2022/2023 PBCME Opiate OD Deaths Partial year comparison 01/01-10/15

- ▶ PBC Medical Examiner –2022 **no pending cases**
 - ► Total drug overdose cases 553
 - ▶ Total opioid OD deaths 421 (76% of total OD cases)
 - ▶ Total Fentanyl & Fentanyl analog cause or presence 391 (93%)**
 - ▶ Decline in Opioid OD deaths 2021/2022 (19%)
- > PBC Medical Examiner –2023 (01/01-12/22) <u>snapshot</u>- 125 pending cases
 - > Total drug overdose cases 522
 - > Total opioid OD deaths 363 (70% of total OD cases) projected total 450 > 6%
 - > Total fentanyl & fentanyl analog cause or presence 276 (93%)
 - * Xylazine: "tranq" non-opioid animal tranquilizer 2022- 34/ 2023-26
 - ** New Fentanyl analogues:
 - N-Pyrrolidino Etonitazene (NPE) 20x more potent than Fentanyl 2022-9/2023-0
 - Fleurofentanyl similar potency to Fentanyl 2022-87/2023-58

PBCFR TRANSPORTS 2017-2023 January 1 – December 31

YEAR	#CALLS	# PATIENTS	%CHANGE/CALLS
2017	2675	2785	
2018	1509	1541	< 44 %
2019	1483	1510	< 2 %
2020	1771	1806	> 16 %
2021	1702	1743	< 4%
2022	1446	1471	< 15%
2023	1283	1309	< 11%

Net change 2017-2023 53% reduction in transports



FLORIDA STATE OXFORD HOUSES

Total Houses	164
Total Beds	1492
Men Houses	102
Men Beds	924
Women Houses	25
Women Beds	218
Women with Children Houses	34
Women with Children Beds	321
Men with Children Houses	3
Men with Children Beds	29

Cities that currently have Oxford Houses

West Palm Beach	Vero Beach	Lakeland	Bradenton
Lantana	Port St. Lucie	St. Petersburg	Ft. Myers
Riviera Beach	Ft. Lauderdale	Temple Terrace	Cape Coral
Lake Park	Lauderhill	Largo	Pensacola
Fort Pierce	Tampa	Clearwater	Panama City
Seminole	Winter Haven	Land O' Lakes	Panama City Beach
Tallahassee	Altamonte Springs	Apopka	Orlando
Winter Park	Jacksonville	Orange Park	St. Augustine
Palm Coast	Daytona	Port Orange	Ocala

Gainesville Miami Gardens Miami Kissimmee Deltona New Port Richie Jacksonville Beach Ft. Walton Sarasota Deland Palm Bay Sanford Crestview Naples Spring Hill Maitland

Cities we are looking to expand to in the near future

Destin
New Smyrna
Clermont
Punta Gorda
Port Charlotte

Overdoses since the last meeting: 1 (non-fatal)

The Oxford House initiative in Florida is continuing to gain momentum, with various chapters actively engaged in community-building and recovery efforts. Across the chapters, there is a clear emphasis placed on education, with initiatives like newcomer orientations and workshops geared towards deepening members' understanding of the Oxford House model. Alongside these educational endeavors, there's a strong focus on community and unity-building activities, ranging from fundraisers to holiday celebrations, creating an energetic and supportive atmosphere within the network of homes.

The commitment to outreach is a standout feature of the Oxford House network in Florida. Members and staff actively participate in local recovery events, have collaborated with law enforcement through presentations with different agencies, and contribute to community resources. This collective effort underscores a shared dedication to creating a sense of togetherness, evident in events like Thanksgiving & Christmas dinners as well as regional trainings. As the chapters look ahead, there's a proactive approach to sustaining this supportive environment, with future goals of continuing to open new houses in new areas, event hosting aimed at continual community building and ongoing training activities.

Some fast facts from our annual survey:

Average length of sobriety (days) - 333

Members who have reported opioid misuse - 60.8%

Members who reported use of Medically Assisted Treatment - 41.2%

Members who have reported stimulant misuse - 80.7%

Average Number of times members tried to get clean or sober - 7.2

Average number of times a member has been to detox without continuing to

Treatment - 2.9

Average amount of 12-Step meetings attended per week - 4.5

Members attending Counseling AND 12-Step Meetings - 40.4%

Importance of Oxford House to members' recovery (1 Unimportant - 5 Important) - 4.7

Upcoming Events:

Oxford House Florida State Workshop 22nd - 24th March 2024 (Paisley, FL) Oxford House World Convention Fall 2024 (Rosen Shingle Creek, Orlando)

All chapters in the state continue to do Narcan Administration trainings. All chapters are working with DCF to become Narcan providers for their areas.

National Website

www.oxfordhouse.org

Vacancy Website

www.oxfordvacancies.org

State Website

www.oxfordhousefl.org

Contacts:

Lori Holtzclaw-Hunt
Director of National Field Services
504-430-8554
lori.holtzclaw@oxfordhouse.org
Michael McKeogh
Regional Manager
601-402-6864
michael.mckeogh@oxfordhouse.org



PALM BEACH COUNTY 2024 UPDATE ON THE CONTINUUM OF CARE

STATE ATTORNEY ADDICTION RECOVERY TASK FORCE

JANUARY 4, 2024

PROVIDER REIMBURSEMENTS JULY - NOVEMBER 2023

COMMUNITY BASED TREATMENT \$28,058,722(5 YTD MONTHS) \$70,340,932 ANNUALIZED

ACUTE CARE

RESIDENTIAL TREATMENT

COMMUNITY BASED NON-TREATMENT SERVICES

\$7,845,740

28%

\$10,009,121

36%

\$5,504,040

20%

\$4,699,825

16%

Covered Service / Project	Total Paid
Assessment Total	\$409,886.94
BNET Total	\$35,011.41
CAT Team Total	\$662,323.14
Child Welfare FIT Team Total	\$264,337.75
Day Treatment Total	\$418,416.00
FACT Team Total	\$659,358.93
First Episode Team Total	\$80,486.02
Forensic Multidisciplinary Team Total	\$161,644.30
In-Home and On-Site Services Total	\$344,101.04
Intensive Case Management Total	\$351,009.43
Medical Services Total	\$1,367,008.57
Medication-Assisted Treatment Total	\$426,295.83
Outpatient - Group Total	\$521 <i>,</i> 769.79
Outpatient - Individual Total	\$1,717,841.61
Provider Level Care Coordination Total	\$426 , 247.63

PALM BEACH COUNTY

COMMUNITY BASED
TREATMENT

\$7,845,738.39

28%

Covered Service / Project	Total Paid		
Crisis Stabilization Total	\$6,894,585.20		
Crisis Support-Screening & Intake Total	\$294,997.50		
	·		
Short-term Residential TX Total	\$610,500.00		
	*1		
South County's Mobile Response Teams Total	\$1,677,023.67		
Substance Abuse Detaution Tatal	¢522.015.00		
Substance Abuse Detoxification Total	\$532,015.00		

PALM BEACH COUNTY JULY- NOVEMBER 2023 ACUTE CARE \$10,009,121.37 36%



Covered Ser	vice / Project	Total Paid
Resdie	ntial 2 Total	\$37,100.00
Reside	ntal 2 Total	\$48,251.14
Reside	ntial 2 Total	\$2,929,621.92
Reside	ntial 3 Total	\$1,003,663.50
Reside	ntial 4 Total	\$585,741.89
Reside	ntial I Total	\$288,400.00
Reside	ntiual 4 Total	\$240,498.21
Room	& Board Level 2 Total	\$99,534.42
Room	& Board Level 3 Total	\$271,225.82

PALM BEACH COUNTY

JULY-NOVEMBER 2023

RESIDENTIAL TREATMENT

\$5,504,036

20%

Provider	City	 al Contracted Service	Program Area	Service	Bed Capacity	Funded Capacity
Mandala Healing Center	West Palm Beach	\$ 535,200	Adult Mental Health	Residential Level I	40	3.5
Mandala Healing Center	West Palm Beach	\$ 350,400	Adult Substance Abuse	Residential Level I	40	3.5
SP Behavioral LLC DBA SandyPine	Tequesta	\$ 200,000	Childrens Mental Health	Residential Level I	149	1.0
Drug Abuse Foundation of Palm Beach County, Inc.	Delray Beach	\$ 3,424,119	Adult Substance Abuse	Residential Level II	84	38.0
Drug Abuse Treatment Association	West Palm Beach	\$ 1,049,038	Childrens Substance Abuse	Residential Level II	20	20.0
WaySide House	Delray Beach	\$ 845,100	Adult Substance Abuse	Residential Level II	28	14.0
Mandala Healing Center	West Palm Beach	\$ 2,175,600	Adult Mental Health	Residential Level III	16	16.0
Mandala Healing Center	West Palm Beach	\$ 1,912,400	Adult Substance Abuse	Residential Level III	16	16.0
Drug Abuse Foundation of Palm Beach County, Inc.	Delray Beach	\$ 176,400	Adult Substance Abuse	Residential Level IV	16	7.0
Jeff Industries, Inc.	Hypoluxo	\$ 46,055	Adult Mental Health	Residential Level IV	4	4.0
Sunset House, Inc.	Palm Beach Gardens	\$ 181,065	Adult Substance Abuse	Residential Level IV	34	6.0
The Lord's Place, Inc.	West Palm Beach	\$ 108,513	Adult Mental Health	Residential Level IV	50	4.0
The Lord's Place, Inc.	West Palm Beach	\$ 1,225,706	Adult Mental Health	Residential Level IV	24	24.0
South County Mental Health Center	Delray Beach	\$ 979,200	Adult Mental Health	Room and Board with Supervision Level II	37	16.0
WaySide House	Delray Beach	\$ 104,544	Adult Substance Abuse	Room and Board with Supervision Level II	28	2.0
Ebb Tide Treatment, LLC	Palm Beach Gardens	\$ 135,460	Adult Substance Abuse	Room and Board with Supervision Level III	28	8.0
Transpire Help	Lake Worth	\$ 173,996	Adult Substance Abuse	Room and Board with Supervision Level III	15	15.0
WaySide House	Delray Beach	\$ 	Adult Substance Abuse	Room and Board with Supervision Level III	28	8.0
		\$ 13,842,902				

Residential treatment programs funded in FY 23-24

PUBLICLY FUNDED BEHAVIORAL HEALTH SERVICES



PALM BEACH COUNTY

Numbers Served thru November, 2023						
		Year To Date Unduplicated				
Program -	Service Category	Serve⊽				
Adult Mental Health	Residential Care	191				
	Outpatient Care	2,717				
	Crisis Care	1,064				
	State Hospital Discharges	76				
	Peer Support Services	154				
Adult Substance						
Abuse	Residential Care	370				
	Outpatient Care	2,380				
	Detoxification	569				
	Women Specific Services	17				
	Injecting Drug Users	503				
	Peer Support Services	604				
Children Mental						
Health	Residential Care	5				
	Outpatient Care	988				
	Crisis Care	44				
Children Substance						
Abuse	Residential Care	48				
	Outpatient Care	594				

PERFORMANCE OUTCOME MEASURES



PALM BEACH COUNTY

✓ Performance Measures Outcomes

Measure	Description	Program	Program Type	Score	Target	Result
CSU	Percent of CSU Readmissions within 30 days		All	N/A	8.00%	ď
DETOX	Percent of DETOX Readmissions within 30 days	All	All	N/A	15.00%	Ó
M0003	Average annual days worked for pay for adults with severe and persistent mental illness	Adult	Mental Health	82.65	40.00	ß
M0375	Percent of adults with severe and persistent mental illnesses who improve their level of functioning	Adult	Mental Health	84.00%	64.00%	ß
M0376	Percent of adults with serious mental illnesses who improve their level of functioning	Adult	Mental Health	100.00%	65.00%	ß
M0703	Percent of adults with serious mental illness who are competitively employed	Adult	Mental Health	40.00%	24.00%	Ó
M0742	Percent of adults with severe and persistent mental illnesses who live in stable housing environment	Adult	Mental Health	92.00%	90.00%	ß
M0743	Percent of adults in forensic involvement who live in stable housing environment	Adult	Mental Health	98.00%	67.00%	ß
M0744	Percent of adults in mental health crisis who live in stable housing environment	Adult	Mental Health	83.00%	86.00%	\bar{C}
M0753	Percentage change in clients who are employed from admission to discharge	Adult	Substance Abuse	26.00%	10.00%	Ů

SEFBHN EXPANDED CAPACITY

Central Receiving System

988 / Mobile Response Teams

System Care Coordination

Service Addition/Expansion for Children, Youth, & Families:

Early Childhood Community Action Treatment (CAT) Team

Family Intensive Treatment (FIT) Expansion

Multisystemic Therapy Team (MST)

Expanded Capacity for the Adult Behavioral Health Continuum of Care:

Short-Term Residential Treatment

Conditional Release Residential Treatment

FACT Team Expansion

Expansion of Med Management/Medical Services

Transitional & Respite Care - Residential Services

Supported Employment – Clubhouses





EXPANDED CAPACITY FOR THE ADULT BEHAVIORAL HEALTH CONTINUUM OF CARE

Forensic Multi- Disciplinary Team	Provide services to individuals determined by the Court to be Incompetent to Proceed (ITP) or Not Guilty by Reason of Insanity (NGI).
FACT	Added a second FACT team to Palm Beach County to serve 100 individuals with serious mental illness successfully reside in the community. The goal is to reduce Baker Acts, divert from state hospitalization, eliminate incarceration, improve social connectedness.
Short Term Residential Treatment (SRT)	An acute 15-bed program serving <u>adults</u> who are high utilizers of acute care and/or are being diverted from the State Mental Health Facilities. Average length of stay 90-120 days
Conditional Release Beds	Expand residential conditional release placements for adults involved in the Criminal Justice System and help to divert individuals from placement in county jails
Respite & Transitional Residential Care	These services address persons experiencing an acute or immediately sub-acute crisis who, in the absence of a suitable alternative, would require hospitalization.
Clubhouse	An EBP that builds on people's strengths and provides mutual support, along with professional staff support for people to receive prevocational work training, educational opportunities, and social support. Locations: West Palm Beach and Belle Glade.

CENTRAL RECEIVING SYSTEM

- RECURRING FUNDS OF \$2.97 MILLION
- EXPANSION OF THE NO WRONG DOOR POLICY
- DEVELOPED IN COLLABORATION WITH NEUROBEHAVIORAL HOSPITAL AT THE 45TH STREET FACILITY ADJACENT TO ST. MARY'S HOSPITAL
- ADDS FUNDING FOR 12 OF 20 CHAIRS FOR CRISIS
 RECEIVING, UP TO 23 HOURS OF PSYCHIATRIC
 ASSESSMENT, PLACEMENT IN THE APPROPRIATE LEVEL OF
 CARE, AND CONNECTION TO TREATMENT
- SOFT LAUNCH DECEMBER 1ST
- FINAL REMODELING TO BE COMPLETED ~APRIL 2024
- GOAL: 8 MINUTE DROP OFF TIME FOR LAW ENFORCEMENT







LAUNCHED 988 WITH ENHANCED MOBILE RESPONSE TEAMS CAPACITY

FAMILY INTENSIVE TREATMENT (FIT) TEAM EXPANSION

Benefits...

• The model provides intensive, community-based services to families in the child welfare system with a parent who has relapse-prone substance misuse.

This funding created ...

• A new FIT team to reduce the placement of children in out of home care, foster care, and residential treatment.

Why is this important?

• An important component of the model is family advocacy across systems - child welfare, judicial, and behavioral health systems.

