The risk of fetal harm is low. Acetaminophen is analgesic drug of choice while breast feeding.

Initial Dosing

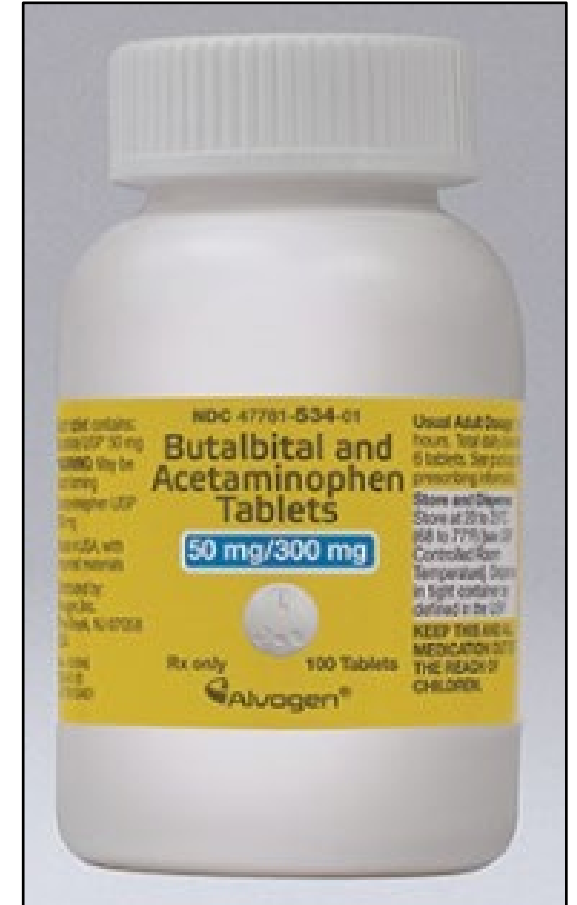
Butalbital/Acetaminophen

Brand Name: Bupap, Allzital

DEA Schedule II-V, status by state laws

MOA: Butalbital produces sedation/Selective COX-2 inhibitor. **Half-life 35 hours.**

Tension HA: 1-2 tabs PO q 4h prn. Limit to <300mg/day butalbital, 1g/4hrs and 4 g/day acetaminophen



Initial Dosing

Butalbital/Acetaminophen/Caffeine/Codeine

Brand Name: Fioricet with Codeine, 50/300/40/30

DEA Schedule III status by state laws

MOA: Butalbital produces sedation/Selective COX-2 inhibitor/binds to opioid receptors.

Tension HA: 1-2 caps PO q 4hrs prn. **Max 6 caps a day.** Limit to <300mg/day butalbital, 1g/4hrs and 4 g/day acetaminophen, 360 mg/day codeine.

Taper: 25-50% q 2-4 days



Initial Dosing

Acetaminophen/Codeine

Brand Name: Tylenol No. 3 (300/30mg)

DEA Schedule III with Black Box warnings

MOA: Selective COX-2 inhibitor /Binds to Opioid receptors

Mild to Mod pain:15-60mg of codeine PO q 4-6hrs PRN, >60mg not more effective.



Initial Dosing

Tramadol

Brand Name: Ultram (50mg, ER 100, 200,300mg)

DEA schedule IV, Black Box warnings

Half-life: 6-8 hours, Renal Dosing CrCl < 30.

MOA: Binds to MU Opioid receptors (Central opioid agonist) and inhibits Norepinephrine/serotonin reuptake.

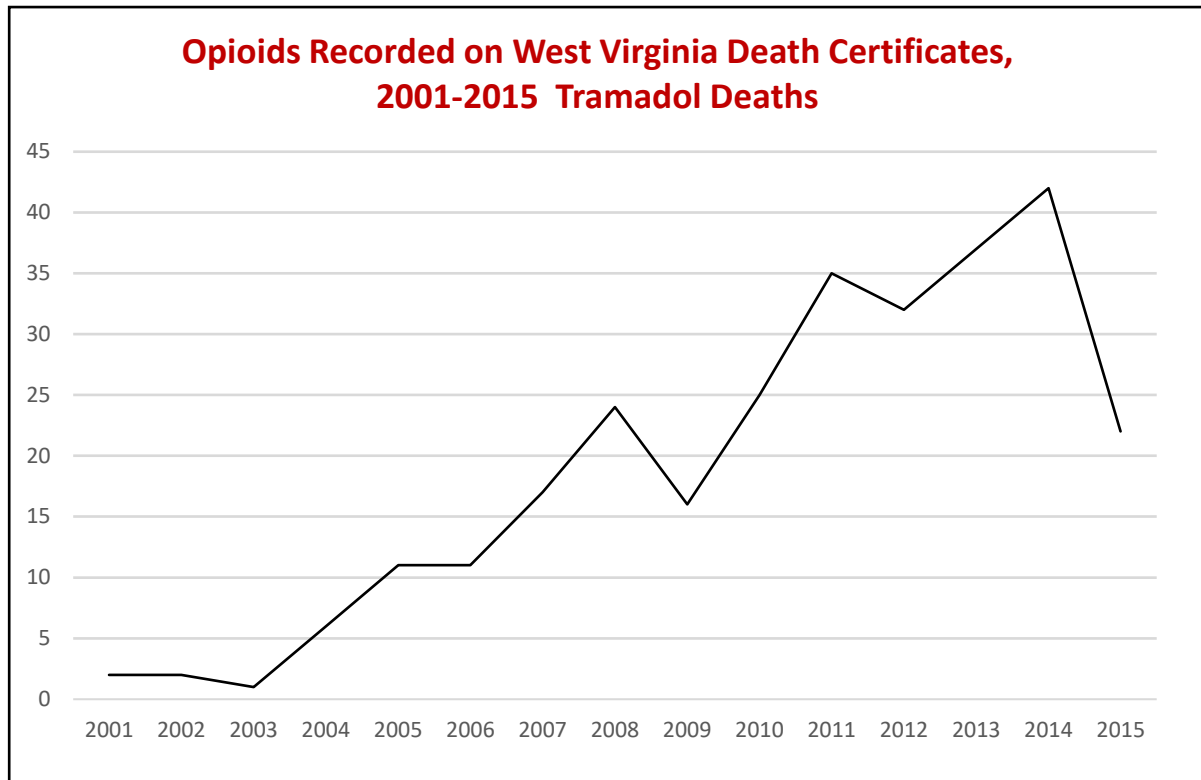
Mod-severe pain: Start 25 mg PO q AM. Increase 25 mg Q 3 days. Titrate to Q 4 hrs., **Max 400mg.** PRN, >60mg not more effective.

Chronic, Mod-severe Pain: ER PO Q day increase by 100 mg/day q 5 days, **Max of 300 Q day.** Conversion from IR 1:1



Initial Dosing

Tramadol-Associated Deaths



The opioid painkiller Tramadol has increased from less than **1 million pills in 2011 to 35.7 million in 2016.**

“It’s way less potential for abuse, way less diversion,” (Mike) Goff said.

- Eric Eyre WV Gazette.
Jan 21, 2017

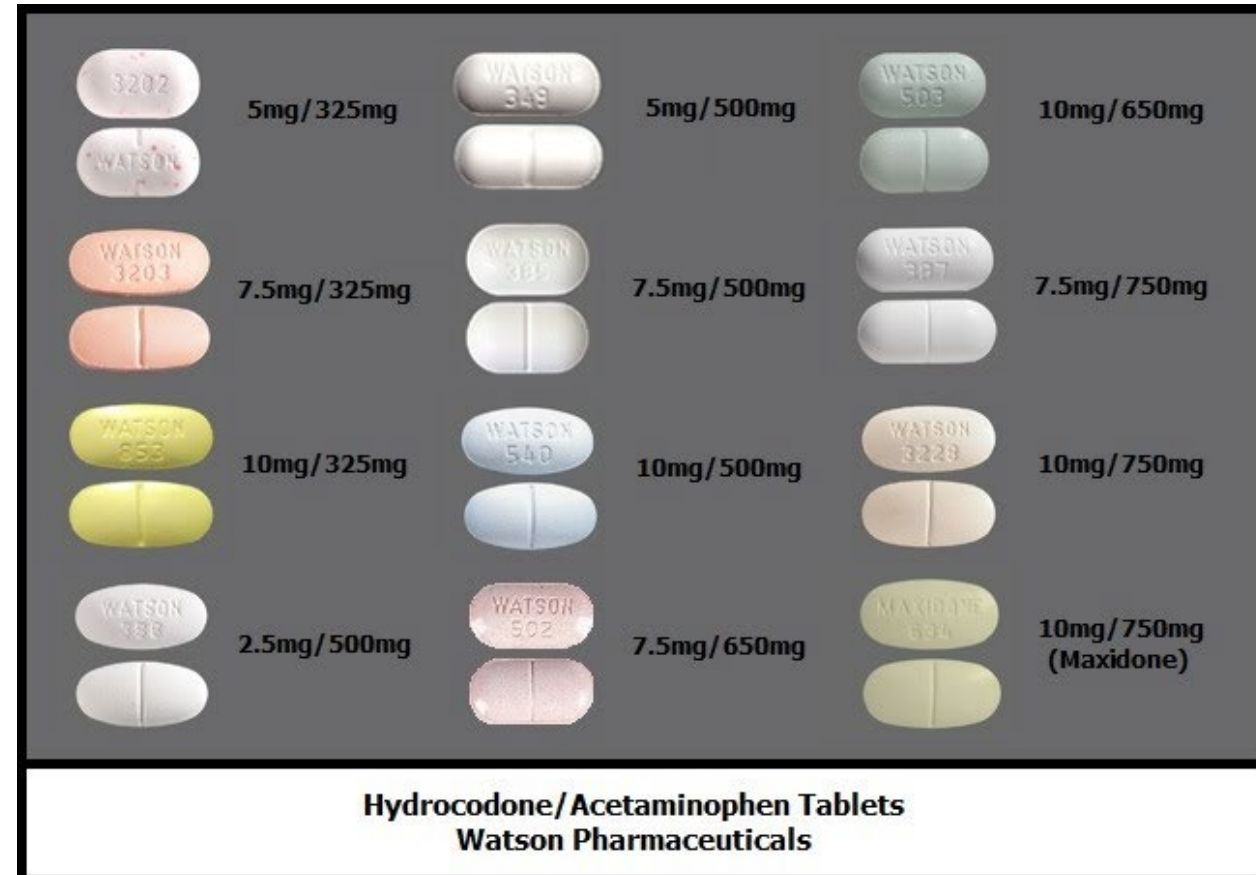
Initial Dosing

Hydrocodone/Acetaminophen

Brand Name: Norco, Vicoden, Lortab, etc.

DEA Schedule II

Pain, Moderate-Moderate Severe
2.5-10mg hydrocodone PO q4-6hrs
PRN. DO NOT Exceed **1g/4hr** or **4g/day** of acetaminophen from all sources



Initial Dosing

Oxycodone/Acetaminophen

Brand Name- Percocet. An alternative for patients with morphine intolerance or allergy.

DEA Schedule II.

Pain, Moderate-Moderate Severe 2.5-10mg oxycodone **PO 6hrs PRN.** May use Q 4 four hours for uncontrolled pain/tolerance.



Initial Dosing

Oxycodone

An alternative for patients with morphine intolerance or allergy.

Has a higher risk of abuse use with caution in patients with higher risk scores.

Long-acting oxycodone: not recommended for patients with chronic pain because it is not truly long-acting, is expensive, and has a high street value.



Initial Dosing

Oxycodone

Brand Names: Roxicodone, Oxycontin

DEA Schedule II

Pain, Mod-Severe: 5-15mg PO q 4-6hrs, PRN, Titrate slowly in elderly, (renal). Taper dose 25-50% q 2-4 days.

Pain Severe, Chronic: Start 10 mg ER PO q 12 increase 25-50% q1-2 days. Titrate slowly in elderly. Reduce dose 50% in debilitated pt > 65YOA. >40mg ER for use in opioid tolerant patients only.



Initial Dosing

Transdermal Fentanyl

More steady, may be a better alternative, it is expensive and can produce tolerance relatively quickly. **DO NOT use on opioid naive patients.**

DEA/FDA schedule II

Fentanyl is lipophilic, and absorption is affected in patients with little subcutaneous fat and in those prone to edema at application sites



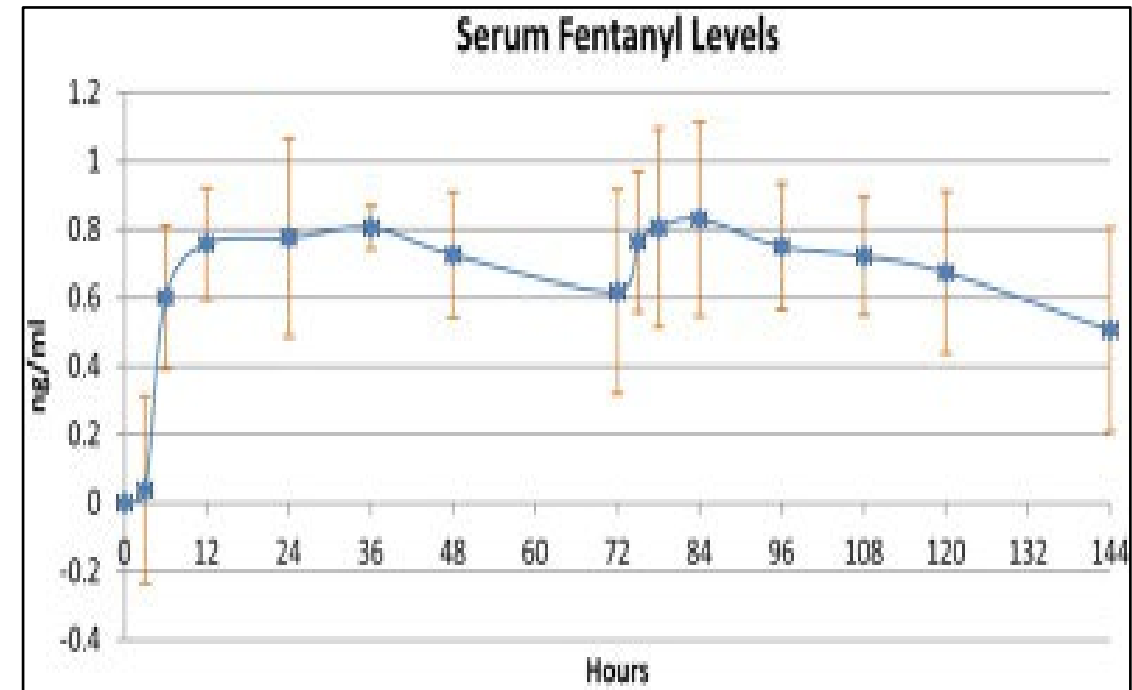
Initial Dosing

Transdermal Fentanyl

The pkg insert has conversion tables. **Adjust dose after three days, then no more frequently than Q 6 days.** Do not cut patch.

Start lower dose in the elderly.
Renal/Hepatic dosing-moderate impairment, start dose at 50%.

To discontinue: Taper dose 50% q 6days.



Initial Dosing

Methadone

Effective for many patients, may produce less tolerance than other opioids.

DEA/FDA schedule II,

It is inexpensive, long-acting, and has a combination of opioid and *N*-methyl-D-aspartate receptor activity that may make it **a good choice for patients with mixed somatic and neuropathic pain.**

However, **physicians who prescribe methadone must be familiar with its use.**

Initial Dosing

Methadone

Unique pharmacokinetics!

Very **long elimination half-life**(12 hrs.), and its MME conversion ratio increases as dosages increase.

Starting dosages in opioid-naive patients are 2.5 to 5 mg q 8-12 hours.



Initial Dosing

Methadone

Brand Name: Methadose, 40mg dispersible tab

Can be used for treatment of **opioid dependence**

Start: 15-30 mg PO X1 **then** 5-10 mg PO q 2-4 hours PRN.

Max: 40 mg on day 1, stabilize dose Q 2-3 days, **then decrease dose** by up to 20% q 24-48 hours. Document if using > 40 mg.

Initial Dosing

Methadone

After initial control of symptoms, dosages should be titrated slowly and no more than once per week.

Side Effects: **Methadone can prolong the QT interval,** avoid co-use with other QT-prolonging medications. Serum drug levels can be used for monitoring.

Methadone does not interfere with urine testing for other opioids.



Initial Dosing

Methadone

Onset of Action-0.5-1 hour.

Peak effect: 1-7.5 hours.