SEFBHN funds \$1.2 million to serve:

• 120 adults and families (\$10,000 per family)

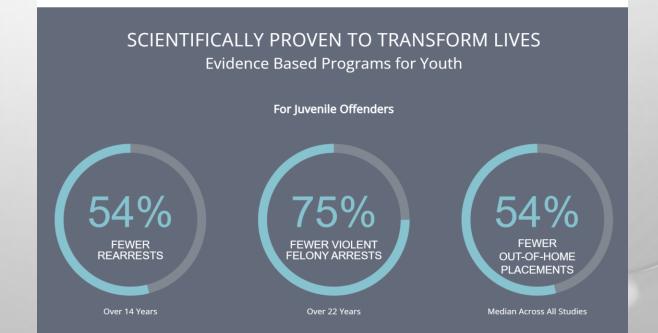
MULTISYSTEMIC THERAPY (MST)

- MULTISYSTEMIC THERAPY (MST) IS AN INTENSIVE, EVIDENCE-BASED TREATMENT THAT EMPOWERS YOUTH (AGED 12-17) AND THEIR FAMILIES TO FUNCTION RESPONSIBLY OVER THE LONG-TERM WITHIN THE COMMUNITY.
- MST REDUCES DELINQUENT AND ANTISOCIAL BEHAVIOR BY ADDRESSING CORE CAUSES OF SUCH CONDUCT.
- CASELOADS ARE SMALL AND SERVICES
 ARE PROVIDED IN THE HOME AT TIMES
 CONVENIENT TO THE FAMILY.
- AVERAGE LENGTH OF TREATMENT IS 3-5 MONTHS.

COMMUNITIES SAW REDUCTION IN



MST IS THE ONLY INTERVENTION FOR HIGH RISK YOUTH WHERE RESULTS HAVE BEEN REPEATEDLY REPLICATED BY INDEPENDENT RESEARCH TEAMS







We support and generate innovative programs and solutions to address the behavioral health needs of our community.

Each year, over 13,000 individuals in Palm Beach County with mental health and substance use disorders are directly assisted by SEFBHN's diverse network of community providers.

They receive critical services like crisis intervention, outpatient counseling, medication assisted treatment (MAT), and much more. As a behavioral health managing entity, we work collaboratively with school systems, law enforcement, and more to effectively direct resources to support and help create programs which serve the unique needs of the communities. Here are just a few of the ways our services touch the lives of those in our community:



Behavioral Health Services for adults, children, and families treat mental health and substance use disorders.

Mobile Response Teams work with 211 helpline to provide support and hope to people experiencing crises.

Peer support specialists engage with victims of overdoses to provide recovery support and connections to treatment.

To learn more, visit us at sefbhn.org by scanning the QR code.





Southeast Florida Behavioral Health Network is a managing entity contracted with the Department of Children and Families. The Florida Department of Children Families is committed to its mission of protecting the vulnerable, promoting strong and economically self-sufficient families, and advancing personal and family recovery and resiliency.





Ann M. Berner, CEO 561-484-5148 Direct Line

Ann Berner@SEFBHN.ORG

QUESTIONS?

December 2023 Feasibility Study Executive Summary

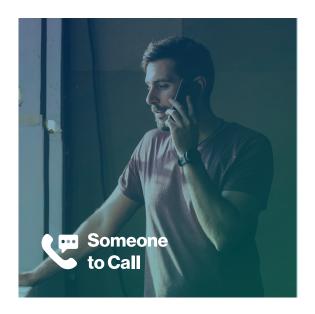
initium

Implementation of the Crisis Now Model in Palm Beach County

The residents of Palm Beach County deserve a high quality, high functioning behavioral health crisis system. The National Guidelines for Behavioral Health Crisis Care delineate best practices for crisis care systems. These guidelines ensure residents receive timely, effective, and compassionate care.



Spearheaded by the National Association of State Mental Health Program Directors, the Crisis Now Model serves as a framework for communities to implement the National Guidelines. The Crisis Now Model is best known for its three key programmatic components:



1. Someone to Call

Regional Crisis Call Centers provide 24/7 access to trained professionals for risk assessment and support through the 988 Suicide & Crisis Lifeline. Best practice guidelines include implementing GPS-enabled technology to dispatch mobile crisis teams to those in need. In the Crisis Now Model, crisis call centers utilize real-time bed registries and have the ability to schedule outpatient appointments and efficiently link people to needed resources. As the hub for services, the crisis call center functions as "air traffic control" for people in crisis, which means call center staff always know where a person is in the continuum of care and facilitate hand-offs and ensure follow-up occurs.

Georgia is widely recognized as a national leader for developing the first statewide crisis line capable of real-time tracking of available crisis beds and utilizing GPS data to dispatch mobile teams. The Georgia Crisis Access Line (GCAL) developed specialized software to support its comprehensive system, which is now made available to other communities in partnership with Behavioral Health Link.

2. Someone to Respond

Mobile Crisis Teams provide community-based intervention services to individuals experiencing a crisis, and offer support wherever the person is. According to the National Guidelines, teams should be staffed by licensed/credentialed clinicians and peers, and should respond within one hour.

Teams should be deployed by the crisis call center on a 24/7 basis. Mobile teams can respond without law enforcement accompaniment unless inclusion is warranted. This approach can lessen the burden of behavioral health crisis response for law enforcement.

The Crisis Now Resource Calculator identifies that Palm Beach County requires 20 mobile teams (each working 40 hours per week) to meet the needs of residents. Today, there are only 3 mobile teams in Palm Beach County.



In Oregon, the Crisis Assistance Helping Out On the Streets (CAHOOTS) program has been widely recognized as a model for mobile crisis and law enforcement collaboration. Appropriate behavioral health calls that come in through 911 are channeled to CAHOOTS teams for response. In 2019, out of nearly 18,000 calls, the CAHOOTS team requested police back up from the City of Eugene just 311 times (2%).

3. Somewhere to Go



Crisis Receiving and Stabilization Facilities provide 24/7 intensive, short-term stabilization in a warm and welcoming environment. As outlined by the National Guidelines, these facilities accept both walk-ins and individuals brought in by first responders.

Crisis receiving facilities provide individual stays of up to 23 hours and 59 minutes, so are often referred to as 23-hour observation units. Short-term stabilization beds for individuals who are not sufficiently stabilized within 24 hours are often co-located with crisis receiving facilities. The Crisis Resource Need Calculator identifies that 82 receiving chairs and 69 short-term beds are needed in Palm Beach County, while currently there are just 20 chairs and 35 short-term beds.

Connections Health Solutions is a nationally-recognized leader in transforming crisis care. Its Crisis Response Center (CRC) in Tucson, Arizona, is distinguished by its unique combination of medical and recovery-focused treatments. The CRC implements best practices, including offering a dedicated first responder drop-off area with a drop-off process of less than 10 minutes and a no rejection policy for first responders. Connections reports that 60-70% of clients achieve stabilization and are successfully discharged back into the community through the 23-hour stabilization service at the CRC.

The Business Case for Evidence-Based Crisis Care

The business case for implementation of the Crisis Now Model is clear. By investing in lower-cost alternatives to care through the expansion and upgrade of crisis call centers, deployment of mobile response teams, and the centralization of specialized facility-based crisis care, communities will save money.

The costly interventions of incarceration and hospitalization will be reduced, and the upstream investments in evidence-based approaches to behavioral health crises will better optimize limited financial resources across the community.



The purpose of this feasibility study is to present recommendations on the crisis care approach best suited for Palm Beach County. While the Crisis Now Model has specific elements that are required for success, there are a variety of ways to implement and operationalize these components. The cost estimates provided below are based on health care costs for Palm Beach County specific needs and service gaps according to the National Guidelines. Projected costs of implementation of the Crisis Now model are based on estimates derived from nationally-recognized services and facilities. Costs are projected without assigning responsibility to a particular community partner for implementation. This report is the starting point for the discussions with stakeholders and community partners on the optimal way to implement the Crisis Now Model in Palm Beach County.

The Crisis Resource Need Calculator provides an overview of the estimated cost reduction associated with transforming the existing crisis care system in Palm Beach County from a starting point focused solely on ED and inpatient psychiatric services (Figure 1).

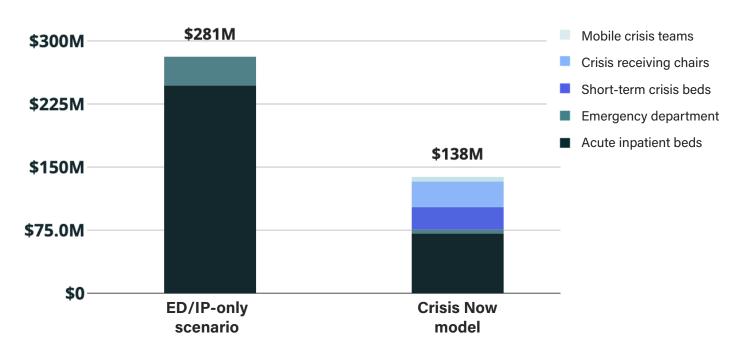


Figure 1. Crisis Resource Need Calculator for Palm Beach County

The ED and inpatient psychiatric services only scenario is a starting point for communities to estimate their cost reduction potential. By implementing the full continuum of Crisis Now services, Palm Beach County can build on its existing crisis services and realize significant savings.

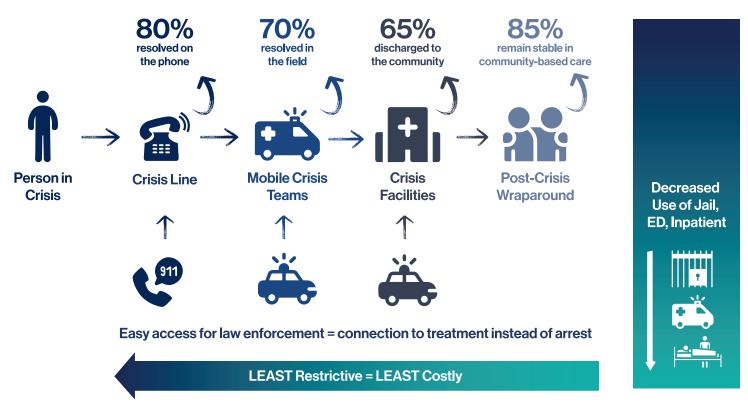
We collected data on the current state of crisis services and compared it to the recommended level of services in the Crisis Now model. By doing so, we identified the gap that needs to be addressed in order to provide a crisis system aligned with the National Guidelines (Table 1).

As shown below, the implementation of a fully equipped crisis system is anticipated to reduce the demand for acute inpatient beds to less than the number available today. This occurs as more crisis care is delivered in alternative settings, and crises are resolved earlier in their progression.

Table 1: Current and Recommended Crisis Service Levels

Site	Current State: Palm Beach County	Crisis Now Model Projections	Gap
Mobile Crisis Teams	3	20	17
Crisis Receiving Chairs	20	82	62
Short-term Crisis Beds	35	69	34
Acute Inpatient Beds	279	186	N/A

A pioneering study cited in the National Guidelines and conducted by the National Action Alliance for Suicide Prevention found that crisis services were the preferred and most efficient care for people in distress, not hospital-based care. An analysis of crisis data in Tucson, Arizona, found that the vast majority of people (80%) had their crisis resolved over the phone when calling a crisis call center, and 70% of those who needed mobile response had their crisis resolved in the community, without the need to visit a crisis facility. Of those who visited a crisis facility, 65% were discharged to levels of care other than inpatient, ED, or jail. Further, 85% of individuals with a mobile crisis team or crisis facility encounter did not have a subsequent ED visit or hospitalization within 45 days. A person-centered crisis system delivers services in the most effective, least restrictive settings, minimizing the use of locked facilities, restraint, force, and seclusion.





As an existing medical provider and a taxpayer funded entity, the Health Care District is well positioned to leverage its resources to facilitate the implementation of the Crisis Now Model in Palm Beach County.

Successful implementation will require the coordination and cooperation of all parties involved in the delivery of behavioral health care in Palm Beach County. The Health Care District is equipped to leverage its electronic health record system to enhance care coordination across various sites of care. The Health Care District can serve as a convener to support accountability and data transparency and ensure sustainability.



Best Practices for a Care Continuum

The National Guidelines for Behavioral Health Crisis Care establish minimum expectations and best practices for each programmatic component of the care continuum.

Crisis Call Center: Minimum Expectations

Minimum Expectations

Operate every moment of every day (24/7/365)

Answer every call or coordinate overflow coverage with a resource that also meets all of the minimum crisis call center expectations

Assess risk of suicide in a manner that meets NSPL standards and danger to others within each call

Coordinate connections to crisis mobile team services in the region

Be staffed with clinicians overseeing clinical triage and other trained team members to respond to all calls received

Connect individuals to facility-based care through warm hand-offs and coordination of transportation as needed

Crisis Call Center: Best Practices

Best Practices (must meet minimum expectations AND):

Incorporate Caller ID functioning

Implement GPS-enabled technology in collaboration with partner crisis mobile teams to more efficiently dispatch care to those in need

Utilize real-time regional bed registry technology to support efficient connection to needed resources

Schedule outpatient follow-up appointments in a manner synonymous with a warm handoff to support connection to ongoing care following a crisis episode

Mobile Crisis Teams: Minimum Expectations

Minimum Expectations

Include a licensed and/or credentialed clinician capable of assessing the needs of individuals within the region of operation

Respond where the person is (home, work, park, etc.) and not restrict services to select locations within the region or particular days/times

Connect individuals to facility-based care as needed through warm hand-offs and coordinating transportation when and only if situations warrant transition to other locations

Mobile Crisis Teams: Best Practices

Best Practices (must meet minimum expectations AND):

Incorporate peers within the mobile crisis team

Respond without law enforcement accompaniment unless special circumstances warrant inclusion in order to support true justice system diversion

Implement real-time GPS technology in partnership with the region's crisis call center hub to support efficient connection to needed resources and tracking of engagement

Schedule outpatient follow-up appointments in a manner synonymous with a warm handoff in order to support connection to ongoing care

Crisis Receiving Facility: Minimum Expectations

Minimum Expectations

Accept all referrals

Not require medical clearance prior to admission but rather assessment and support for medical stability while in the program

Design their services to address mental health and substance use crisis issues

Employ the capacity to assess physical health needs and deliver care for most minor physical health challenges with an identified pathway in order to transfer the individual to more medically staffed services if needed

Be staffed at all times with a multidisciplinary team capable of meeting the needs of individuals experiencing all levels of crisis in the community

Offer walk-in and first responder drop-off options

Be structured in a manner that offers capacity to accept all referrals at least 90% of the time with a no rejection policy for first responders

Screen for suicide risk and complete comprehensive suicide risk assessments and planning when clinically indicated

Screen for violence risk and complete more comprehensive violence risk assessments and planning when clinically indicated

Crisis Receiving Facility: Best Practices

Best Practices (must meet minimum expectations AND):

Function as a 24 hour or less crisis receiving and stabilization facility

Offer a dedicated first responder drop-off area

Incorporate some form of intensive support beds into a partner program to support flow for individuals who need additional support

Include beds within the real-time regional bed registry system operated by the crisis call center hub to support efficient connection to needed resources

Coordinate connection to ongoing care

References

Balfour, M. E., Hahn Stephenson, A., Winsky, J., et al. (2020). Cops, Clinicians, or Both? Collaborative Approaches to Responding to Behavioral Health Emergencies. Alexandria, VA: National Association of State Mental Health Program Directors. Retrieved from https://www.nasmhpd.org/sites/default/files/2020paper11.pdf

Behavioral Health Link. https://behavioralhealthlink.com/crisis-services/georgia-crisis-and-access-line/ Accessed on 12/5/2023.

Connections Health Solutions. https://connectionshs.com/ Accessed on 12/5/2023.

Crisis Resource Need Calculator. https://calculator.crisisnow.com/Accessed on 11/8/2023.