Steady state w/ continuous dosing: 3-5 days

Methadone can prolong the QT interval, avoid co-use with other QT-prolonging medications. Serum drug levels can be used for monitoring.

Initial Dosing

Methadone:

Elimination of methadone is **by hepatic metabolism**, followed by renal and fecal excretion.

Transformed to :

2-ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine (**EDDP**) and 9 other metabolites.

Metabolized by CYP 3A4

So plasma concentrations are **increased** by:

Macrolides, fluoroquinolones, SSRIS, TCAs (decrease dose by 25%), and

Decreased by:

Antiepileptics, Antipsychotics, antiretrovirals. (encourage rescue medications)

[https://www.painweek.org/assets/PAINWeekEnd%20Slides/2018/Dallas/Saturday_06_IV%20Methadone%20\(Aljassem\).pdf](https://www.painweek.org/assets/PAINWeekEnd%20Slides/2018/Dallas/Saturday_06_IV%20Methadone%20(Aljassem).pdf)

Initial Dosing

Buprenorphine Transdermal

Brand Names: Butrans

DEA/FDA schedule III

Dosed by Equivalents:

Opioid Naïve: 5 mcg patch Q 7 days.

<30 MME mg/day: Start 5 mcg patch, increase q 72 hours. Max is 20 mcg.

30-80 MME mg/day: Start 10mcg patch, increase q 72 hours. Max is 20 mcg.

Half Life: 26 hours. Watch Resp Depression onset-24-72 hours



Initial Dosing

Buprenorphine

DEA/FDA schedule III partial opioid agonist that is less likely to produce tolerance.

It is effective for treatment of pain, has lower abuse potential, and is easily monitored.

It is expensive, and its use requires special prescriber training (**except for the transdermal patch**).

May use while breast feeding.



Initial Dosing

Gabapentin

Brand Name: Neurontin

Scheduled-IV. States regulate for association with overdoses.

MOA: blocks voltage dependent calcium channel

Post Herpetic Neuralgia: Start 300 mg PO Q Day X 1 day, then 300 mg PO BID, then 300 Mg TID, Max 1800 mg. Taper dose over 7 days to D/C.



Initial Dosing

Gabapentin

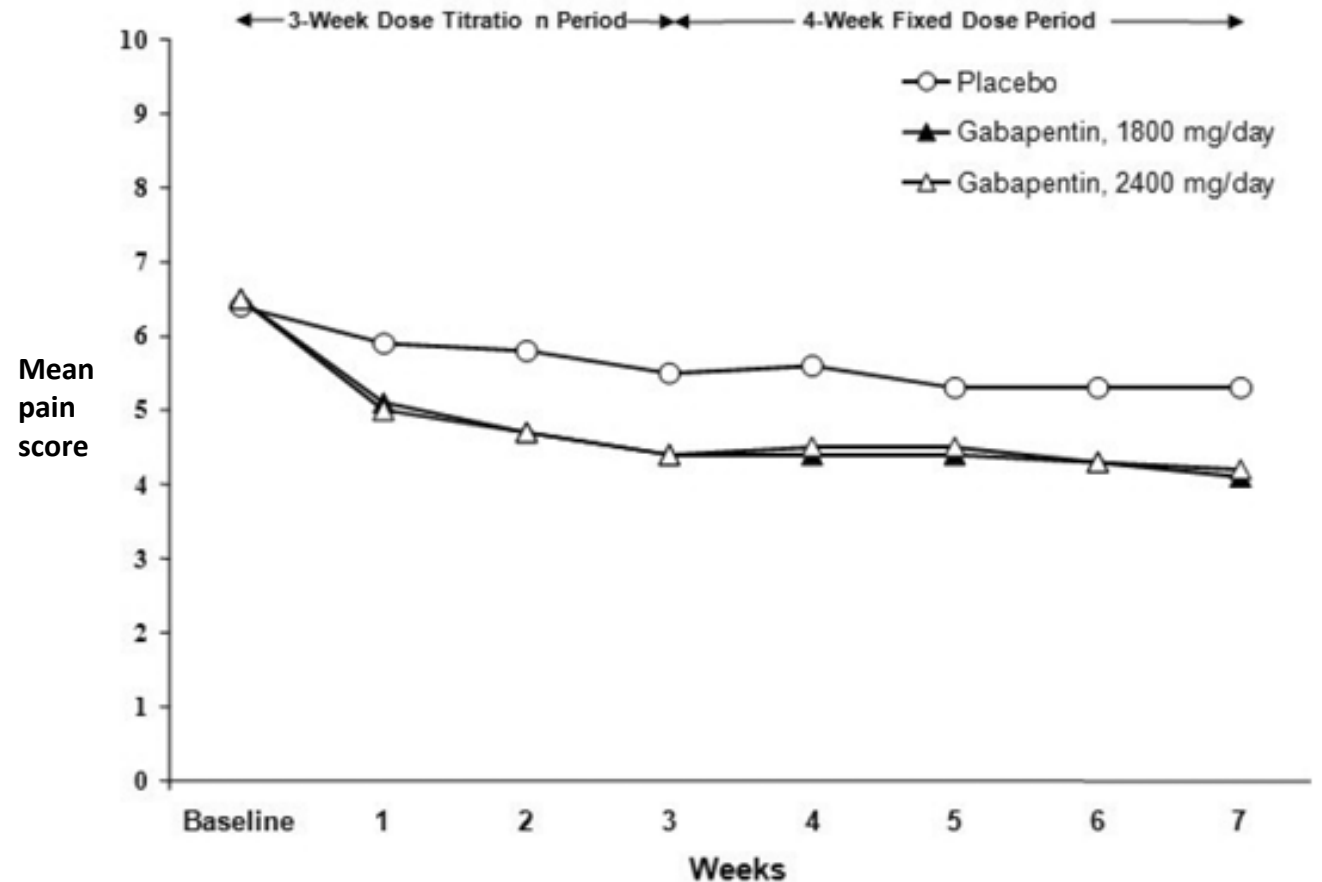
Neuropathic Pain:

Start 300 mg PO Q Day X 1 day,

Then 300 mg PO BID,

Then 300 mg TID. **Max 3600 mg.**

Taper dose over 7 days to D/C.



Initial Dosing

Neurontin Associated w/ Opioid Overdose?

Smith et al (2016) completed a [literature review on gabapentin misuse](#).

- The authors defined misuse as taking a larger dosage than prescribed or use without a prescription, or diversion.
- 33 varied peer-reviewed papers demonstrating gabapentin misuse were reviewed.
- **Prevalence of misuse in the general population was reported to be 1%,**
- 40-65% among individuals with prescriptions and
- **15 and 22% within populations of people who abuse opioids.**
- Gabapentin was misused for recreational purposes, self-medication or intentional self-harm and was misused alone or in combination with other substances, especially opioids, benzodiazepines and/or alcohol.

They concluded that **gabapentin is being misused internationally, within substance abuse populations**

Initial Dosing

WV Gabapentin Associated Deaths

Gabapentin-Related Overdose Deaths —
West Virginia, 2001-2015 (N=325)