Eugene Police Department Crime Analysis Unit. CAHOOTS Program Analysis. 2020. Retrieved from https://www.eugene-or.gov/DocumentCenter/View/56717/CAHOOTS-Program-Analysis

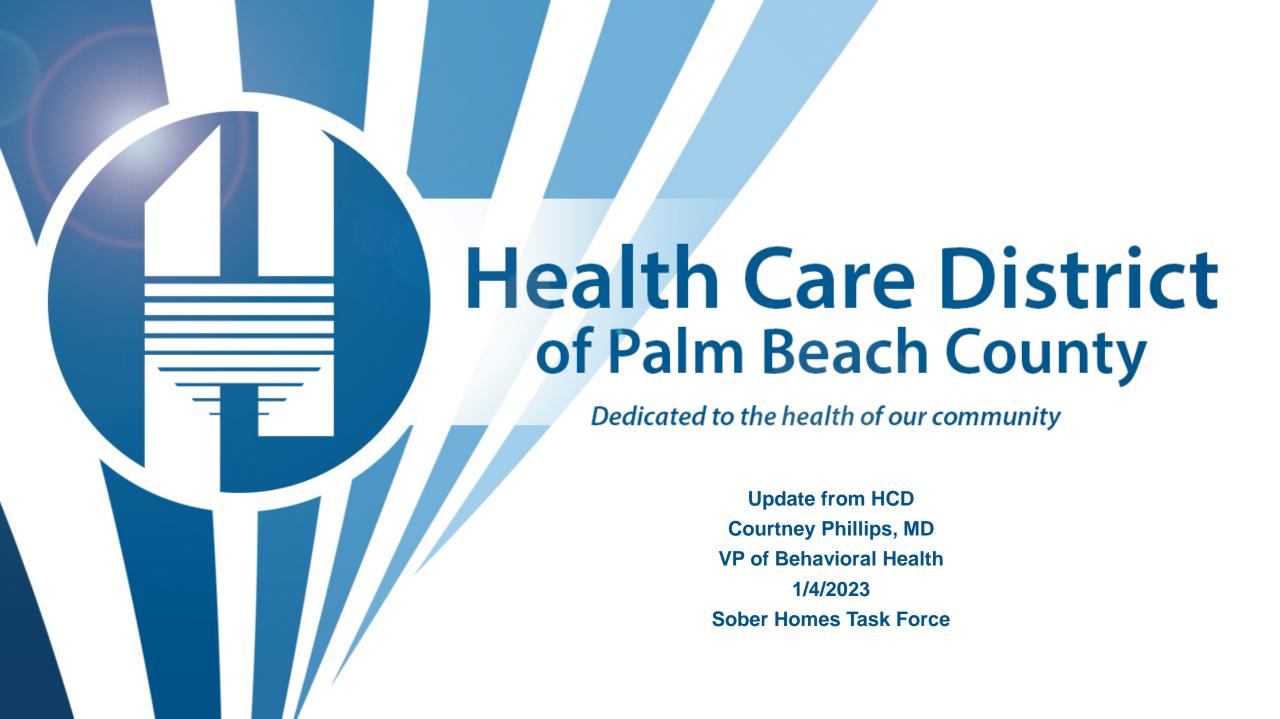
Georgia Crisis and Access Line. The Georgia Collaborative ACO. https://www.georgiacollaborative.com/providers/georgia-crisis-and-access-line-gcal/. Accessed on 12/5/2023.

Georgia Department of Behavioral Health and Developmental Disabilities. The Crisis System of Georgia | Georgia Department of Behavioral Health and Developmental Disabilities. Accessed on 12/5/2023.

Hepburn, S. (2021). Georgia's Crisis System Transformation and Lessons Learned in Anticipation of 988. Crisis Talk. https://talk.crisisnow.com/georgias-crisis-system-transformation-and-lessons-learned-in-anticipation-of-988/

National Action Alliance for Suicide Prevention: Crisis Services Task Force. (2016). Crisis now: Transforming services is within our reach. Washington, DC: Education Development Center, Inc.

Substance Abuse and Mental Health Services Administration. (2020). National Guidelines for Behavioral Health Crisis Care Best Practice Toolkit. Substance Abuse and Mental Health Services Administration. Retrieved from https://www.samhsa.gov/data/





Outline

- Current service update
 - SUD services
 - BH Integrated services
 - BH/Psych Services
- Future plans/collaboration for Crisis Now





SUD services

- Added Delray Beach site October 1 to offer same services as Mangonia Park
 - SUD care anytime with a lot of walk in capacity
 - Primary care
 - Psychiatry
 - Group therapy
- Have a lot of capacity
- New 24-7 Mangonia Park clinic
- SEFBHEN collaboration
- Fully using injectable buprenorphine through insurance and grant.





Primary Care Behavioral Health Integration Update

- Working on developing pediatric integration in 1 pediatric clinic
- Added Integrated BHC in primary care in Belle Glade to assess behavioral health...All clinics except Boca and Jupiter have bhc in adult primary care
- Working on adding more individual and group therapy





Psychiatry update

- FAU psychiatry residency is with the District at Delray and Mangonia 4 days per week
- We have a Psychiatric APRN at Lantana full time and West Palm Beach clinic full time seeing teens to adults (both trained in substance use as well)
- We have a LOT of capacity to see more psychiatric patients, please send by calling 561-642-1000. Many patients can get appointments within a week right now.
- Working on getting injectable antipsychotics as a regular practice

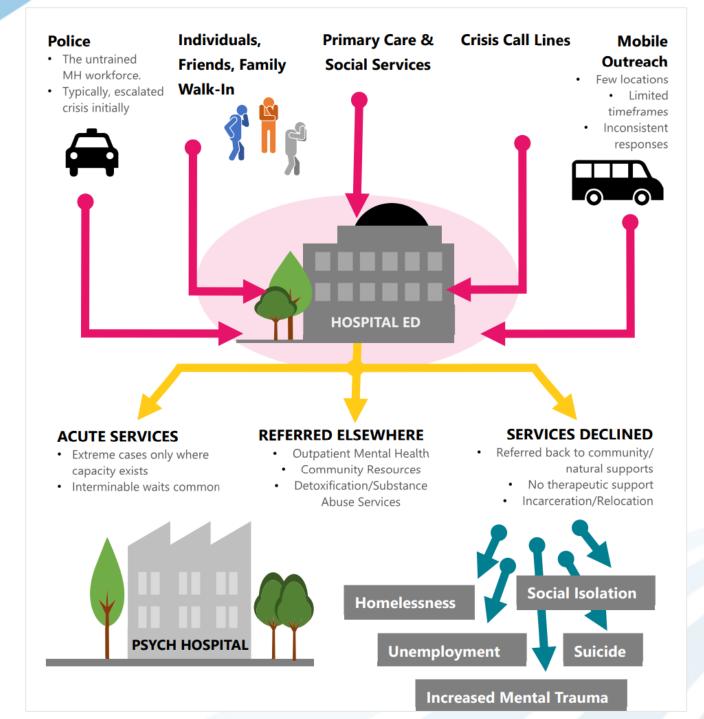




Future plan collaborating and assisting in coordinating a "Crisis Now" Model for Palm Beach County



Traditional Crisis Flow







HCD BEHAVIORAL HEALTH INITIATIVE

- Emerged from discussions with Palm Beach County Administration that started pre-COVID, paused during the COVID response, and restarted with greater urgency upon COVID stabilization.
- Need for evidence based, real-world-tested best practices guidance to the BH field (same as physical health)

SAMHSA National Guidelines for Behavioral Health Crisis Care

U.S Department of Health and Human Services agency that leads public health efforts to advance BH; The mission is to reduce the impact of Substance Abuse and Mental Illness on American communities.

Initium Health – Public Benefit Corp (Denver, CO)

 Assess and provide potential ownership and governance models for a new system of care based on SAMHSA guidelines



Four Core Elements For Transforming Crisis Services



High-Tech Crisis Call Centers

These programs use technology for real-time coordination across a system of care and leverage big data for performance improvement and accountability across systems. At the time, they provide high-touch support to individuals and families in crisis.



24/7 **Mobile Crisis**

Mobile crisis offers outreach and support where people in crisis are. Programs should include contractually required response times and medical backup.



Crisis Stabilization Programs

These programs offer short-term "sub-acute" care for individuals who need support and observation, but not ED holds or medical inpatient stay, at lower costs and without the overhead of hospital-based acute care.



Essential Principles and **Practices**

These must include a recovery orientation, trauma-informed care, significant use of peer staff, a commitment to Zero Suicide/Suicide Safer Care, strong commitments to safety for consumers ans staff, and collaboration with lay enforcement.



Crisis Care Continuum



988 is the preferred, nowrong-door place to access crisis care Mobile response teams available 24/7

Crisis receiving and stabilization facilities



Crisis Care Continuum



Decreased
Use of
Jail, ED,
Inpatient



Proposed Mental Health Model for Palm Beach County



REHABILITATIVE CARE





HOUSING

- . Congregate or Individual
- Family-based
- Support that provides in-home care
- Transitional Housing

PEOPLE

SOCIAL CONNECTION

- Peer support
- Clubhouses

Entire Continuum of Care in Single System

Centralized Case
Management and Care
Coordination

- Online community
- Recovery Centers



PURPOSE

EDUCATIONAL AND EMPLOYMENT SUPPORT

(Restoring a sense of purpose and relevance)



Inpatient/ED versus Crisis Now Model

Projected costs of ED/IP system vs Crisis Now Model by Site of Service

Site	ED/IP	Crisis Now			
Mobile Crisis Teams	\$O	\$5.5M			
Crisis Receiving Chairs	\$O	\$30.4M			
Short-term Crisis Beds	\$0	\$26.6M			
Emergency Department	\$34.2M	\$4.8M			
Acute Inpatient Beds	\$247M	\$71.0M			
TOTAL	\$281M	\$138M			

Crisis Now reduces costs through better allocation of resources, resolving more crises in the community without the need for emergency department or acute inpatient care. (Keep in mind that we have some of these services already in PBC that might need to be coordinated or aligned with SAMHSA model)

Where are we now with execution?

- <u>Initium</u> completed feasibility study, has full draft that will be finalized based on the community meeting from December 19 as well as the follow up from the meeting. (Executive summary is out).
- HCD has committed to make Mangonia Park a 24-7 designated clinic in its current site for substance use crisis and mental health crisis after conversations with West Palm Beach Police Department. Aiming for q2/q3 implementation. This will help hold patients diverted/cleared from ASU as well as give crisis MAT to do a 24-7 warm handoff and attempt to prevent baker act.
- HCD is exploring collaboration with many county partners currently as well as taking a lead to helping elevate standards to SAMHSA guidelines for different aspects of this system.
 - Currently in conversation with PBSO to collaborate on Mobile Crisis Units





Thank you@

Questions?



Homelessness, Addiction & Mental Illness

presented by Detective Jennifer Jones Rivera Beach Police Department

Partners in Care Program (Police Department)

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The Partners in Care program strives to improve access to care for individuals experiencing a behavioral health crisis, divert consumers from the criminal justice system and into the health care system, all while improving the communication and coordination across all systems of care, and increasing the safety of all involved in these crisis interactions.

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Each clinician who is paired with an officer receives 16 hours of job specific training to prepare them for their role in this co-response program.

https://www.nashville.gov/departments/police/investigative-services/alternative-policing-strategies/partners-care



MNPD Crisis Intervention Team Weekly Report Week Ending 03/11/2023 South Precinct

	Prior Week	Last Week	% Change	Prior 4 Weeks	Last 4 Weeks	% Change	Prior YTD	YTD	% Change	Since Inception	
CIT Incidents											
All CIT Incidents	41	58	41.5%	171	212	24.0%	0	491	-	855	
Meeting Criteria for CIT Response	11	17	54.5%	55	59	7.3%	0	138	-	234	
How Officer was Notified											
DEC Dispatched	9	14	55.6%	41	46	12.2%	0	104	-	172	
CIT Requested by Other Officer	2	3	50.0%	12	12	0.0%	0	31		48	
CIT Initiated	0	0		2	1	-50.0%	0	3		31	
Follow-up	0	0		0	0		0	0		3	
Time Spent on CIT Responses											
Total Duration in Hours	10	17	70.0%	215	135	-37.2%	0	435	-	756	
Officer Activites											
Subject Could be Charged w Crime	2	2	0.0%	10	9	-10.0%	0	24		35	
Subject Arrested	0	0	-	3	1	-66.7%	0	6	-	8	
Force Used	0	0	-	3	0	-100.0%	0	5	-	6	
Subject Armed	0	0	-	5	0	-100.0%	0	6	740	10	
Subject Injured	0	0		.4	4	0.0%	0	9		18	
Subject Injured Prior to Contact	0	0	-	4	3	-25.0%	0	8	-	17	
Officer Injured	0	0	-	0	0	*	0	0	+	0	
Officer Injured Prior to Contact	0	0		0	0		0	0		0	
Other Party Injured	0	0	-	0	1	100	0	1	-	1	
Other Party Injured Prior to Contact	0	0	-	0	1		0	1	-	1	
Responses by Officer Assignment											
Central	0	0	-	0	0		0	0		0	
East	0	0	-	0	0		0	0		0	
Hermitage	0	0		0	0	-	0	0		0	
Madison	0	0		0	0	-	0	0	-	0	
Midtown Hills	0	0	-	0	0	-	0	٥		0	
North	0	0	-	0	0	-	0	0	-	0	
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Other Division	0	0		0	0		0	0	-	0	

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In 1999, MHC recognized the need to support the whole family and expanded services to include children and youth.

https://www.mhc-tn.org

States with the most homelessness





Risk vs Protective Factors

RISK FACTORS

► Aggressive behavior in childhood ► Belief in self control (self esteem)

PROTECTIVE FACTORS

- ► Lack of parental supervision
- ► Low peer refusal skills
- Drug experimentation
- Availability of drugs
- Community poverty

- Parental support
- Positive relationships
- Goals
- Healthy coping skills
- Neighborhood resources

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Discussion

- What are the challenges we are facing with housing in Palm Beach County?
- What challenges have you or someone you know faced in finding housing for yourself?
- ► How might these challenges impact those who are unhoused?
- What success have you had with serving individuals who are unhoused?
- What failures have you seen?
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- What is the prevalence of severe mental illness/substance abuse? Why do you think that is?
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- 1 Section 3. subsection (20) is added to section 397.321, Florida
- 2 Statutes, to read:
- 3 397.321 Duties of the department. The department shall:
- 4 (20) Prominently display and make available on its website No no
- 5 later than January 1, 2025, all information, including but not
- 6 limited to information contained in the department's Provider
- 7 Licensure and designations System, pertaining to the following:
- 8 (a) Service provider applications for licensure and license9 renewal.
- (b) Policies and procedures provided to the department by anapplicant for service provider licensure or license renewal.
- 12 (c) The name and location of each recovery residence engaged 13 in a referral relationship with a licensed service provider or
- 14 service provider applicant, as required under ss. 397.4104, and
- **15** 397.403(1)(j).
- 16 (c) A licensed service provider's organizational chart
- 17 identifying medical, clinical, managerial and operational
- 18 positions.
- 19 (d) All complaints, investigative reports and findings
- 20 pertaining to service providers received by the department that
- 21 result in a violation classified under ss. 397.411(7), 397.4104(2)
- 22 or 397.4873(6). Complainants' names and other identifying
- 23 information shall be redacted.
- 24 (e) Fines assessed for violations pursuant to ss.
- 25 397.411(7), 397.4104(2) or 397.4873(6), Florida Statutes.
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- 27 license suspension or revocation.
- 28 (g) All inspection reports for service provider licenses.

29

Page **1** of **2**

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Page 1 of 2

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Page **2** of **2**

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1/4/24, 10:42 AM AHCA: Document Results



Search	
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Search Page > Provider Results

Click 'Select' to display a document.

Export Results
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Inspection Details for This Provider
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FloridaHealthFinder Profile

Search Criteria Selected:

Provider Name: ABBEY DELRAY Provider Type: Nursing Home

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Please check your browser popup blocker settings if you have trouble viewing documents.

The Statement of Deficiencies Public Record Search displays a complete list of inspections. Documents on this page are redacted per 45 Code of Federal Regulations (CFR) 164.514 through the use of an automated redaction software, which may over-redact to protect from the potential release of confidential information. Manually redacted documents can be obtained by contacting the Public Records Office at PublicRecordsReq@ahca.myflorida.com.

Users will be directed to the federal Nursing Home Compare website at www.medicare.gov/care-compare for nursing home standard and complaint inspections with deficiencies cited that were conducted within the last three years.