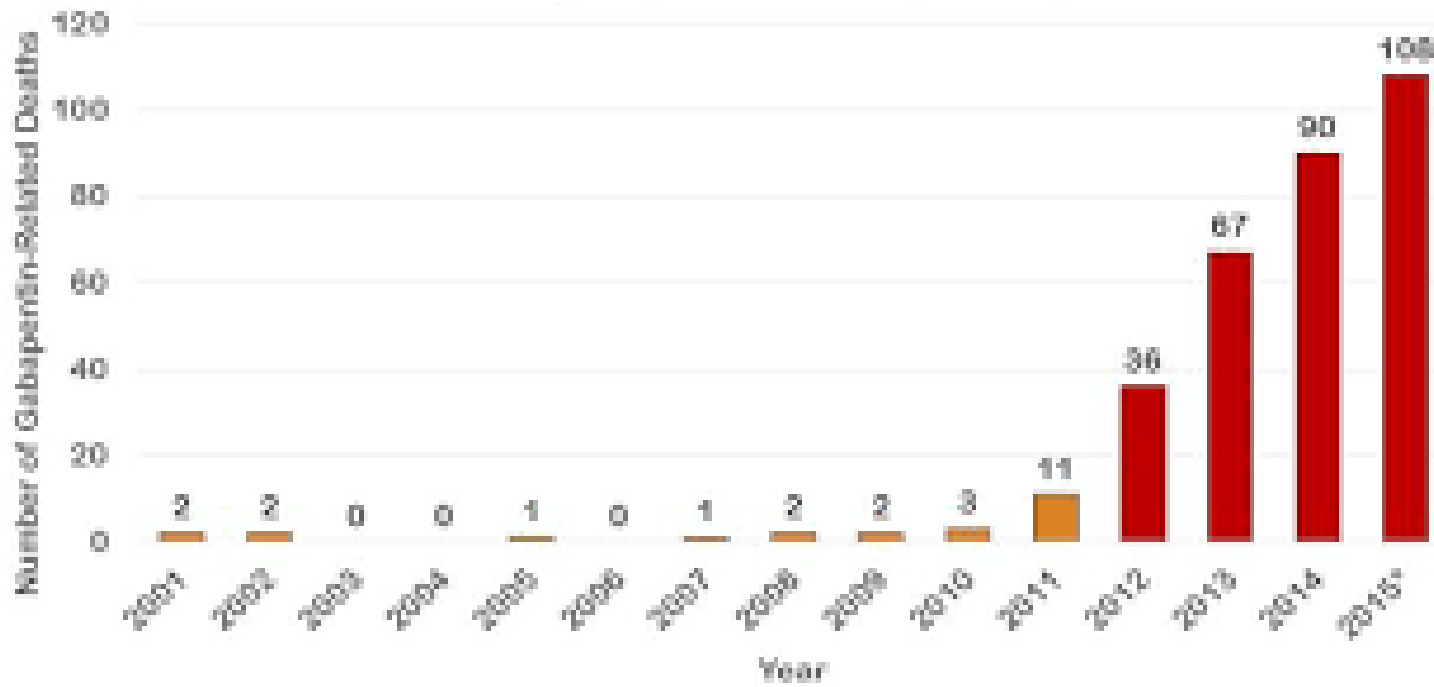
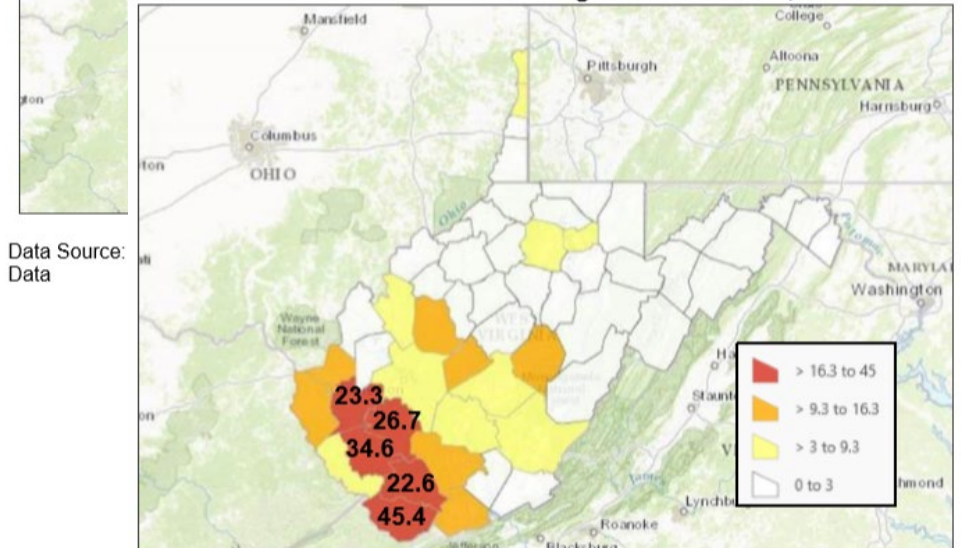


Figure 25: County-Level Distribution of Gabapentin-Related Overdose Deaths West Virginia Occurrences, 2001-2015



Figure 26: County-Level Distribution of Gabapentin-Related Overdose Deaths West Virginia Occurrences, 2001-2015



Data Source:
Data

Data Source: WV Health Statistics Center, Vital Statistics System 2015 Preliminary Data

Initial Dosing

Pregabalin

Brand Name: Lyrica

DEA Scheduled V. States may regulate for association with overdoses.

MOA: Binds Alpha2-Delta subunit of voltage dependent calcium channel reducing neurotransmitter release

Diabetic Neuropathic Pain Start 50 mg PO TID X 1 week, then 100 mg PO TID if needed. Taper dose over 7 days to D/C.

Post Herpetic Neuralgia or Other Neuropathic Pain Start 75 mg PO BID X 1 week, then 150 mg PO BID X 2 week, then 300 mg BID if needed. Taper dose over 7 days to D/C.

Initial Dosing

Recent Lyrica Concerns

A 2018 study describe **Australian patterns of pregabalin use** and intentional poisoning and identify people potentially at high risk of misuse.

Pregabalin dispensing **increased by between 2013 and 2016** and there were 88 pregabalin-associated deaths, a **57.8% yearly increase** per year of intentional pregabalin poisonings.

Initial Dosing

Recent Lyrica Concerns

Patients overdosing on pregabalin commonly co-ingested opioids, benzodiazepines, and illicit drugs, and had high rates of psychiatric and substance use comorbidities.

14.7% of pregabalin users were at high-risk of misuse.



<http://prescriptionassistance123.com/blog/lyrica-prescription-assistance-programs/>

Initial Dosing

Recent Lyrica Concerns

Those at high-risk of misuse were more likely to be:

- Younger
- Male
- Co-prescribed benzodiazepines or opioids
- Have more individual prescribers, and
- Have higher pregabalin strengths dispensed

Initial Dosing

Naloxone

Routes: Pre-filled syringe, Vial, intranasal routes

MOA: opioid antagonist, blocks opioids. $\frac{1}{2}$ life is 60-90 minutes.

Opioid Overdose (child or adult):

Narcan Nasal Spray: 4 mg dose sprayed into one nostril

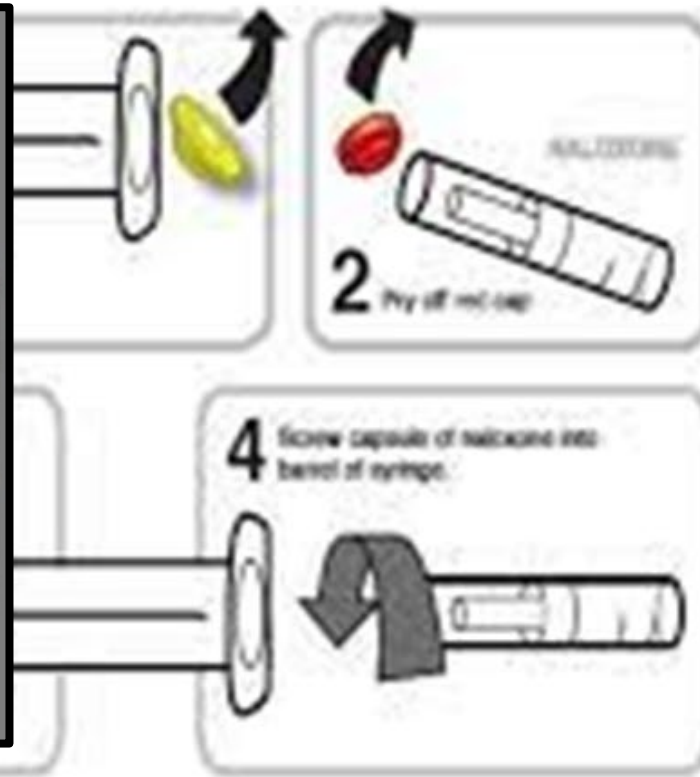
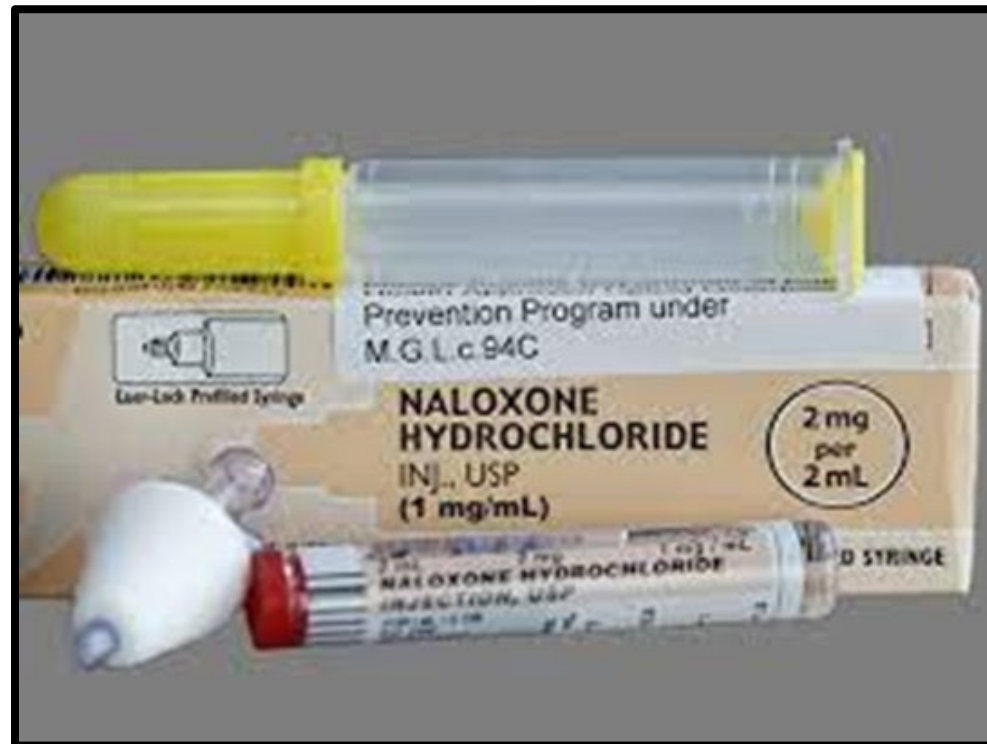
Naloxone Nasal via Atomizer: 2 mg dose sprayed into the nostrils (half in each nostril)