1 2 3 4 5	<u>678</u>				
	<u>Inspection Type</u>	<u>Document Type</u>	<u>Visit Date</u>	<u>Pages</u>	Inspection Status

<u>Select</u>	Complaint	Statement of Deficiencies	12/06/2023	2	No Deficiencies
<u>Select</u>	Complaint	Statement of Deficiencies	08/02/2023	2	No Deficiencies
<u>Select</u>	Complaint	Statement of Deficiencies	08/02/2023	2	Deficiencies Corrected
Select	Complaint	Statement of Deficiencies	06/21/2023	15	Deficiencies Cited
<u>Select</u>	Standard	Statement of Deficiencies	04/21/2023	2	Deficiencies Corrected
<u>Select</u>	Fire/Life/Safety	Statement of Deficiencies	04/21/2023	2	Deficiencies Corrected
<u>Select</u>	Complaint	Statement of Deficiencies	04/21/2023	2	No Deficiencies
<u>Select</u>	Standard	Statement of Deficiencies	03/17/2023	54	Deficiencies Cited
<u>Select</u>	Fire/Life/Safety	Statement of Deficiencies	03/14/2023	13	Deficiencies Cited
<u>Select</u>	Complaint	Statement of Deficiencies	02/08/2023	2	No Deficiencies
12345	5 6 7 8				•

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Medicaid Licensure & Regulation

Find a Facility

Report Fraud









Contact Us

(888) 419-3456 (800) 955-8771 (TDD)



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Select	Complaint	Statement of Deficiencies	04/21/2023	2	No Deficiencies
Select	Standard	Statement of Deficiencies	03/17/2023	54	Deficiencies Cited
Select	Fire/Life/Safety	Statement of Deficiencies	03/14/2023	13	Deficiencies Cited
Select	Complaint	Statement of Deficiencies	02/08/2023	2	No Deficiencies

Return to Provider Results

PRINTED: 12/12/2023

STATEMENT	r Health Care Admini of DEFICIENCIES CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CO A. BUILDING:	INSTRUCTION	(X3) DATE 8 COMPL		
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NAME OF PR	OVIDER OR SUPPLIER	STREET	ADDRESS, CITY, STATE,	ZIP CODE			
ABBEY DE	LRAY		V 11TH COURT V BEACH, FL 33445				
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ICA Form 30:	20-0001						
		USUPPLIER REPRESENTATIVE'S SIGNATU	RE	TITLE		(X6) DATE	
	ally Signed					12/11/23	

STATEMENT OF DEFICIENCIES (XI) PROVIDER AND PLAN OF CORRECTION (DENTIFICATION NUMBER 108335 **NING** **INSTATEMENT OF DEFICIENCIES NAME OF PROVIDER OR SUPPLIER **ABBEY DELRAY* **STREET ADDRESS, CITY, STATE, ZIP CODE **2105 BY 11TH COURT **DELRAY BEACH, FL. 33445 **OPENING SUMMARY STATEMENT OF DEFIDENCIES **PREFIX **TAG** (RACH DEFIDENCY MUST 8E REFECEDED BY FULL **PREFIX **PREFIX **PREFIX **PREFIX **TAG** **INITIAL COMMENTS An unannounced Complaint survey, Complaint **RZ023011781, Complaint #*Z023013264, Complaint #*Z023013464, Complaint **RZ0230134640 was conducted on 1200623 at Abbey Delray. The facility is in compliance with 42 CFR Part 483, Requirements for Long Term Gare Facilities. **ABBEY DELCAY:** **PREFIX **P			ID HUMAN SERVICES MEDICAID SERVICES				FOR	D: 12/12/2023 M APPROVED D: 0938-0391
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		#2023011781, Comp Complaint #2023013- #2023014604 was co Abbey Delray. The fa CFR Part 483, Requi	laint #2023013264, 484, Complaint Inducted on 12/06/23 at cility is in compliance with 42					
Electropically Signed 12/11/2022			SUPPLIER REPRESENTATIVE'S SIGNATUR	RE		TITLE		

Executoritically Signed

Any deficiency statement ending with an asterisk (") denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguarist provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

FORM CMS-2567(02-99) Previous Versions Obsolete Event ID: V1LM11 Facility ID: 95051

If continuation sheet Page 1 of 1

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A bill to be entitled An act relating to substance abuse treatment; amending s. 212.02, F.S.; eliminating certain tax liabilities imposed on certified recovery residences; amending s. 397.311, F.S.; providing the levels of care at certified recovery residences and their respective levels of care for residents; amending s. 397.321, F.S.; requiring the Department of Children and Families to display and make available on its website certain information pertaining to service providers and recovery residences by a specified date; requiring the department to display on its website certain documents pertaining to service providers; amending s. 397.335, F.S.; revising the membership of the Statewide Council on Opioid Abatement to include additional members; amending s. 397.487, F.S.; extending the deadline for certified recovery residences to retain a replacement for a certified recovery residence administrator who has been removed from his or her position; authorizing, rather than requiring, the credentialing entity to revoke the certificate of compliance if a certified recovery residence fails to meet specified standards; requiring certified recovery residences to remove certain individuals from their positions if they are arrested

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and awaiting disposition for, are found guilty of, or enter a plea of quilty or nolo contendere to certain offenses, regardless if adjudication is withheld; requiring the certified recovery residence to retain a certified recovery residence administrator if the previous certified recovery residence administrator has been removed due to any reason; prohibiting certified recovery residences, on or after a specified date, from denying an individual access to housing solely for being prescribed federally approved medications from licensed health care professionals; prohibiting local laws, ordinances, or regulations adopted on or after a specified date from regulating the duration or frequency of a resident's stay in a certified recovery residence in certain zoning districts; providing applicability; amending s. 397.4871, F.S.; authorizing, rather than requiring, credentialing entities to revoke a certificate of compliance if a recovery residence fails to meet specified standards; authorizing certain Level IV certified recovery residences owned or controlled by a licensed service provider and managed by a certified recovery residence administrator approved for a specified number of residents to manage a specified greater number of residents, provided that certain

Page 2 of 13

criteria are met; prohibiting a certified recovery residence administrator who has been removed by a certified recovery residence from taking on certain other management positions without approval from a credentialing entity; defines the term "community housing"; providing an effective date.

Be It Enacted by the Legislature of the State of Florida:

Section 1. Paragraph (k) is added to subsection (10) of section 212.02, Florida Statutes, to read:

212.02 Definitions.—The following terms and phrases when used in this chapter have the meanings ascribed to them in this section, except where the context clearly indicates a different meaning:

(10) "Lease," "let," or "rental" means leasing or renting of living quarters or sleeping or housekeeping accommodations in hotels, apartment houses, roominghouses, tourist or trailer camps and real property, the same being defined as follows:

(k) For purposes of this chapter, recovery residences certified pursuant to s. 397.487 which rent properties are not subject to any taxes imposed on transient accommodations, including taxes imposed under s. 212.03; any locally imposed discretionary sales surtax or any convention development tax imposed under s. 212.0305; any tourist development tax imposed

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76 <u>under s. 125.0104; or any tourist impact tax imposed under s.</u>
77 125.0108.

Section 2. Subsection (5) of section 397.311, Florida Statutes, is amended to read:

397.311 Definitions.—As used in this chapter, except part VIII, the term:

- (5) "Certified recovery residence" means a recovery residence that holds a valid certificate of compliance and is actively managed by a certified recovery residence administrator. The levels of care within a certified recovery residence are as follows:
- (a) Level I recovery residences that house individuals in recovery who are post-treatment, with a minimum of 9 months of sobriety. Level I certified homes are democratically run by the members who reside in the home.
- (b) Level II recovery residences encompass the traditional perspectives of sober living homes. There is oversight from a house manager with lived experience, typically a senior resident. Residents are expected to follow rules outlined in a resident handbook, pay dues, if applicable, and work toward achieving milestones within a chosen recovery path.
- (c) Level III recovery residences offer higher supervision by staff with formal training to ensure resident accountability.

 These homes offer peer-support services and are staffed 24 hours a day. Clinical services are not performed at the residence. The

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101	services offered may include, but are not limited to, life skill
102	mentoring, recovery planning, and meal preparation. This support
103	structure is most appropriate for residents who require a more
104	structured environment during early recovery from addiction.
105	(d) A Level IV certified recovery residence are dwellings
106	offered, referred to, or provided by, a licensed service
107	provider to its patients who are required to reside at the
108	residence while receiving intensive outpatient and higher levels
109	of outpatient care. Level IV recovery residences are staffed 24
110	hours a day and combine outpatient licensable services with
111	recovery residential living. Residents are required to follow a
112	treatment plan, attend group and individual sessions, in
113	addition to developing a recovery plan within the social model
114	of recovery spectrum. No clinical services are provided at the
115	residence and all licensable services are provided off-site.
116	Section 3. Subsection (20) is added to section 397.321,
117	Florida Statutes, to read:
118	397.321 Duties of the department.—The department shall:
119	(20) Prominently display and make available on its website
120	no later than January 1, 2025, all documents in the department's
121	Provider Licensure and Designations System pertaining to the
122	<pre>following:</pre>
123	(a) Service provider applications for licensure and
124	<u>license renewal.</u>
125	(b) Policies and procedures provided by the department to

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126	an applicant for service provider licensure or license renewal.
127	(c) The name and location of each recovery residence
128	engaged in a referral relationship with a licensed service
129	provider or service provider applicant, as required under ss.
130	397.4104 and 397.403(1)(j).
131	(d) All complaints pertaining to service providers
132	received by the department, and all investigative reports and
133	findings, whether founded or unfounded. Complainant names and
134	other identifying information shall be redacted.
135	(e) Fines assessed for violations pursuant to ss.
136	397.411(7), 397.4104(2), and 397.4873(7).
137	(f) All reports or other documentation pertaining to
138	service provider license suspension or revocation.
139	(g) All inspection reports for service provider licenses
140	and recovery residences.
141	Section 4. Paragraph (a) of subsection (2) of section
142	397.335, Florida Statutes, is amended to read:
143	397.335 Statewide Council on Opioid Abatement
144	(2) MEMBERSHIP.—
145	(a) Notwithstanding s. 20.052, the council shall be
146	composed of the following members:
147	1. The Attorney General, or his or her designee, who shall
148	serve as chair.
149	2. The secretary of the department, or his or her
150	designee, who shall serve as vice chair.

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CODING: Words $\frac{\text{stricken}}{\text{stricken}}$ are deletions; words $\frac{\text{underlined}}{\text{ore additions}}$.

3. One member appointed by the Governor.

- 4. One member appointed by the President of the Senate.
- 5. One member appointed by the Speaker of the House of Representatives.
 - 6. Two members appointed by the Florida League of Cities who are commissioners or mayors of municipalities. One member shall be from a municipality with a population of fewer than 50,000 people.
 - 7. Two members appointed by or through the Florida Association of Counties who are county commissioners or mayors. One member shall be appointed from a county with a population of fewer than 200,000, and one member shall be appointed from a county with a population of more than 200,000.
 - 8. One member who is either a county commissioner or county mayor appointed by the Florida Association of Counties or who is a commissioner or mayor of a municipality appointed by the Florida League of Cities. The Florida Association of Counties shall appoint such member for the initial term, and future appointments must alternate between a member appointed by the Florida League of Cities and a member appointed by the Florida Association of Counties.
 - 9. Two members appointed by or through the State Surgeon

 General. One shall be a staff member from the department who has

 experience coordinating state and local efforts to abate the

 opioid epidemic, and one shall be a licensed physician who is

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- 177 <u>10. One member appointed by the Florida Association of</u> 178 Recovery Residences.
 - $\underline{\mbox{11.}}$ One member appointed by the Florida Association of EMS Medical Directors.
 - 12. One member appointed by the Florida Society of
 Addiction Medicine who is a medical doctor board certified in
 addiction medicine.
 - 13. One member appointed by the Florida Behavioral Health Association.
 - 14. One member appointed by Floridians for Recovery.
 - $\underline{\mbox{15.}}$ One member appointed by the Florida Certification Board.
 - Section 5. Present paragraphs (c), (d), and (e) of subsection (8) of section 397.487, Florida Statutes, are redesignated as subsections (d), (e), and (f), respectively, and amended, a new paragraph (c) is added to that subsection, subsections (13) and (14) are added to that section, and paragraph (b) of subsection (8) of that section is amended, to read:
 - 397.487 Voluntary certification of recovery residences.-
 - (8) Onsite followup monitoring of a certified recovery residence may be conducted by the credentialing entity to determine continuing compliance with certification requirements. The credentialing entity shall inspect each certified recovery

Page 8 of 13

residence at least annually to ensure compliance.

- (b) A certified recovery residence must notify the credentialing entity within 3 business days after the removal of the recovery residence's certified recovery residence administrator due to termination, resignation, or any other reason. The <u>certified</u> recovery residence has <u>90</u> 30 days to retain a certified recovery residence administrator. The credentialing entity shall revoke the certificate of compliance of any <u>certified</u> recovery residence that fails to comply with this paragraph.
- (c) If a certified recovery residence's administrator has been removed due to termination, resignation, or any other reason and had been previously approved to actively manage more than 50 residents pursuant to s. 397.4871(8)(b), the certified recovery residence has 90 days to retain another certified recovery residence administrator pursuant to that section. The credentialing entity shall revoke the certificate of compliance of any certified recovery residence that fails to comply with this paragraph.
- (d) (e) If any owner, director, or chief financial officer of a certified recovery residence is arrested and awaiting disposition for or found guilty of, or enters a plea of guilty or nolo contendere to, regardless of whether adjudication is withheld, any offense listed in s. 435.04(2) while acting in that capacity, the certified recovery residence must shall

immediately remove the person from that position and shall notify the credentialing entity within 3 business days after such removal. The credentialing entity shall revoke the certificate of compliance of a <u>certified</u> recovery residence that fails to meet these requirements.

- (e)(d) A credentialing entity shall revoke a <u>certified</u> recovery residence's certificate of compliance if the <u>certified</u> recovery residence provides false or misleading information to the credentialing entity at any time.
- (f)(e) Any decision by a department-recognized credentialing entity to deny, revoke, or suspend a certification, or otherwise impose sanctions on a certified recovery residence, is reviewable by the department. Upon receiving an adverse determination, the certified recovery residence may request an administrative hearing pursuant to ss. 120.569 and 120.57(1) within 30 days after completing any appeals process offered by the credentialing entity or the department, as applicable.
- (13) Effective January 1, 2025, a recovery residence may not deny an individual access to housing solely on the basis that he or she has been prescribed federally approved medication that assists with treatment for substance use disorders by a licensed physician, a physician's assistant, or an advanced practice registered nurse registered under s. 464.0123.
 - (14) A local law, ordinance, or regulation may not

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regulate the duration or frequency of a resident's stay in a certified recovery residence located within a multifamily zoning district. This subsection does not apply to any local law, ordinance, or regulation adopted on or before February 1, 2024.

Section 6. Paragraphs (b) and (c) of subsection (6) of section 397.4871, Florida Statutes, are amended, and paragraph (c) is added to subsection (8) of that section, to read:

397.4871 Recovery residence administrator certification.

- (6) The credentialing entity shall issue a certificate of compliance upon approval of a person's application. The certification shall automatically terminate 1 year after issuance if not renewed.
- (b) If a certified recovery residence administrator of a recovery residence is arrested and awaiting disposition for or found guilty of, or enters a plea of guilty or nolo contendere to, regardless of whether adjudication is withheld, any offense listed in s. 435.04(2) while acting in that capacity, the certified recovery residence must shall immediately remove the person from that position and shall notify the credentialing entity within 3 business days after such removal. The certified recovery residence shall have 30 days to retain a certified recovery residence administrator within 90 days after such removal. The credentialing entity shall revoke the certificate of compliance of any recovery residence that fails to meet these requirements.

(c) A credentialing entity <u>may shall</u> revoke a <u>certified</u> recovery residence administrator's certificate of compliance if the recovery residence administrator provides false or misleading information to the credentialing entity at any time.

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(c) Notwithstanding paragraph (b), a Level IV certified recovery residence with a community housing component, which residence is actively managed by a certified recovery residence administrator approved for 100 residents under this section and is wholly owned or controlled by a licensed service provider, may actively manage up to 150 residents so long as the licensed service provider maintains a service provider personnel-topatient ratio of 1 to 8 and maintains onsite supervision at the residences 24 hours a day, 7 days a week, with a personnel-toresident ratio of 1 to 10. A certified recovery residence administrator who has been removed by a certified recovery residence due to termination, resignation, or any other reason may not continue to actively manage more than 50 residents for another service provider or certified recovery residence without being approved by the credentialing entity. For purposes of this paragraph, the term "community housing" means a certified recovery residence offered, referred to, or provided by, a licensed service provider that provides housing to its patients who are required to reside at the residence while receiving intensive outpatient and higher levels of outpatient care. A

HB 1065 2024

301	certified recovery residence as defined in s. 397.311(5) used by
302	a licensed service provider that meets the definition of
303	community housing shall be classified as a Level IV level of
304	support,.
305	Section 7. This act shall take effect July 1, 2024.

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By Senator Harrell

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A bill to be entitled

An act relating to substance abuse treatment; amending s. 212.02, F.S.; eliminating certain tax liabilities imposed on certified recovery residences; amending s. 397.311, F.S.; providing the levels of care at certified recovery residences and their respective levels of care for residents; defining the term "community housing"; amending s. 397.321, F.S.; requiring the Department of Children and Families to display and make available on its website certain information pertaining to service providers and recovery residences by a specified date; requiring the department to display on its website certain documents pertaining to service providers; amending s. 397.335, F.S.; revising the membership of the Statewide Council on Opioid Abatement to include additional members; amending s. 397.487, F.S.; extending the deadline for certified recovery residences to retain a replacement for a certified recovery residence administrator who has been removed from his or her position; requiring certified recovery residences to remove certain individuals from their positions if they are arrested and awaiting disposition for, are found guilty of, or enter a plea of quilty or nolo contendere to certain offenses, regardless if adjudication is withheld; requiring the certified recovery residence to retain a certified recovery residence administrator if the previous certified recovery residence administrator has been removed due to any reason; conforming

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provisions to changes made by the act; prohibiting certified recovery residences, on or after a specified date, from denying an individual access to housing solely for being prescribed federally approved medications from licensed health care professionals; prohibiting local laws, ordinances, or regulations adopted on or after a specified date from regulating the duration or frequency of a resident's stay in a certified recovery residence in certain zoning districts; providing applicability; amending s. 397.4871, F.S.; conforming provisions to changes made by the act; authorizing certain Level IV certified recovery residences owned or controlled by a licensed service provider and managed by a certified recovery residence administrator approved for a specified number of residents to manage a specified greater number of residents, provided that certain criteria are met; prohibiting a certified recovery residence administrator who has been removed by a certified recovery residence from taking on certain other management positions without approval from a credentialing entity; providing an effective date.

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Be It Enacted by the Legislature of the State of Florida:

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Section 1. Paragraph (k) is added to subsection (10) of section 212.02, Florida Statutes, to read:

212.02 Definitions.—The following terms and phrases when used in this chapter have the meanings ascribed to them in this

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section, except where the context clearly indicates a different meaning:

- (10) "Lease," "let," or "rental" means leasing or renting of living quarters or sleeping or housekeeping accommodations in hotels, apartment houses, roominghouses, tourist or trailer camps and real property, the same being defined as follows:
- (k) For purposes of this chapter, recovery residences certified pursuant to s. 397.487 which rent properties are not subject to any taxes imposed on transient accommodations, including taxes imposed under s. 212.03; any locally imposed discretionary sales surtax or any convention development tax imposed under s. 212.0305; any tourist development tax imposed under s. 125.0104; or any tourist impact tax imposed under s. 125.0108.

Section 2. Present subsections (9) through (50) of section 397.311, Florida Statutes, are redesignated as subsections (10) through (51), respectively, a new subsection (9) is added to that section, and subsection (5) of that section is amended, to read:

397.311 Definitions.—As used in this chapter, except part VIII, the term:

- (5) "Certified recovery residence" means a recovery residence that holds a valid certificate of compliance and is actively managed by a certified recovery residence administrator.
- (a) A Level I certified recovery residence houses individuals in recovery who have completed treatment, with a minimum of 9 months of sobriety. A Level I certified recovery residence is democratically run by the members who reside in the

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home.

(b) A Level II certified recovery residence encompasses the traditional perspectives of sober living homes. There is oversight from a house manager who has experience with living in recovery. Residents are expected to follow rules outlined in a resident handbook, which is provided by the certified recovery residence administrator. Residents must pay dues, if applicable, and work toward achieving realistic and defined milestones within a chosen recovery path.

- (c) A Level III certified recovery residence offers higher supervision by staff with formal training to ensure resident accountability. Such residences are staffed 24 hours a day, 7 days a week, and offer residents peer-support services, which may include, but are not limited to, life skill mentoring, recovery planning, and meal preparation. No clinical services are performed at the residence. Such residences are most appropriate for persons who require a more structured environment during early recovery from addiction.
- (d) A Level IV certified recovery residence is a residence offered, referred to, or provided by, a licensed service provider to its patients who are required to reside at the residence while receiving intensive outpatient and higher levels of outpatient care. Such residences are staffed 24 hours a day and combine outpatient licensable services with recovery residential living. Residents are required to follow a treatment plan and attend group and individual sessions, in addition to developing a recovery plan within the social model of living a sober lifestyle. No clinical services are provided at the residence, and all licensable services are provided off-site.

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(9) "Community housing" means a certified recovery residence offered, referred to, or provided by a licensed service provider that provides housing to its patients who are required to reside at the residence while receiving intensive outpatient and higher levels of outpatient care. A certified recovery residence used by a licensed service provider that meets the definition of community housing shall be classified as a Level IV level of support, as described in subsection (5).

Section 3. Subsection (20) is added to section 397.321, Florida Statutes, to read:

- 397.321 Duties of the department.—The department shall:
- (20) Prominently display and make available on its website no later than January 1, 2025, all documents in the department's Provider Licensure and Designations System pertaining to the following:
- (a) Service provider applications for licensure and license renewal.
- (b) Policies and procedures provided to the department by an applicant for service provider licensure or license renewal.
- (c) The name and location of each recovery residence engaged in a referral relationship with a licensed service provider or service provider applicant, as required under ss. 397.4104 and 397.403(1)(j).
- (d) All complaints pertaining to service providers received by the department, and all investigative reports and findings, whether founded or unfounded. Complainant names and other identifying information shall be redacted.
- (e) Fines assessed for violations pursuant to ss. 397.411(7), 397.4104(2), and 397.4873(7).

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(f) All reports or other documentation pertaining to service provider license suspension or revocation.

- (g) All inspection reports for service provider licenses and recovery residences.
- Section 4. Paragraph (a) of subsection (2) of section 397.335, Florida Statutes, is amended to read:
 - 397.335 Statewide Council on Opioid Abatement.-
 - (2) MEMBERSHIP.-
- (a) Notwithstanding s. 20.052, the council shall be composed of the following members:
- 1. The Attorney General, or his or her designee, who shall serve as chair.
- 2. The secretary of the department, or his or her designee, who shall serve as vice chair.
 - 3. One member appointed by the Governor.
 - 4. One member appointed by the President of the Senate.
- 5. One member appointed by the Speaker of the House of Representatives.
- 6. Two members appointed by the Florida League of Cities who are commissioners or mayors of municipalities. One member shall be from a municipality with a population of fewer than 50,000 people.
- 7. Two members appointed by or through the Florida Association of Counties who are county commissioners or mayors. One member shall be appointed from a county with a population of fewer than 200,000, and one member shall be appointed from a county with a population of more than 200,000.
- 8. One member who is either a county commissioner or county mayor appointed by the Florida Association of Counties or who is

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a commissioner or mayor of a municipality appointed by the Florida League of Cities. The Florida Association of Counties shall appoint such member for the initial term, and future appointments must alternate between a member appointed by the Florida League of Cities and a member appointed by the Florida Association of Counties.

- 9. Two members appointed by or through the State Surgeon General. One shall be a staff member from the department who has experience coordinating state and local efforts to abate the opioid epidemic, and one shall be a licensed physician who is board certified in both addiction medicine and psychiatry.
- 10. One member appointed by the Florida Association of Recovery Residences.
- $\underline{\mbox{11. One member appointed by the Florida Association of EMS}}$ Medical Directors.
- 12. One member appointed by the Florida Society of Addiction Medicine who is a medical doctor board certified in addiction medicine.
- 13. One member appointed by the Florida Behavioral Health Association.
 - 14. One member appointed by Floridians for Recovery.

Section 5. Present paragraphs (c), (d), and (e) of subsection (8) of section 397.487, Florida Statutes, are redesignated as paragraphs (d), (e), and (f), respectively, a new paragraph (c) is added to that subsection, subsections (13) and (14) are added to that section, and paragraphs (b) and present paragraphs (c), (d), and (e) of subsection (8) of that section are amended, to read:

397.487 Voluntary certification of recovery residences.

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(8) Onsite followup monitoring of a certified recovery residence may be conducted by the credentialing entity to determine continuing compliance with certification requirements. The credentialing entity shall inspect each certified recovery residence at least annually to ensure compliance.

- (b) A certified recovery residence must notify the credentialing entity within 3 business days after the removal of the recovery residence's certified recovery residence administrator due to termination, resignation, or any other reason. The certified recovery residence has 90 30 days to retain a certified recovery residence administrator. The credentialing entity shall revoke the certificate of compliance of any certified recovery residence that fails to comply with this paragraph.
- (c) If a certified recovery residence's administrator has been removed due to termination, resignation, or any other reason and had been previously approved to actively manage more than 50 residents pursuant to s. 397.4871(8)(b), the certified recovery residence has 90 days to retain another certified recovery residence administrator pursuant to that section. The credentialing entity shall revoke the certificate of compliance of any certified recovery residence that fails to comply with this paragraph.
- (d) (e) If any owner, director, or chief financial officer of a certified recovery residence is arrested and awaiting disposition for or found guilty of, or enters a plea of guilty or nolo contendere to, regardless of whether adjudication is withheld, any offense listed in s. 435.04(2) while acting in that capacity, the certified recovery residence must shall

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immediately remove the person from that position and shall notify the credentialing entity within 3 business days after such removal. The credentialing entity may shall revoke the certificate of compliance of a certified recovery residence that fails to meet these requirements.

- (e) (d) A credentialing entity shall revoke a <u>certified</u> recovery residence's certificate of compliance if the <u>certified</u> recovery residence provides false or misleading information to the credentialing entity at any time.
- (f) (e) Any decision by a department-recognized credentialing entity to deny, revoke, or suspend a certification, or otherwise impose sanctions on a certified recovery residence, is reviewable by the department. Upon receiving an adverse determination, the certified recovery residence may request an administrative hearing pursuant to ss. 120.569 and 120.57(1) within 30 days after completing any appeals process offered by the credentialing entity or the department, as applicable.
- (13) On or after January 1, 2025, a recovery residence may not deny an individual access to housing solely on the basis that he or she has been prescribed federally approved medication that assists with treatment for substance use disorders by a licensed physician, a physician's assistant, or an advanced practice registered nurse registered under s. 464.0123.
- (14) A local law, ordinance, or regulation may not regulate the duration or frequency of a resident's stay in a certified recovery residence located within a multifamily zoning district. This subsection does not apply to any local law, ordinance, or regulation adopted on or before February 1, 2025.

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Section 6. Paragraphs (b) and (c) of subsection (6) of section 397.4871, Florida Statutes, are amended, and paragraph (c) is added to subsection (8) of that section, to read:

397.4871 Recovery residence administrator certification.-

- (6) The credentialing entity shall issue a certificate of compliance upon approval of a person's application. The certification shall automatically terminate 1 year after issuance if not renewed.
- (b) If a certified recovery residence administrator of a recovery residence is arrested and awaiting disposition for or found guilty of, or enters a plea of guilty or nolo contendere to, regardless of whether adjudication is withheld, any offense listed in s. 435.04(2) while acting in that capacity, the certified recovery residence must shall immediately remove the person from that position and shall notify the credentialing entity within 3 business days after such removal. The certified recovery residence shall have 30 days to retain a certified recovery residence administrator within 90 days after such removal. The credentialing entity shall revoke the certificate of compliance of any recovery residence that fails to meet these requirements.
- (c) A credentialing entity shall revoke a <u>certified</u> recovery residence administrator's certificate of compliance if the recovery residence administrator provides false or misleading information to the credentialing entity at any time.

(8)

(c) Notwithstanding paragraph (b), a Level IV certified recovery residence with a community housing component, which residence is actively managed by a certified recovery residence

31-00370C-24 20241180 291 administrator approved for 100 residents under this section and is wholly owned or controlled by a licensed service provider, 292 293 may actively manage up to 150 residents so long as the licensed 294 service provider maintains a service provider personnel-to-295 patient ratio of 1 to 8 and maintains onsite supervision at the 296 residences 24 hours a day, 7 days a week, with a personnel-to-297 resident ratio of 1 to 10. A certified recovery residence 298 administrator who has been removed by a certified recovery 299 residence due to termination, resignation, or any other reason 300 may not continue to actively manage more than 50 residents for 301 another service provider or certified recovery residence without 302 being approved by the credentialing entity. 303

Section 7. This act shall take effect July 1, 2024.

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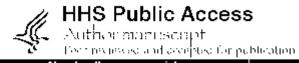
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Clinical Trial Design Challenges and Opportunities for Emerging Treatments for Opioid Use Disorder A Review

Brian D. Kiluk, PhD, Bethea A. Kleykamp, PhD, Sandra D. Comer, PhD, Roland R. Griffiths, PhD, Andrew S. Huhn, PhD, MBA, Matthew W. Johnson, PhD, Kyle M. Kampman, MD, Marco Pravetoni, PhD, Kenzie L. Preston, PhD, Ryan Vandrey, PhD, Cecilia L. Bergeria, PhD, Michael P. Bogenschutz, MD, Randall T. Brown, MD, PhD, Kelly E. Dunn, PhD, MBA, Robert H. Dworkin, PhD, Patrick H. Finan, PhD, Peter S. Hendricks, PhD, Elisabeth J. Houtsmuller, PhD, Thomas R. Kosten, MD, Dustin C. Lee, PhD, Frances R. Levin, MD, Aimee McRae-Clark, PharmD, Charles L. Raison, MD, Kurt Rasmussen, PhD, Dennis C. Turk, PhD, Roger D. Weiss, MD, and Eric C. Strain, MD

Abstract

IMPORTANCE

Novel treatments for opioid use disorder (OUD) are needed to address both the ongoing opioid epidemic and long-standing barriers to existing OUD treatments that target the endogenous μ -opioid receptor (MOR) system. The goal of this review is to highlight unique clinical trial design considerations for the study of emerging treatments for OUD that address targets beyond the MOR system. In November 2019, the Analgesic, Anesthetic, and Addiction Clinical Trial Translations, Innovations, Opportunities, and Networks

(ACTTION) public-private partnership with the US Food and Drug Administration sponsored a meeting to discuss the current evidence regarding potential treatments for OUD, including cannabinoids, psychedelics, sedative-hypnotics, and immunotherapeutics, such as vaccines.

OBSERVATIONS

Consensus recommendations are presented regarding the most critical elements of trial design for the evaluation of novel OUD treatments, such as: (1) stage of treatment that will be targeted (eg, seeking treatment, early abstinence/detoxification, long-term recovery); (2) role of treatment (adjunctive with or independent of existing OUD treatments); (3) primary outcomes informed by patient preferences that assess opioid use (including changes in patterns of use), treatment retention, and/or global functioning and quality of life; and (4) adverse events, including the potential for opioid-related relapse or overdose, especially if the patient is not simultaneously taking maintenance MOR agonist or antagonist medications.

CONCLUSIONS AND RELEVANCE

Applying the recommendations provided here as well as considering input from people with lived experience in the design phase will accelerate the development, translation, and uptake of effective and safe therapeutics for individuals struggling with OUD.