Evzio Auto-injector: 2 mg dose of naloxone injected into the outer thigh

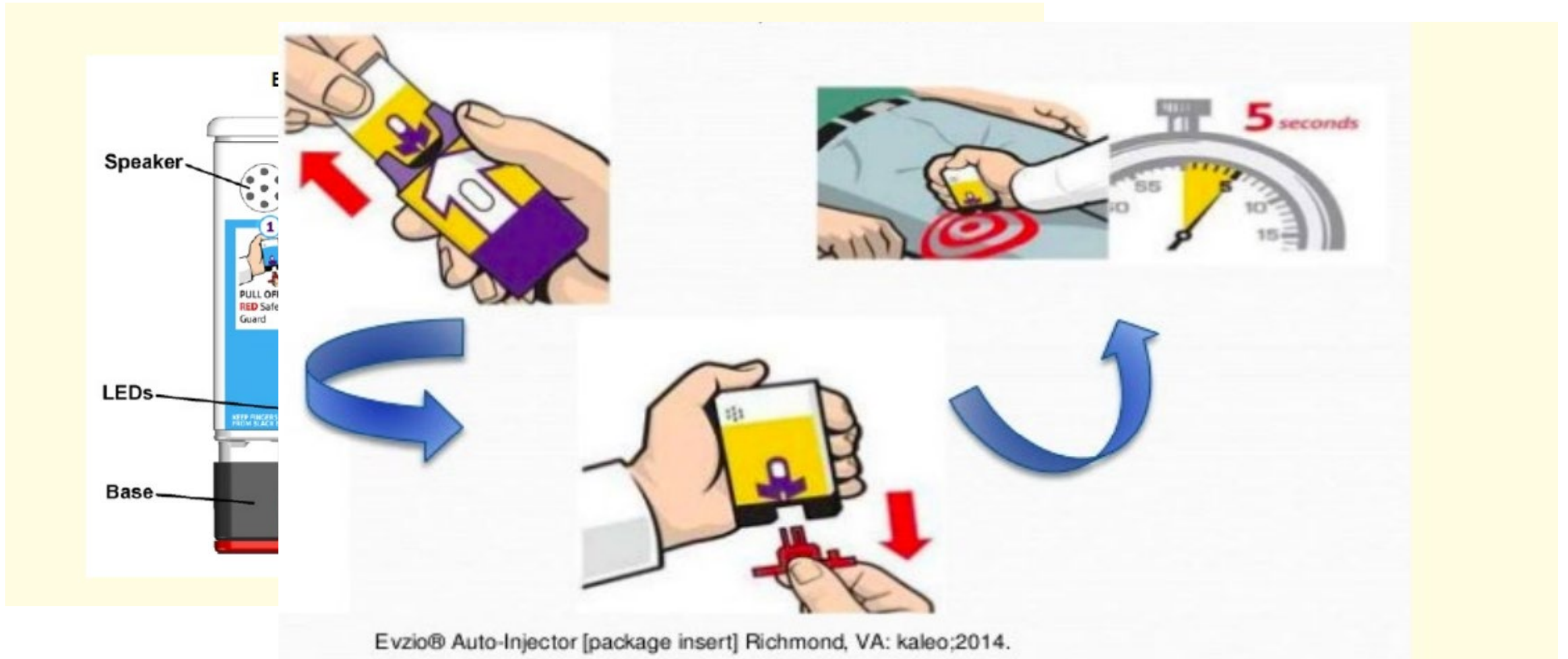
Call 911, Repeat after 4 min if symptoms reoccur.



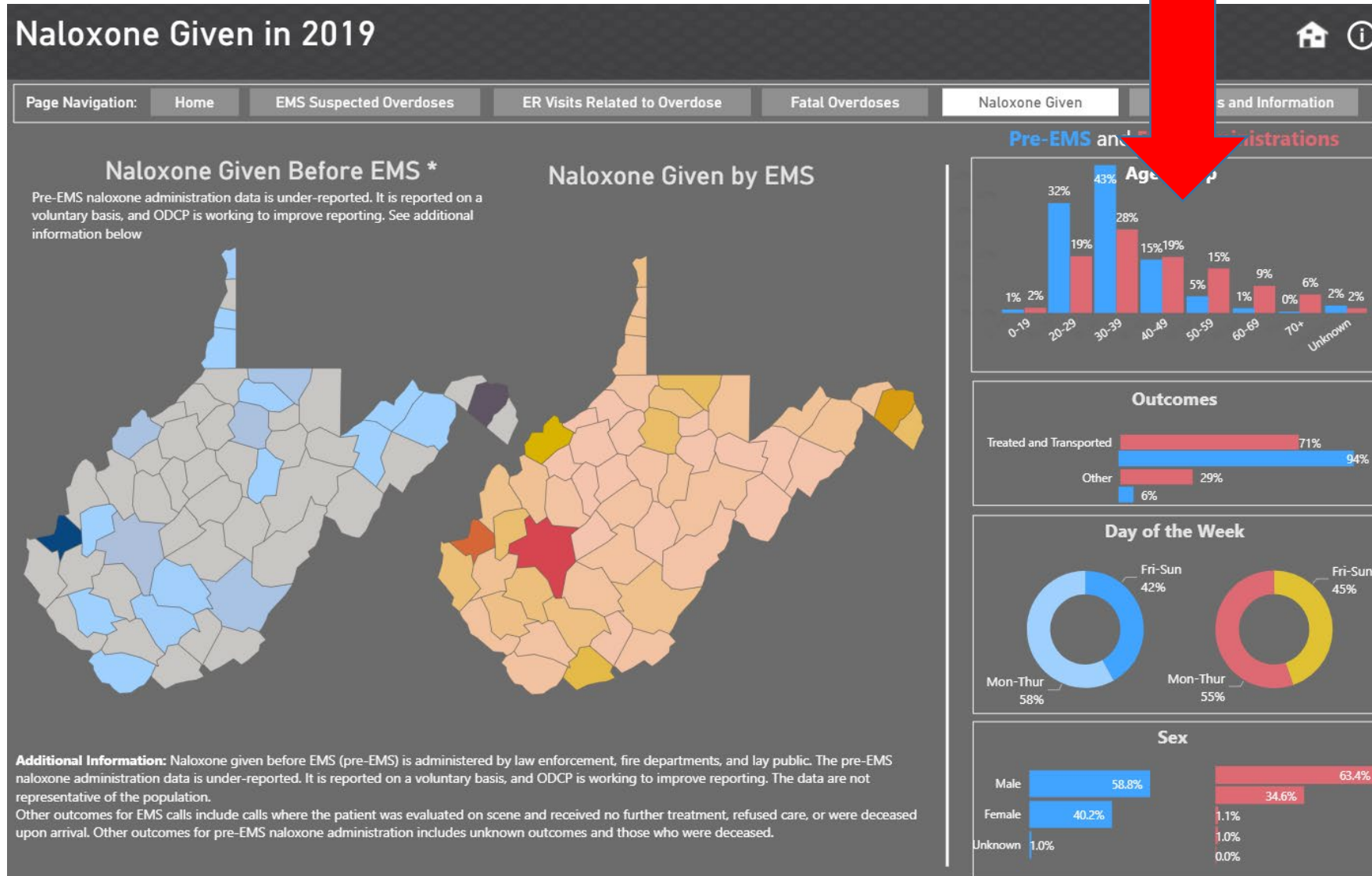
Initial Dosing-Naloxone



Initial Dosing-Naloxone



Initial Dosing



Of note:
If older than 40YOA EMS more likely to administer first dose.