Opioid use disorder (OUD) is a major cause of disease burden, leading to increased pregnancy or birth complications, viral infections, and fatal overdoses. $^{1-3}$ The 3 effective and safe medications for treating OUD (MOUD) act through the μ -opioid receptor (MOR), the primary target for opioids misused for their rewarding effects. The MOR agonists methadone or buprenorphine and the MOR antagonist naltrexone are the standard of care for OUD because they reduce risk of relapse, overdose deaths, infections, and criminal behavior, but discontinuation and relapse still exceed 50% within 6 months. Furthermore, each of these MOUDs have different induction and dosing procedures as well as regulatory, policy, and patient-level barriers that have hindered patient access and retention. Thus, OUD treatment options need expansion through development of novel stand-alone therapies or adjuncts to existing MOR-based MOUDs. $^{10-12}$

A critical step in developing novel treatments for OUD is the completion of randomized clinical trials (RCTs). However, the inherent features of OUD, including a pronounced physical dependence and a high risk of overdose, suggest the design of these trials will likely need to differ from designs used to evaluate existing treatments for OUD. There is not a strong consensus in the OUD field concerning standardized key trial design decisions or outcome measures. Given the importance of this topic and the need for new and novel OUD treatments, a meeting sponsored by the Analgesic, Anesthetic, and Addiction ClinicalTrialTranslations, Innovations, Opportunities, and

Networks (ACTTION) public-private partnership with the US Food and Drug Administration (FDA) was convened in November 2019 to discuss study design considerations unique to 4 candidate medication categories for OUD that do not directly target the MOR system: cannabinoids, psychedelics, sedative-hypnotics, and immunotherapeutics; a summary of highlights from the meeting has been previously published. This article reviews the key trial considerations derived from that meeting and provides consensus considerations and recommendations for studies of non-MOR-based treatments for OUD.

Methods

The ACTTION Consortium for Addiction Research on Efficacy and Safety (CARES) meeting included participants from academia, government, and nonprofit organizations selected on the basis of their research, clinical, or administrative expertise relevant to the candidate medication categories or clinical trials of OUD treatments. There was no direct participation from any pharmaceutical company. Meeting details, including agenda, goals, list of attendees, presentations, and transcripts of discussion, are available on the CARES website. The following considerations and recommendations were informed by the meeting presentations and discussions, literature reviews, and coauthors' feedback on iterative revisions of drafts of this article.

Discussion

Study Planning

Study planning should begin by specifying the stage in OUD treatment targeted by the intervention(s), as this decision will influence all subsequent design decisions. The core stages in the OUD treatment and recovery trajectory can be conceptualized as (1) current active use of opioids; (2) acute abstinence, nonmedically supervised withdrawal, and/or supervised medical withdrawal; (3) early recovery (eg, less than 6 months of abstinence with or without opioid agonist or antagonist treatment); and (4) sustained recovery (eg, abstinence from illicit opioid use for at least 6 months). Each stage has unique treatment needs, and study planning should consider whether the novel treatment will be adjunctive to existing regulatory agency-approved MOUD, which may be essential for those with physical dependence and withdrawal symptoms. The need for adjunctive treatment, including harm reduction strategies, such as naloxone training to prevent fatal overdose, would be essential for clinically unstable patients.

Historically, few OUD trials have incorporated the preferences of patients, and per patient-focused drug development, $\frac{15}{15}$ we recommend using input from people with lived experience to guide the choice of primary and secondary outcomes. For instance, although treatment retention was found as the most reported outcome across 60 OUD trials, many patients report an eagerness to complete

therapy and end agonist treatment as a main goal. Strategies for incorporating patient perspectives into study planning include focus groups, interviews, online surveys, workshops, social media listening, and community-based participatory research strategies. Tr.18 Guidance on methods for engaging patients and other relevant stakeholders are described elsewhere.

Study Design

Intervention (Including Randomization, Blinding, and Dosing) Trial designs will be dictated to a large extent by the stage of treatment that the intervention is targeting as well as the unique properties of the intervention under evaluation. The <u>Table</u> gives an overview of specific considerations for the 4 types of emerging medication treatments reviewed here. The National Institute on Drug Abuse has identified additional emerging areas of interest for OUD treatment development that target a range of novel pharmacological mechanisms of action, such as respiratory stimulants, γ -aminobutyric acid metabotropic receptor family B agonists, and ghrelin antagonists. Discussing all emerging treatments, including nonmedication interventions (eg, repetitive transcranial magneticstimulation $\frac{55}{2}$), was beyond the scope of this meeting, yet many of the considerations and recommendations described here also apply to these other approaches. Each of these emerging treatments has specific characteristics that influence study design choices, including dosing, mode of administration, and timing of intervention relative to treatment stage.

Table.

Unique Considerations Associated With 4 Emerging Treatments for Opioid Use Disorder

				Immunotherapeutics
	Cannabis and			(vaccines and monoclonal
Consideration	cannabinoids	Psychedelics	Sedative-hypnotics	antibodies)
Rationale	The endocannabinoid and	Classic psychedelics (serotonin 2A	Sleep is a basic biological	Active (vaccination) and
	opioid systems interact	receptor agonists) have been	system that can be affected	passive (transfer of premade
	with some subtypes of	associated with reduced substance	by opioid use and can also	antibodies) immunization
	cannabinoid receptors that	use in naturalistic and clinical	affect the trajectory of	strategies rely on the presence
	influence the rewarding	settings, with the strongest	opioid use. Sleep	of drug-specific antibodies to
	effects of opioids. Some	evidence for LSD as a treatment of	dysfunction is a common	selectively bind to target
	uncontrolled observational	alcoholism. Candidate	issue across all substance	opioids in plasma and prevent
	research has suggested that	psychological mechanisms of	use disorders. Sleep	drugs from crossing the blood-
	cannabis and cannabinoids	action include awe, cognitive	disturbance can have	brain barrier and reaching the
	can have a substitution	flexibility, and insight; candidate	profound effects on a	brain. By reducing the
	effect on opioid use	biological mechanisms include	patient's life, including	concentration of free
	behavior $\frac{20-27}{}$	inflammation and brain network	ability to cope with craving,	(unbound) opioids in the brain,
		functioning. ²⁸ - ³⁴	and can affect the cognitive	vaccines and monoclonal
			effort associated with	antibodies reduce opioids'
			opioid abstinence. 35-41	pharmacological effects. 42-50
Types	Cannabis is a complex	LSD, psilocybin (found in <i>Psilocybe</i>	Orexin-1 or 1/2 antagonists;	Individual and multivalent
	chemical entity that	mushrooms), mescaline (found in	tricyclics (Doxepin);	vaccines targeting specific
	contains >100 botanically	peyote and other cacti), and	antipsychotics	types of opioids, including
	derived phytocannabinoids,	dimethyltryptamine (found in	(Quetiapine); melatonin;	oxycodone, heroin, and
	each of which can be	ayahuasca).	mirtazapine; or ramelteon.	fentanyl. Individual and
	synthesized or isolated.		Benzodiazepines or	multivalent monoclonal
			benzodiazepinelike drugs	antibodies formulations against
			(eg, Zolpidem) are often not	various opioids.
			used due to risk for misuse.	•

Abbreviations: LSD, lysergic acid diethylamide; OUD, opioid use disorder; PSQI, Pittsburgh Sleep Quality Index.

For drug development, the criterion-standard efficacy and safety studies are double-blind, placebo-controlled RCT designs. However, for OUD, these designs face ethical concerns of a placeboonly condition and challenges in blinding treatment groups. Additional research designs that could be considered include adaptive or pragmatic trials and the use of real-world data as primary or secondary outcomes. Feedomers of the specifics of blinding and ran-domization, we recommend that efforts to examine novel compounds be paired with some form of standardized and efficacious psychosocial support, including in-person or digital treatment modules, to mitigate the risk that patients are left with no treatment if a compound fails.

Comparators

The severe nature of the opioid physical dependence syndrome means that a placebo-controlled trial in the absence of an agonist MOUD might be unsafe or unfeasible for patients who are in early abstinence and at risk of opioid withdrawal symptoms, relapse, or overdose. Relevant alternative types of comparators include (1) low or subtherapeutic doses of study medication, (2) ascending doses of study medication, (3) standard-of-care pharmacologic or nonpharmacologic treatments in a comparative effectiveness trial design, or (4) a combination of different comparators.

The type of comparator will also influence whether the objective of the clinical trial is to test superiority or noninferiority between different treatment conditions. Investigators may choose to provide an MOUD as a plat form therapy for all participants while comparing an active vs placebo adjunctive medication using a superiority trial design (eg, a sleep agent compared with placebo for those stabilized with methadone). Ethical concerns related to place bodosing could also increase the appeal of noninferiority trials, although these are more complex in design and analysis than superiority trials, with challenges described elsewhere. ⁵⁹

Study Setting

RCTs of MOR-based MOUDs have been traditionally completed on an outpatient basis in settings, such as opioid treatment programs or medical offices, because of inherent restrictions on MOUD prescribing and dispensing. Some emerging treatments, such as sedative-hypnotics or vaccines, may have fewer regulatory or medical requirements compared with MOR-based treatments and therefore may afford more flexibility in the study designs and open opportunities for novel approaches. Methods for remote data collection have advanced considerably during the SARS-CoV-2 pandemic, expanding possible approaches to collecting substance use outcomes (eg, remotely collected breathalyzer data for alcohol or tobacco use). Recent parallel efforts to leverage nonspecialized care professionals to expand the OUD treatment infrastructure, including health care professionals, 45,66 may further bolster innova-

tion. However, these approaches may not be useful in all cases; the study of some agents may require even more intensive in-person designs compared with traditional OUD clinical trials. The in-person interactions and monitoring required for safe delivery and evaluation of some novel treatments present challenges to conducting clinical trials on a larger scale, an issue the field has acknowledged and begun to address with more scalable intervention paradigms. 67,68

Participant Characteristics

Participant selection in the form of inclusion and exclusion criteria are essential for ensuring that a trial targets the population of interest, minimizes variance in outcomes because of factors other than the intervention, and supports future meta-analyses. At minimum, we recommend that the following categories be addressed in the study inclusion and exclusion criteria and/or baseline data collection associated with the study: (1) opioid use variables, including historical (lifetime) and current (past year) opioid use behavior, including type, timing, amount, and route of administration of opioid(s), previous experience with opioid overdose, including hospitalization, OUD treatment history, and degree of OUD severity; (2) historical or current alcohol and other substance use disorders, including prior use of target medication; (3) medical history, including prescribed medications in past 90 days and concomitant medical and psychiatric conditions; and (4) psychosocial variables (eg, problems resulting from opioid use, including incarceration). In addition, basic patient demographic characteristics (eg, age, sex, gender, race and ethnicity, and socioeconomic status) should be collected with awareness of specific populations that are at risk of developing OUD or those who experience disparate consequences, including individuals with mental health disorders, ⁶⁹/₅ youth and young adults, ⁷⁰/₇ military veterans, ⁷¹/₇ pregnant women, ⁷²/₇ racial and ethnic minority populations, ^{73,74}/₇ and individuals from particular geographic regions (eg, US Appalachian and Southern states). ^{75,76}/_{75,76} Limitations should be considered when selecting eligibility criteria depending on specific safety considerations associated with the intervention under study.

Outcome Measures

The type of efficacy outcomes chosen for a trial depends on the goal of the trial (eg, targeted phase of OUD treatment, key comparators). Literature reviews have noted that primary and secondary outcomes and their associated measures vary widely across clinical trials for OUD. 16,77 Opioid abstinence and treatment retention have been the most common primary end points in clinical trials for OUD and other substance use disorders. However, there is an evolving understanding of the importance of continuous measures of opioiduse, including changes in use patterns, such as the frequency, duration, and amount of use.

The degree to which these different, but important, outcomes are clinically meaningful is still being debated. L16,79,80 Currently there are no criterion-standard outcomes in OUD trials. Thus, the below recommendations are meant to functionas guide posts when choosing outcomes.

Primary Outcomes

The dichotomous outcome of opioid abstinence, defined as no detected or self-reported use within an assessment window, has been the most common measure of opioid use behavior in clinical trials. $\frac{78}{2}$ According to the FDA Guidance for Industry regarding end points for demonstrating effectiveness of drugs for treatment of OUD, $\frac{78}{2}$ drug use patterns other than abstinence can be used as thresholds to define treatment response. Measurement of such response-defining thresholds must be specified, and evidence from clinical trials, longitudinal observation studies, or other sources are needed to support the clinical benefit of a given drug use pattern (ie, reduction). $\frac{78}{2}$ We recommend that both abstinence and patterns of opioid use be measured and that clear responder criteria be specified for each, with the potential for a grace period. For trials that identify opioid abstinence as the primary outcome, we recommend opioid use be assessed using objective (eg, urinalysis) and subjective (eg, patient, clinician, and/or observer) measures. $\frac{77}{2}$ The field is currently moving to less frequent objective testing of these outcomes for practical reasons and to reduce the burden on participants. We recommend that decisions regarding frequency of testing be based on the clinical stability of the patient population, the pharmacological properties of treatment, and participation burden.

Trials of MOR-based treatments demonstrate that retention in treatment longer than 6 months is associated with better treatment outcomes compared with shorter durations of treatment or no treatment. However, neither we northe FDA 78 recommend that treatment retention be a stand-alone clinical end point, as retention can be easily influenced or driven by factors external to the intervention being examined. We recommend that at least 1 outcome consider general patient functioning as assessed through prepost changes in *DSM* OUD diagnostic status or symptom criteria, quality of life assessment tools, or other patient-centered outcomes that can better capture how a treatment is affecting a patient's life beyond acute opioid exposure.

Secondary Outcomes

Key secondary outcomes, which could be primary outcomes depending on the aims of the study, include: (1) opioid withdrawal signs and symptoms; (2) opioidcraving; (3) treatment adherence; (4) treatment satisfaction; (5) physical health (eg, comorbid diagnoses, including chronicpain); (6) mental health (eg, anxiety, depression, and other substance use); (7) cognitive and physical functioning (eg, memory, attention, sleep duration and quality, and pain severity); (8) personal and social functioning (eg, family and social relations, criminal behavior, employment, schooling, relationships, and housing and food stability); (9) health risk behavior (eg, hospitalizations, overdoses), and (10) risk of medication misuse (eg, rewarding or reinforcing effects of medication).

Risk and Adverse Events

A critical outcome in OUD trials includes opioid-related overdose or death, which is at increased risk during treatment initiation and the first several weeks after initiating abstinence or attempting opioid withdrawal. We recommend that trials, especially early treatment trials, include frequent assessment of these opioid-related adverse events, which include hospitalization, naloxone administration, and emergency department visits. Trials should also include counseling on opioid overdose risk knowledge at the onset of enrollment (eg, Brief Opioid Overdose Knowledge tutorial or the Overdose Education and Naloxone Distribution training) and provide naloxone.

Additional opioid-specific risks that might be monitored include infectious disease exposure and seroconversion rates (eg, HIV and hepatitis C). Emerging treatments may have unique adverse effects and events that should be monitored. For example, immunotherapeutics, such as vaccines and monoclonal antibodies specific for opioids, should be carefully evaluated for immune-related adverse effects in immunocompromised patients. In contrast, some sedative-hypnotic medications and cannabinoids have risks, including acute psychiatric and/or physical health consequences, misuse risk, drug-drug interactions, and diversion that should be monitored. Examples of potential risks of emerging treatments covered in the present review are included in the Table.

Challenges and Opportunities

Regulatory requirements and quality control issues, including variations in regulation at the regional and national levels in the US and other countries, can make large-scale clinical trials challenging. For example, cannabis (and other cannabinoids) and psilocybin (and other psychedelics) are all classified as schedule I drugs according to the Federal US Controlled Substances Act (ie, drugs with no currently accepted medical use and a high potential for misuse), making it more challenging and administratively burdensome to conduct clinical trials. Relatedly, both classes of drugs have a controversial history, including issues with social acceptance and legality. Meanwhile, state-level regulation of cannabinoids has led to variable (if any) manufacturing standards across states, resulting in intervariations and intravariations in potency and dosing across cannabinoid products. This makes it difficult to generalize research findings across some marketed consumer products.

These challenges and perspectives are slowly changing, as evidenced by the recent FDA breakthrough therapy designation for psilocybin in the treatment of depression, and 3,4-methylenedioxy-methamphetamine (MDMA) in the treatment of posttraumatic stress disorder. In contrast, opioid vaccines are not designated as controlled substances by the US Drug Enforcement Administration (DEA), and therefore, DEA regulations would not complicate treatment per se. However, opioid conjugate vaccines consist of multiple components, including an opioid-based small molecule hapten, which could be regulated by the DEA as either a schedule I or II drug, thereby affecting research and manufacturing. Manufacturing challenges related to DEA drug scheduling apply to a broad range of compounds currently in development, including synthetic cannabinoids, psychedelics, and nontraditional opioid receptor agonists and antagonists.

Another challenge is that the types of opioids being used has expanded from commercially produced opioids and heroin to also include fentanyl and/or its structural analogs, resulting in a dynamic opioid marketplace for which research may lag street-level use, type of drug, and availability. Recent data suggest increased exposure to fentanyl and its structural analogs across the US. 92,93 Opioids produce diverse effects on the development and nature of opioid physical dependence and withdrawal, and fentanyl appears to be engendering a unique and particularly severe withdrawal syndrome. Establishing a treatment's efficacy becomes especially challenging when the type of substance being targeted has such wide variability in terms of potency, route of administration, detectability, and potential for adverse outcomes.

A third challenge is that the complexity of OUD and its different stages of development are likely to have different (albeit over-lapping) underlying mechanisms that require different types of or combinations of treatments. For example, early sporadic use is a different stage in the life cycle from years of chronic, daily use. Furthermore, medication alone is often not a sufficient treatment for OUD, and it is important to include psychosocial and behavioral interventions and to tailor these nonpharmacological interventions to the stage of opioid use. There remain gaps in our understanding of how best to combine pharmacological and behavioral treatments. $\frac{58}{100}$

Despite these and other challenges, there are valuable opportunities for clinical trials with emerging treatments. Research methods are developing quickly, especially in sleep measurement, wearable devices for drug detection, remote data collection (eg, telehealth and wearable technology), and the development of genetic bio-markers for selection of phenotypes and endophenotypes that may better reflect underlying neurobiological mechanisms. The present review focused on study design considerations for clinical trials and did not discuss other relevant types of research, including pre-clinical studies, laboratory-based within-subject human studies, and observational/epidemiological studies.

Conclusions

The Box provides a summary of the key considerations and recommendations for clinical trials evaluating emerging non-MOR treatments for OUD. Promoting a unifying structure of best research practices as described in the present review will help the field build consensus as to the appropriate methodological strategies and prevent otherwise promising targets from languishing or being abandoned because of problematic study designs rather than true lack of efficacy or lack of uptake. In the context of a continually evolving and escalating opioid crisis, research must prioritize both innovation and efficiency. The field and the patients with whom we work will be best served by maintaining an open dialogue to develop a consistent methodological framework for the assessment and treatment of OUD.

Box.

Key Recommendations and Considerations

Study Objectives

- Prospective trial registration prior to the start data collection in publicly accessible database, including primary and secondary outcomes, hypotheses, and study objectives.
- Priority should be given to specifying the stage in OUD treatment that will be targeted with the intervention (eg, current active use of opioids, acute abstinence, nonmedically supervised withdrawal, and/or supervised withdrawal, early recovery, or long-term recovery) and determining whether the emerging treatment will be adjunctive to or independent of existing OUD treatments.

Clinical Trial Design

- Study design will ideally be double-blind randomized clinical trial.
- Comparators should include a placebo group (when ethically appropriate) and/or an active control comparison(s).
- If the novel treatment is a stand-alone intervention, then comparison should include an existing, evidence-based OUD treatment (eg, methadone, buprenorphine, naltrexone, or behavioral/psychosocial support).

Sample

- Participants should be a representative, diverse population of patients (ie, age, sex, sexual orientation, race and ethnicity, socioeconomic status, and history of substance use).
- Exclusion criteria that are too restrictive and may negatively affect the generalizability of the study should be carefully evaluated and included on the basis of safety or another enhanced rationale considered (eg, exclusion of participants with concurrent medical, physical, or mental health issues).

Primary End Point

- Primary outcomes should be chosen to align with the study objectives and the phase of treatment that is to be targeted (eg, symptoms of opioid withdrawal or craving will be more important to measure in early recovery rather than during long-term recovery). In addition, primary outcomes will need to be tailored to the expected treatment indication (eg, sleep measures for a sleep intervention).
- At minimum, we recommend that primary outcomes for trials beyond phase I include opioid use behavior, treatment retention, and at least 1 outcome that addresses global functioning (eg, change in *DSM* criteria, quality of life).
- A dichotomous measure to define responder (based on opioid abstinence or reduction in opioid use) should be a primary outcome, but also consider continuous measures of opioid use (ie, quantity, frequency).
- Selection of end points should be informed by input from patients and family members to determine the most salient OUD symptoms/experiences and outcomes.

Secondary Outcomes

• Potential secondary outcomes should include opioid withdrawal and/or craving, treatment adherence and satisfaction, physical and mental health, risk of misuse of study intervention, patient-focused outcomes, such as psychosocial functioning (including employment and legal issues), sleep, pain, and cognitive functioning, and health outcomes (eg, viral load if positive for HIV or hepatitis C virus).

Assessment of Harms

• Adverse events, including opioid-related adverse events (eg, hospitalization, naloxone administration, visits to emergency department), and reasons for premature terminations from trial should be collected and carefully reviewed with sensitivity to relapse risk and overdose.

Abbreviation: OUD, opioid use disorder.

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Conflict of Interest Disclosures:

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Footnotes

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REFERENCES

- 1. Volkow ND, Woodcock J, Compton WM, et al. Medication development in opioid addiction: meaningful clinical end points. *Sci Transl Med*. 2018; 10(434):eaan2595. doi: 10.1126/scitranslmed.aan2595 [PubMed] [CrossRef] [Google Scholar]
- 2. Degenhardt L, Charlson F, Mathers B, et al. The global epidemiology and burden of opioid dependence: results from the Global Burden of Disease 2010 study. *Addiction*. 2014;109(8):1320–1333. doi: 10.1111/add.12551 [PubMed] [CrossRef] [Google Scholar]
- 3. Hser Y-I, Mooney LJ, Saxon AJ, et al. High mortality among patients with opioid use disorder in a large healthcare system. *J Addict Med.* 2017;11 (4):315–319. doi: 10.1097/ADM.00000000000312 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 4. Volkow ND, Blanco C. Medications for opioid use disorders: clinical and pharmacological considerations. *J Clin Invest*. 2020;130(1):10–13. doi: 10.1172/JCI134708