Kanawha: 1905

Cabell: 1608

Berkeley: 1154

Opioid Equivalents

Determine the total daily amount of each opioid the patient takes.

Convert each to MMEs—multiply the dose for each opioid by the conversion factor.

Add them together.

Warning: Do not use the calculated dose in MMEs to determine dosage for converting one opioid to another—**the new opioid should be lower to avoid unintentional overdose** caused by incomplete cross-tolerance and individual differences in opioid pharmacokinetics. Consult the medication label.

Calculating morphine milligram equivalents (MME)

OPIOID (doses in mg/day except where noted)	CONVERSION FACTOR
Codeine	0.15
Fentanyl transdermal (in mcg/hr)	2.4
Hydrocodone	1
Hydromorphone	4
Methadone	
1-20 mg/day	4
21-40 mg/day	8
41-60 mg/day	10
≥ 61-80 mg/day	12
Morphine	1
Oxycodone	1.5
Oxymorphone	3

These dose conversions are estimated and cannot account for all individual differences in genetics and pharmacokinetics.

Opioid Equivalents

50 MME/day:

50mg Hydrocodone: 10 tablets of hydrocodone/ acetaminophen 5/300

33mg of oxycodone: 2 tablets of oxycodone sustained-release 15 mg

12mg of methadone: <3 tablets of methadone 5 mg

90 MME/day:

90mg Hydrocodone: 9 tablets of hydrocodone/ acetaminophen 10/300

60 mg of oxycodone: 2 tablets of oxycodone sustained-release 30 mg

20 mg of methadone: 4 tablets of methadone 5 mg

Case Study:

Your employer has recently lost three long established family physicians and management has informed you that, as a physician, you will have to assume the care of their patients on controlled medications. Your area has lost it's only non-interventionist pain specialist to a recent drug sweep.

On Monday, after this conference, the first patient presents. He's a 56 years of age former timber worker on disability for the last 22 years after being crushed by a tree and suffering a flailed chest and multiple vertebral fractures. He smokes, has COPD, his A1C is 9.8% and his LDL is 140.

He's on Metformin, Pro-air, and **hydrocodone 5 mg TID**. He's been on these meds for the last 6 years. He has refused preventative efforts and your EMR informs you of multiple care gaps including colonoscopy and immunizations.

Case Study:

Preparing for the 15 minute visit (he's established in the system) you prioritize your efforts.

Which of the following is the most worrisome health concern?

- A) His socioeconomic status
- B) His lack of preventative care
- C) His uncontrolled diabetes
- D) His chronic opioid use

Rx Opioid Overdose 17.2 per 100,000
Diabetes mellitus 47.6 per 100,000
Diseases of heart 267.0
Malignant neoplasms 256.3
Chronic lower respiratory diseases 92.6
Cerebrovascular diseases 58.3

Case Study:

You decide to focus on his diabetes during today's visit, but recognize you must address his opioid use as well.

You decide to make an effort to reduce his opioid use by limiting his number of pills but not his total MMEs.

What is his current MME use:

- A) 50 MME
- B) 30.5 MME
- C) 15 MME
- D) 5 MME

Case Study:

What choice or choices offer an equivalent MME?

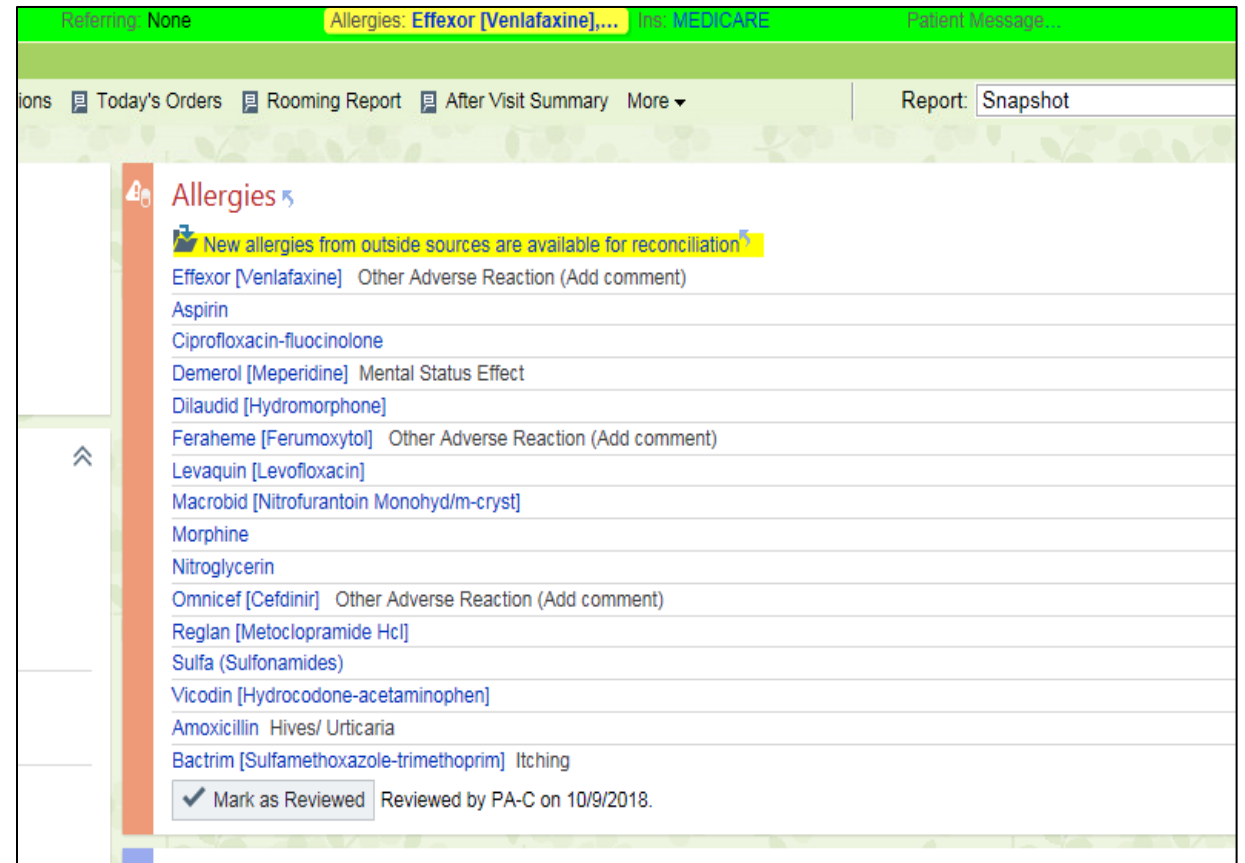
- A) Oxycodone 7.5mg BID
- B) Hydrocodone 7.5 mg BID
- C) Oxycodone 10mg QHS
- D) Hydrocodone 10mg Q AM

Behaviors Suspicious for Misuse or Dependency

- Taking a controlled substance for a long period of time (new patients)
- Refusing to grant permission to obtain old records or communicate with previous physicians
- Demonstrating reluctance to undergo a comprehensive history, physical examination, or diagnostic testing (especially urine drug screening)

Behaviors Suspicious for Misuse or Dependency

- Requesting a specific drug (often because of the higher resale value of a brand name)
- Professing **multiple allergies to recommended medications**
- Resisting other treatment options



The screenshot displays a medical software interface with a green header bar. The header contains the text "Allergies: Effexor [Venlafaxine],..." and "MEDICARE". Below the header, there are navigation tabs for "Today's Orders", "Rooming Report", "After Visit Summary", and "More". A "Report: Snapshot" dropdown is visible on the right. The main content area is titled "Allergies" and features a list of medications with their associated reactions. A yellow highlight is present over the text "New allergies from outside sources are available for reconciliation". The list includes:

- Effexor [Venlafaxine] Other Adverse Reaction (Add comment)
- Aspirin
- Ciprofloxacin-fluocinolone
- Demerol [Meperidine] Mental Status Effect
- Dilaudid [Hydromorphone]
- Feraheme [Ferumoxytol] Other Adverse Reaction (Add comment)
- Levaquin [Levofloxacin]
- Macrobid [Nitrofurantoin Monohyd/m-cryst]
- Morphine
- Nitroglycerin
- Omnicef [Cefdinir] Other Adverse Reaction (Add comment)
- Reglan [Metoclopramide Hcl]
- Sulfa (Sulfonamides)
- Vicodin [Hydrocodone-acetaminophen]
- Amoxicillin Hives/ Urticaria
- Bactrim [Sulfamethoxazole-trimethoprim] Itching

At the bottom of the list, there is a "Mark as Reviewed" button with a checkmark icon and the text "Reviewed by PA-C on 10/9/2018."

Behaviors Suspicious for Misuse or Dependency

Other aberrant behavior

- Issuing threats or displaying anger
- Targeting appointments at the end of the day or during off hours (nights or weekends)
- Giving excessive flattery



Behaviors Suspicious for Misuse or Dependency

- Calling and visiting a physician's associates
- Repeatedly losing a prescription
- Requesting a dose escalation
- Demonstrating noncompliance with prescription instructions
- Demonstrating other evidence of alcohol or illicit drug misuse



Testing:

Schedule patients additional time:

For example, writing a new prescription for a controlled substance would require evaluating the patient for a history of abuse or addiction, and may include screening.

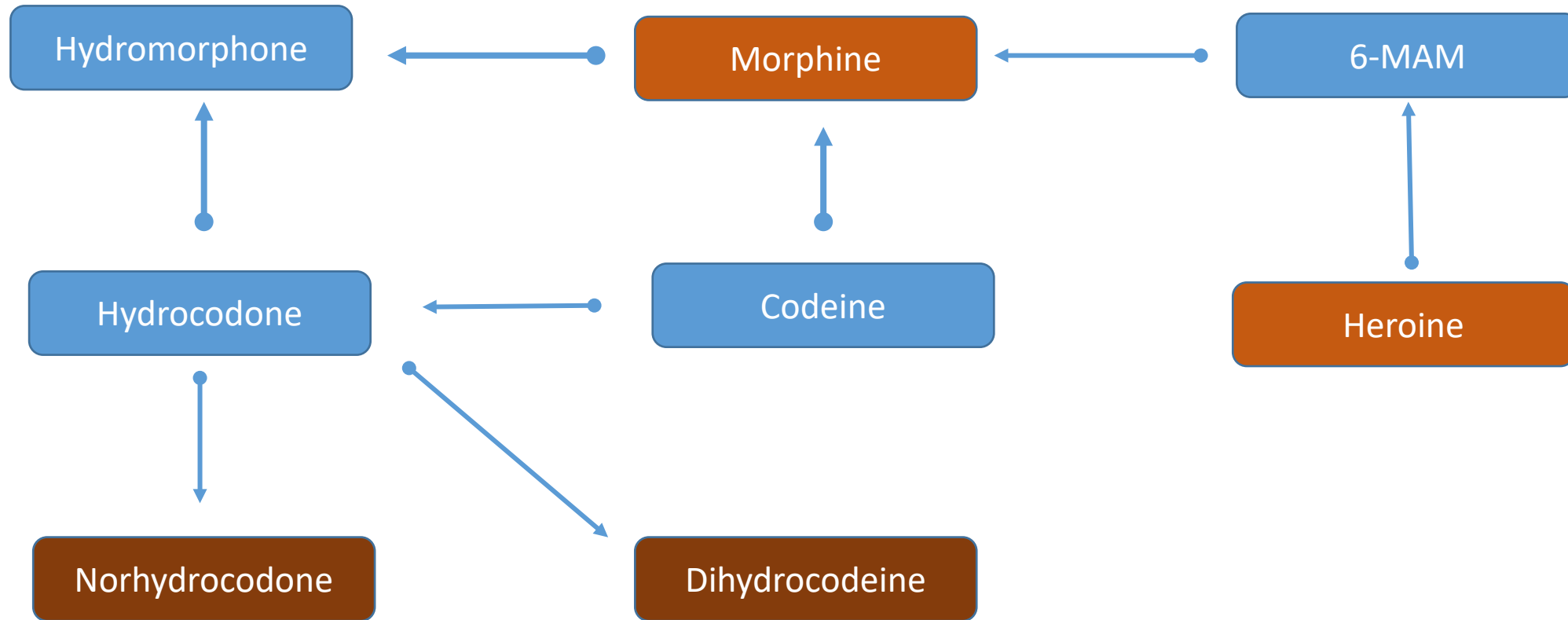


Testing:

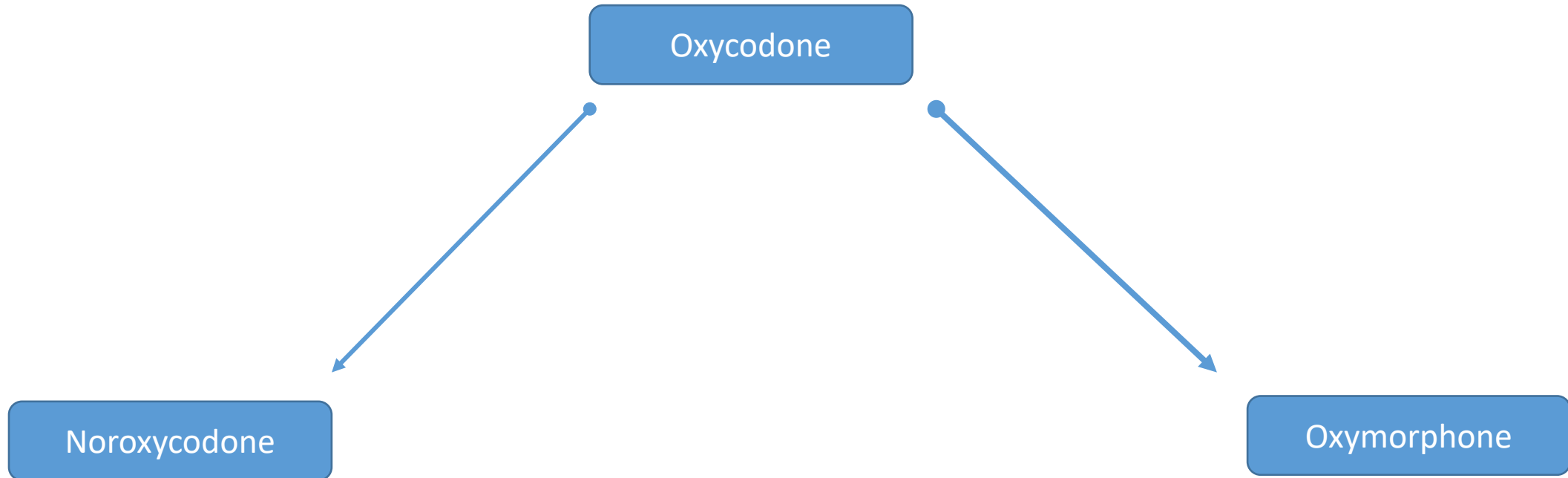
There are two main types of Urine Drug Screening:

- **immunoassay testing**, and
- **chromatography** (i.e., gas chromatography/mass spectrometry [GC/MS] or high-performance liquid chromatography)
- **Immunoassay tests are the preferred initial test for screening.** They use antibodies to detect the presence of drugs. These tests can be processed rapidly, are inexpensive.

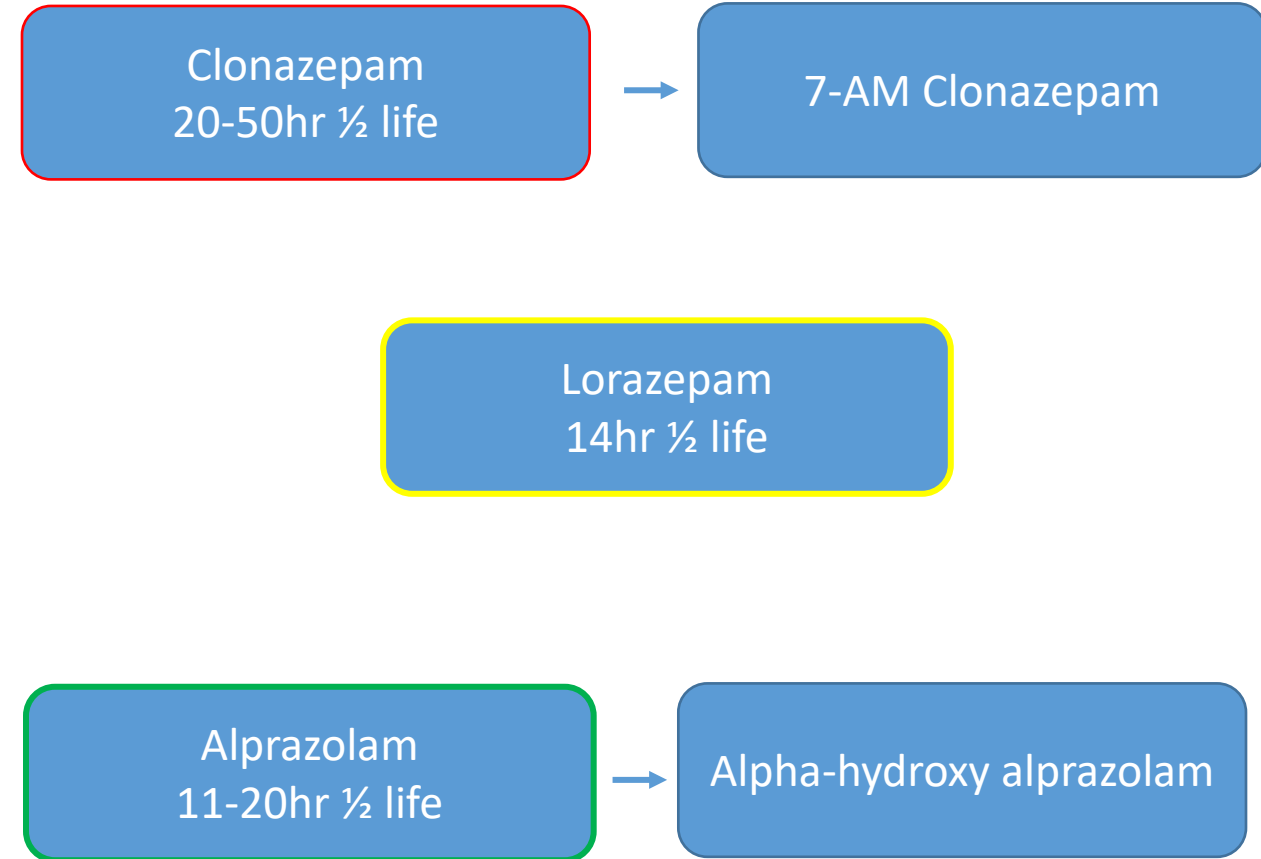
Opioids Metabolism



Opioids Metabolism



Benzodiazepines Metabolism



Urine Drug Screens

The most commonly ordered drug screens are for:

- Cocaine metabolites
- Amphetamines
- Phencyclidine
- Marijuana metabolites
- Opiate metabolites.

The **U.S. Department of Transportation** **requires testing for these five substances** when conducting urine drug screenings for transportation employees



<https://www.protocoldrugtesting.com/>

Urine Drug Screens

The Accuracy of Immunoassay

It varies..

- high predictive value for marijuana and cocaine,
- lower predictive value for opiates and amphetamines.

Many commonly prescribed medications **can cause positive** immunoassay tests



<https://www.wcpo.com/news/national/urine-screens-are-big-business-report-shows-costs-quadrupled-from-2011-to-2014>

Reducing UDS Tampering

- Request removal of any unnecessary outer clothing
- Remove anything in the collection area that could be used to adulterate or substitute a urine specimen
- Request the display and removal of any items in the patient's pockets, coat, hat, etc.



<https://www.synthetix5.com/synthetic-urine-belt-kit/>

<https://peepack.com/product/peepack-sterile-urine-kit-3-pack/>

Reducing UDS Tampering



- Require all other personal belongings (e.g., briefcase, purse) to remain with the outer clothing
- Instruct the patient to wash and dry his or her hands (preferably with liquid soap) under direct observation and not to wash again until after delivering the specimen
- Place a bluing agent in the commode and turn off the water supply to the testing site

Methods and Criteria for UDS

Collection Methods and Criteria

- Direct observation of specimen collection (when required)
- **Sample size:** 30 mL or more
- **Temperature:** between 90°F (32.2°C) and 100°F (37.7°C)
- **Urine pH:** 4.5 to 8.5

- Use of an approved chain of custody form to track specimen handling

Methods and Criteria for UDS

Findings Suggestive of Adulterated, Diluted, or Substituted Specimens

General: Temperature $< 90^{\circ}\text{F}$ or $> 100^{\circ}\text{F}$

Unusual appearance (e.g., bubbly, cloudy, clear, dark)

Adulterated: Nitrite concentration > 5 mg per dL

Urine pH < 3 or ≥ 11

Diluted/Substituted: Creatinine concentration < 2.0 mg per dL

Drugs that cause a false positive on UDS

Opiates

Dextromethorphan,
fluoroquinolones,
quinine,
verapamil‡

diphenhydramine,
poppy seeds,
rifampin,

Duration of Detectability: One to three days

‡—In methadone assays only.

Drugs that cause a false positive on UDS

Phencyclidine (PCP/Angel Dust)

dextromethorphan

ibuprofen

ketamine

thioridazine

venlafaxine

diphenhydramine

imipramine

meperidine

tramadol

Duration of Detectability: 7 to 14 days

Drugs that cause a false positive on UDS

Benzodiazepines

Oxaprozin and Sertraline

Duration of Detectability: 3 days for short-acting agents (e.g., lorazepam). Up to 30 days for long-acting agents (e.g., diazepam).

Cocaine

Topical anesthetics containing cocaine

Duration of Detectability: 2-3 days with occasional use, Up to 8 days with heavy use.

Drugs that cause a false positive on UDS

Amphetamines

Amantadine (Symmetrel),
Bupropion (Wellbutrin),
Chlorpromazine,
Desipramine (Norpramin),
Fluoxetine (Prozac),

L-methamphetamine
(in nasal decongestants*),
Labetalol (Normodyne),
Methylphenidate (Ritalin),
Phentermine,
Phenylephrine,

Phenylpropanolamine,
Promethazine(Phenergan),
Pseudoephedrine,
Ranitidine (Zantac),
Thioridazine,
Trazodone (Desyrel)

Duration of Detectability: Up to 3 days. *Current immunoassays have corrected the false-positive result for nasal decongestants containing L -methamphetamine.

Drugs that cause a false positive on UDS

Tetrahydrocannabinol

Dronabinol -(Marinol),

NSAIDs-ibuprofen, naproxen (Naprosyn), and sulindac (Clinoril),

PPIs- (pantoprazole [Protonix])

Duration of Detectability: 3 days with single use, 5-7 days with use around 4X per week, 10-15 days with daily use, **More than 30 days with long-term, heavy use**

Testing: Pill Counts

The main goal of a pill count is **to prevent diversion, misuse and abuse.**



Testing: Pill Counts

Request that the patient bring all unused pills to an appointment in the original container.

Notify the patient the **day before or the same day** as the appointment.

Check if the number of pills in the container match what the expected number would be if the patient followed the prescribed dosage.

Case Study:

Your 55 YO tree harvester has returned for his second visit.

His A1C is now 10.2%

You refer him to a local endocrinologist and focus on his opioid use.

What is the best way to determine he's using his opioids as ordered?

A) UDS

B) BOP

C) A pill count

D) the word of his wife who is here to attest to his need of opioids and his compliance

Case Study:

To set the appropriate expectations, you complete a UDS today. The results come back the next day and show THC.

At this point the best course of action is to:

- A) Inform the patient he will be dismissed.
- B) Repeat the test.
- C) Order a confirmation on the same sample.
- D) Ask the patient to return for an interview.

Break