 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 5. National Academies of Sciences, Engineering, and Medicine; Health and Medicine Division; Board on Health Sciences Policy; Committee on Medication-Assisted Treatment for Opioid Use Disorder; Mancher M, Leshner AI, eds. *Medications for Opioid Use Disorder Save Lives*. National Academies Press; 2019. [PubMed] [Google Scholar]
- 6. Morgan JR, Schackman BR, Leff JA, Linas BP, Walley AY. Injectable naltrexone, oral naltrexone, and buprenorphine utilization and discontinuation among individuals treated for opioid use disorder in a United States commercially insured population. *J Subst Abuse Treat*. 2018;85:90–96. doi: 10.1016/j.jsat.2017.07.001 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 7. Lee JD, Nunes EV Jr, Novo P, et al. Comparative effectiveness of extended-release naltrexone versus buprenorphine-naloxone for opioid relapse prevention (X:BOT): a multicentre, open-label, randomised controlled trial. *Lancet*. 2018;391 (10118):309–318. doi: 10.1016/S0140-6736(17)32812-X [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 8. O'Connor AM, Cousins G, Durand L, Barry J, Boland F. Retention of patients in opioid substitution treatment: a systematic review. *PLoS One*. 2020;15(5):e0232086. doi: 10.1371/journal.pone.0232086 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 9. Volkow ND, Frieden TR, Hyde PS, Cha SS. Medication-assisted therapies–tackling the opioid-overdose epidemic. *N Engl J Med*. 2014;370 (22):2063–2066. doi: 10.1056/NEJMp1402780 [PubMed] [CrossRef] [Google Scholar]
- 10. Fuehrlein BS, Ross DA. Opioid use disorder: a desperate need for novel treatments. *Biol Psychiatry*. 2017;81(7):e43–e45. doi: 10.1016/j.biopsych.2017.01.014 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 11. Rasmussen K, White DA, Acri JB. NIDA's medication development priorities in response to the opioid crisis: ten most wanted. *Neuropsychopharmacology*. 2019;44(4):657–659. doi: 10.1038/s41386-018-0292-5 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 12. Blanco C, Volkow ND. Management of opioid use disorder in the USA: present status and future directions. *Lancet*. 2019;393(10182):1760–1772. doi: 10.1016/S0140-6736(18)33078-2 [PubMed] [CrossRef] [Google Scholar]
- 13. Strain EC, Kampman KM, Weiss RD. Moving beyond medications that act at the μ receptor in the treatment of opioid use disorder. *JAMA Psychiatry*. 2021;78(7):701–702. doi: 10.1001/jamapsychiatry.2021.0259 [PubMed] [CrossRef] [Google Scholar]
- 14. Analgesic, Anesthetic, and Addiction Clinical Trial Translations, Innovations, Opportunities, and Networks. *Consortium for Addiction Research on Efficacy and Safety (CARES)*. Accessed June 10, 2022. https://www.acttion.org/cares [Google Scholar]
- 15. Food US and Administration Drug. *Patient-focused drug development: collecting comprehensive and representative input*. Accessed July 27, 2021. https://www.fda.gov/regulatory-information/search-fda-guidance-documents/patient-focused-drug-development-collecting-comprehensive-and-representative-input
- 16. Dennis BB, Sanger N, Bawor M, et al. A call for consensus in defining efficacy in clinical trials for opioid addiction: combined results from a systematic review and qualitative study in patients receiving pharmacological assisted therapy for opioid use disorder. *Trials*. 2020;21(1):30. doi: 10.1186/s13063-019-3995-y [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 17. Cook NS, Cave J, Holtorf A-P. Patient preference studies during early drug development: aligning stakeholders to ensure development plans meet patient needs. *Front Med (Lausanne)*. 2019;6:82. doi: 10.3389/fmed.2019.00082 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 18. Wallerstein N, Duran B. Community-based participatory research contributions to intervention research: the intersection of science and practice to improve health equity. *Am J Public Health*. 2010; 100(suppl 1):S40–S46. doi: 10.2105/AJPH.2009.184036 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 19. Sheridan S, Schrandt S, Forsythe L, Hilliard TS, Paez KA; Advisory Panel on Patient Engagement (2013 Inaugural Panel). The PCORI Engagement Rubric: promising practices for partnering in research. *Ann Fam Med.* 2017;15(2):165–170. doi: 10.1370/afm.2042 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 20. Humphreys K, Saitz R. Should physicians recommend replacing opioids with cannabis? *JAMA*. 2019;321(7):639–640. doi: 10.1001/jama.2019.0077 [PubMed] [CrossRef] [Google Scholar]
- 21. Hurd YL. Cannabidiol: swinging the marijuana pendulum from 'weed' to medication to treat the opioid epidemic. *Trends Neurosci*. 2017;40(3):124–127. doi: 10.1016/j.tins.2016.12.006 [PubMed] [CrossRef] [Google Scholar]
- 22. Hurd YL, Yoon M, Manini AF, et al. Early phase in the development of cannabidiol as a treatment for addiction: opioid relapse takes initial center stage. *Neurotherapeutics*. 2015;12(4):807–815. doi: 10.1007/s13311-015-0373-7 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 23. Lucas P Rationale for cannabis-based interventions in the opioid overdose crisis. *Harm Reduct J.* 2017;14(1):58. doi: 10.1186/s12954-017-0183-9 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 24. Bisaga A, Sullivan MA, Glass A, et al. The effects of dronabinol during detoxification and the initiation of treatment with extended release naltrexone. *Drug Alcohol Depend*. 2015;154:38–45. doi: 10.1016/j.drugalcdep.2015.05.013 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 25. Jicha CJ, Lofwall MR, Nuzzo PA, Babalonis S, Elayi SC, Walsh SL. Safety of oral dronabinol during opioid withdrawal in humans. *Drug Alcohol Depend*. 2015;157:179–183. doi: 10.1016/j.drugalcdep.2015.09.031 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 26. Lofwall MR, Babalonis S, Nuzzo PA, Elayi SC, Walsh SL. Opioid withdrawal suppression efficacy of oral dronabinol in opioid dependent humans. *Drug Alcohol Depend*. 2016;164:143–150. doi: 10.1016/j.drugalcdep.2016.05.002 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 27. Wiese B, Wilson-Poe AR. Emerging evidence for cannabis' role in opioid use disorder. *Cannabis Cannabinoid Res.* 2018;3(1):179–189. doi: 10.1089/can.2018.0022 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 28. Nichols DE, Johnson MW, Nichols CD. Psychedelics as medicines: an emerging new paradigm. *Clin Pharmacol Ther*. 2017;101(2):209–219. doi: 10.1002/cpt.557 [PubMed] [CrossRef] [Google Scholar]
- 29. Garcia-Romeu A, Kersgaard B, Addy PH. Clinical applications of hallucinogens: a review. *Exp Clin Psychopharmacol*. 2016;24(4):229–268. doi: 10.1037/pha0000084
 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 30. Krebs TS, Johansen PØ. Lysergic acid diethylamide (LSD) for alcoholism: meta-analysis of randomized controlled trials. *J Psychopharmacol*. 2012;26(7):994–1002. doi: 10.1177/0269881112439253 [PubMed] [CrossRef] [Google Scholar]
- 31. Garcia-Romeu A, Davis AK, Erowid E, Erowid F, Griffiths RR, Johnson MW. Persisting reductions in cannabis, opioid, and stimulant misuse after naturalistic psychedelic use: an online survey. Front Psychiatry. 2020;10:955–955. doi: 10.3389/fpsyt.2019.00955 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 32. Pisano VD, Putnam NP, Kramer HM, Franciotti KJ, Halpern JH, Holden SC. The association of psychedelic use and opioid use disorders among illicit users in the United States. *J Psychopharmacol*. 2017;31(5):606–613. doi: 10.1177/0269881117691453 [PubMed] [CrossRef] [Google Scholar]
- 33. Belouin SJ, Henningfield JE. Psychedelics: where we are now, why we got here, what we must do. *Neuropharmacology*. 2018;142:7–19. doi: 10.1016/j.neuropharm.2018.02.018 [PubMed] [CrossRef] [Google Scholar]
- 34. Argento E, Tupper KW, Socias ME. The tripping point: the potential role of psychedelic-assisted therapy in the response to the opioid crisis. *Int J Drug Policy*. 2019;66:80–81. doi: 10.1016/j.drugpo.2018.11.006 [PubMed] [CrossRef] [Google Scholar]
- 35. Cheatle MD, Webster LR. Opioid therapy and sleep disorders: risks and mitigation strategies. *Pain Med.* 2015;16(suppl 1):S22–S26. doi: 10.1111/pme.12910 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 36. Huhn AS, Harris J, Cleveland HH, et al. Ecological momentary assessment of affect and craving in patients in treatment for prescription opioid dependence. *Brain Res Bull*. 2016;123:94–101. doi: 10.1016/j.brainresbull.2016.01.012 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 37. Lydon-Staley DM, Cleveland HH, Huhn AS, et al. Daily sleep quality affects drug craving, partially through indirect associations with positive affect, in patients in treatment for nonmedical use of prescription drugs. *Addict Behav.* 2017;65:275–282. doi: 10.1016/j.addbeh.2016.08.026 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 38. Fathi HR, Yoonessi A, Khatibi A, Rezaeitalab F, Rezaei-Ardani A. Crosstalk between sleep disturbance and opioid use disorder: a narrative review. *Addict Health*. 2020;12(2):140–158. doi: 10.22122/ahj.v12i2.249 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 39. Freeman LK, Gottfredson NC. Using ecological momentary assessment to assess the temporal relationship between sleep quality and cravings in individuals recovering from substance use disorders. Addict Behav. 2018;83:95–101. doi: 10.1016/j.addbeh.2017.11.001 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 40. Dolsen MR, Harvey AG. Life-time history of insomnia and hypersomnia symptoms as correlates of alcohol, cocaine and heroin use and relapse among adults seeking substance use treatment in the United States from 1991 to 1994. *Addiction*. 2017;112(6):1104–1111. doi: 10.1111/add.13772 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 41. James MH, Fragale JE, Aurora RN, Cooperman NA, Langleben DD, Aston-Jones G. Repurposing the dual orexin receptor antagonist suvorexant for the treatment of opioid use disorder: why sleep on this any longer? *Neuropsychopharmacology*. 2020;45 (5):717–719. doi: 10.1038/s41386-020-0619-x [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 42. Pravetoni M, Comer SD. Development of vaccines to treat opioid use disorders and reduce incidence of overdose. *Neuropharmacology*. 2019; 158:107662. doi: 10.1016/j.neuropharm.2019.06.001 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 43. Baehr C, Pravetoni M. Vaccines to treat opioid use disorders and to reduce opioid overdoses. *Neuropsychopharmacology*. 2019;44(1):217–218. doi: 10.1038/s41386-018-0197-3 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 44. Bremer PT, Kimishima A, Schlosburg JE, Zhou B, Collins KC, Janda KD. Combatting synthetic designer opioids: a conjugate vaccine ablates lethal doses of fentanyl class drugs. *Angew Chem Int Ed Engl.* 2016;55(11):3772–3775. doi: 10.1002/anie.201511654 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 45. Ohia-Nwoko O, Kosten TA, Haile CN. Animal models and the development of vaccines to treat substance use disorders. *Int Rev Neurobiol*. 2016; 126:263–291. doi: 10.1016/bs.irn.2016.02.009 [PubMed] [CrossRef] [Google Scholar]
- 46. Tenney RD, Blake S, Bremer PT, et al. Vaccine blunts fentanyl potency in male rhesus monkeys. *Neuropharmacology*. 2019;158:107730. doi: 10.1016/j.neuropharm.2019.107730 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 47. Townsend EA, Blake S, Faunce KE, et al. Conjugate vaccine produces long-lasting attenuation of fentanyl vs. food choice and blocks expression of opioid withdrawal-induced increases in fentanyl choice in rats. *Neuropsychopharmacology*. 2019;44(10):1681–1689. doi: 10.1038/s41386-019-0385-9 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 48. Banks ML, Olson ME, Janda KD. Immunopharmacotherapies for treating opioid use disorder. *Trends Pharmacol Sci.* 2018;39(11):908–911. doi: 10.1016/j.tips.2018.08.001 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 49. Raleigh MD, King SJ, Baruffaldi F, et al. Pharmacological mechanisms underlying the efficacy of antibodies generated by a vaccine to treat oxycodone use disorder. Neuropharmacology. 2021;195:108653. doi: 10.1016/j.neuropharm.2021.108653 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 50. Baehr C, Kelcher AH, Khaimraj A, et al. Monoclonal antibodies counteract opioid-induced behavioral and toxic effects in mice and rats. *J Pharmacol Exp Ther*. 2020;375(3):469–477. doi: 10.1124/jpet.120.000124 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 51. Savage C, McCabe OL. Residential psychedelic (LSD) therapy for the narcotic addict, a controlled study. *Arch Gen Psychiatry*. 1973;28(6):808–814. doi: 10.1001/archpsyc.1973.01750360040005 [PubMed] [CrossRef] [Google Scholar]
- 52. Stein MD, Kurth ME, Sharkey KM, Anderson BJ, Corso RP, Millman RP. Trazodone for sleep disturbance during methadone maintenance: a double-blind, placebo-controlled trial. *Drug Alcohol Depend*. 2012;120(1–3):65–73. doi: 10.1016/j.drugalcdep.2011.06.026 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 53. Akbarzadeh A, Norouzian D, Farhangi A, et al. Immunotherapy of 347 volunteer outpatient morphine addicts by human therapeutic morphine vaccine in Kermanshah Province of Iran. *J Pharmacol Toxicol*. 2009;4:30–35. doi: 10.3923/jpt.2009.30.35 [CrossRef] [Google Scholar]
- 54. Clinical Trials of Multivalent Opioid Vaccine Components. <u>ClinicalTrials.gov</u> identifier: <u>NCT04458545</u>. Updated July 5, 2022. Accessed August 20, 2022. https://clinicaltrials.gov/ct2/show/NCT04458545
- 55. Ekhtiari H, Tavakoli H, Addolorato G, et al. Transcranial electrical and magnetic stimulation (tES and TMS) for addiction medicine: a consensus paper on the present state of the science and the road ahead. *Neurosci Biobehav Rev*. 2019;104:118–140. doi: 10.1016/j.neubiorev.2019.06.007 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 56. Ford I, Norrie J. Pragmatic trials. N Engl J Med. 2016;375(5):454-463. doi: 10.1056/NEJMra1510059 [PubMed] [CrossRef] [Google Scholar]
- 57. Pallmann P, Bedding AW, Choodari-Oskooei B, et al. Adaptive designs in clinical trials: why use them, and how to run and report them. *BMC Med*. 2018;16(1):29. doi: 10.1186/s12916-018-1017-7 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 58. Dugosh K, Abraham A, Seymour B, McLoyd K, Chalk M, Festinger D. A systematic review on the use of psychosocial interventions in conjunction with medications for the treatment of opioid addiction. *J Addict Med.* 2016;10(2):93–103. doi: 10.1097/ADM.0000000000000193 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 59. Mauri L, D'Agostino RB Sr. Challenges in the design and interpretation of noninferiority trials. *N Engl J Med*. 2017;377(14):1357–1367. doi: 10.1056/NEJMra1510063 [PubMed] [CrossRef] [Google Scholar]
- 60. Uscher-Pines L, Huskamp HA, Mehrotra A. Treating patients with opioid use disorder in their homes: an emerging treatment model. *JAMA*. 2020;324(1):39–40. doi: 10.1001/jama.2020.3940 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 61. Ho C, Argáez C. *Telehealth-Delivered Opioid Agonist Therapy for the Treatment of Adults With Opioid Use Disorder: Review of Clinical Effectiveness, Cost-effectiveness, and Guidelines*. Canadian Agency for Drugs and Technologies in Health; 2018. [PubMed] [Google Scholar]
- 62. Dahne J, Tomko RL, McClure EA, Obeid JS, Carpenter MJ. Remote methods for conducting tobacco-focused clinical trials. *Nicotine Tob Res.* 2020;22(12):2134–2140. doi: 10.1093/ntr/ntaa105 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 63. Crowley D, Delargy I. A national model of remote care for assessing and providing opioid agonist treatment during the COVID-19 pandemic: a report. *Harm Reduct J*. 2020;17(1):49. doi: 10.1186/s12954-020-00394-z [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 64. Mitchell MM, Mendelson J, Gryczynski J, Carswell SB, Schwartz RP. A novel telehealth platform for alcohol use disorder treatment: preliminary evidence of reductions in drinking. *Am J Drug Alcohol Abuse*. 2020;46(3):297–303. doi: 10.1080/00952990.2019.1658197 [PubMed] [CrossRef] [Google Scholar]
- 65. Korthuis PT, McCarty D, Weimer M, et al. Primary care–based models for the treatment of opioid use disorder: a scoping review. *Ann Intern Med.* 2017;166(4):268–278. doi: 10.7326/M16-2149 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 66. Donroe JH, Bhatraju EP, Tsui JI, Edelman EJ. Identification and management of opioid use disorder in primary care: an update. *Curr Psychiatry Rep.* 2020;22(5):23. doi: 10.1007/s11920-020-01149-0 [PubMed] [CrossRef] [Google Scholar]
- 67. Hendricks PS, Copes H, Family N, Williams LT, Luke D, Raz S. Perceptions of safety, subjective effects, and beliefs about the clinical utility of lysergic acid diethylamide in healthy participants within a novel intervention paradigm: qualitative results from a proof-of-concept study. *J Psychopharmacol*. 2022;36(3):337–347. doi: 10.1177/02698811211055855 [PubMed] [CrossRef] [Google Scholar]
- 68. Family N, Hendricks PS, Williams LT, et al. Safety, tolerability, pharmacokinetics, and subjective effects of 50, 75, and 100 μg LSD in healthy participants within a novel intervention paradigm: a proof-of-concept study. *J Psychopharmacol*. 2022;36(3):321–336. doi: 10.1177/02698811211069103 [PubMed] [CrossRef] [Google Scholar]
- 69. Priester MA, Browne T, Iachini A, Clone S, DeHart D, Seay KD. Treatment access barriers and disparities among individuals with co-occurring mental health and substance use disorders: an integrative literature review. *J Subst Abuse Treat*. 2016;61:47–59. doi: 10.1016/j.jsat.2015.09.006 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 70. Olfson M, Zhang VS, Schoenbaum M, King M. Trends in buprenorphine treatment in the United States, 2009–2018. *JAMA*. 2020;323(3):276–277. doi: 10.1001/jama.2019.18913 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 71. Gordon AJ, Drexler K, Hawkins EJ, et al. Stepped Care for Opioid Use Disorder Train the Trainer (SCOUTT) initiative: expanding access to medication treatment for opioid use disorder within Veterans Health Administration facilities. Subst Abus. 2020;41(3):275–282. doi: 10.1080/08897077.2020.1787299 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 72. Short VL, Hand DJ, MacAfee L, Abatemarco DJ, Terplan M. Trends and disparities in receipt of pharmacotherapy among pregnant women in publically funded treatment programs for opioid use disorder in the United States. *J Subst Abuse Treat*. 2018;89:67–74. doi: 10.1016/j.jsat.2018.04.003 [PubMed] [CrossRef] [Google Scholar]
- 73. Eghaneyan BH, Sanchez K, Haeny AM, et al. Hispanic participants in the National Institute on Drug Abuse's Clinical Trials Network: a scoping review of two decades of research. *Addict Behav Rep.* 2020;12:100287. doi: 10.1016/j.abrep.2020.100287 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 74. Pinedo M A current re-examination of racial/ethnic disparities in the use of substance abuse treatment: do disparities persist? *Drug Alcohol Depend*. 2019;202:162–167. doi: 10.1016/j.drugalcdep.2019.05.017 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 75. Abraham AJ, Andrews CM, Yingling ME, Shannon J. Geographic disparities in availability of opioid use disorder treatment for Medicaid enrollees. *Health Serv Res.* 2018;53(1):389–404. doi: 10.1111/1475-6773.12686 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 76. Stahler GJ, Mennis J. Treatment outcome disparities for opioid users: are there racial and ethnic differences in treatment completion across large US metropolitan areas? *Drug Alcohol Depend*. 2018;190:170–178. doi: 10.1016/j.drugalcdep.2018.06.006 [PubMed] [CrossRef] [Google Scholar]
- 77. Biondi BE, Zheng X, Frank CA, Petrakis I, Springer SA. A literature review examining primary outcomes of medication treatment studies for opioid use disorder: what outcome should be used to measure opioid treatment success? *Am J Addict*. 2020;29(4):249–267. doi: 10.1111/ajad.13051 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 78. US Food and Drug Administration. *Opioid use disorder: endpoints for demonstrating effectiveness of drugs for treatment guidance for industry*. Accessed July 27, 2021. https://www.fda.gov/regulatory-information/search-fda-guidance-documents/opioid-use-disorder-endpointsdemonstrating-effectiveness-drugs-treatment-guidance-industry
- 79. Kiluk BD, Fitzmaurice GM, Strain EC, Weiss RD. What defines a clinically meaningful outcome in the treatment of substance use disorders: reductions in direct consequences of drug use or improvement in overall functioning? *Addiction*. 2019;114(1):9–15. doi: 10.1111/add.14289 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 80. Fitzmaurice GM, Lipsitz SR, Weiss RD. Within-treatment frequency of use versus abstinence as a predictor of longitudinal post-treatment follow-up assessments of drug use. *Drug Alcohol Depend*. 2020;208:107857. doi: 10.1016/j.drugalcdep.2020.107857 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 81. Timko C, Schultz NR, Cucciare MA, Vittorio L, Garrison-Diehn C. Retention in medication-assisted treatment for opiate dependence: a systematic review. *J Addict Dis*. 2016;35(1):22–35. doi: 10.1080/10550887.2016.1100960 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 82. Marsden J, Tai B, Ali R, Hu L, Rush AJ, Volkow N. Measurement-based care using DSM-5 for opioid use disorder: can we make opioid medication treatment more effective? *Addiction*. 2019;114(8):1346–1353. doi: 10.1111/add.14546 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 83. Kiluk BD, Frankforter TL, Cusumano M, Nich C, Carroll KM. Change in DSM-5 alcohol use disorder criteria count and severity level as a treatment outcome indicator: results from a randomized trial. *Alcohol Clin Exp Res.* 2018. doi: 10.1111/acer.13807 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 84. Cousins G, Teljeur C, Motterlini N, McCowan C, Dimitrov BD, Fahey T. Risk of drug-related mortality during periods of transition in methadone maintenance treatment: a cohort study. *J Subst Abuse Treat*, 2011;41(3):252–260. doi: 10.1016/j.jsat.2011.05.001 [PubMed] [CrossRef] [Google Scholar]
- 85. Sordo L, Barrio G, Bravo MJ, et al. Mortality risk during and after opioid substitution treatment: systematic review and meta-analysis of cohort studies. *BMJ*. 2017;357:j1550. doi: 10.1136/bmj.j1550 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 86. Dunn KE, Barrett FS, Yepez-Laubach C, et al. Brief Opioid Overdose Knowledge (BOOK): a questionnaire to assess overdose knowledge in individuals who use illicit or prescribed opioids. *J Addict Med*. 2016;10(5):314–323. doi: 10.1097/ADM.000000000000000035 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- 87. McLarnon ME, Monaghan TL, Stewart SH, Barrett SP. Drug misuse and diversion in adults prescribed anxiolytics and sedatives. *Pharmacotherapy*. 2011;31(3):262–272. doi: 10.1592/phco.31.3.262 [PubMed] [CrossRef] [Google Scholar]
- 88. Hasin DS, Sarvet AL, Cerdá M, et al. US adult illicit cannabis use, cannabis use disorder, and medical marijuana laws: 1991–1992 to 2012–2013. *JAMA Psychiatry*. 2017;74(6):579–588. doi: 10.1001/jamapsychiatry.2017.0724 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 89. Sellers EM, Leiderman DB. Psychedelic drugs as therapeutics: no illusions about the challenges. *Clin Pharmacol Ther*. 2018;103(4):561–564. doi: 10.1002/cpt.776

 [PubMed] [CrossRef] [Google Scholar]
- 90. Mitchell JM, Bogenschutz M, Lilienstein A, et al. MDMA-assisted therapy for severe PTSD: a randomized, double-blind, placebo-controlled phase 3 study. *Nat Med*. 2021;27(6):1025–1033. doi: 10.1038/s41591-021-01336-3 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 91. Comer SD, Pravetoni M, Coop A, Baumann MH, Cunningham CW. Potential unintended consequences of class-wide drug scheduling based on chemical structure: a cautionary tale for fentanyl-related compounds. *Drug Alcohol Depend*. 2021;221:108530. doi: 10.1016/j.drugalcdep.2021.108530 [CrossRef] [Google Scholar]
- 92. Palamar JJ, Ciccarone D, Rutherford C, Keyes KM, Carr TH, Cottler LB. Trends in seizures of powders and pills containing illicit fentanyl in the United States, 2018 through 2021. Drug Alcohol Depend. 2022;234:109398. doi: 10.1016/j.drugalcdep.2022.109398 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- 93. Brunetti P, Pirani F, Carlier J, Giorgetti R, Busardò FP, Lo Faro AFA. A 2017–2019 update on acute intoxications and fatalities from illicit fentanyl and analogs. *J Anal Toxicol*. 2021;45(6):537–554. doi: 10.1093/jat/bkaa115 [PubMed] [CrossRef] [Google Scholar]

JAMA Psychiatry | Original Investigation

Cannabis Use Disorder and Subsequent Risk of Psychotic and Nonpsychotic Unipolar Depression and Bipolar Disorder

Oskar Hougaard Jefsen, MD; Annette Erlangsen, PhD; Merete Nordentoft, DMSc; Carsten Hjorthøj, PhD

IMPORTANCE Cannabis use is increasing worldwide and is suspected to be associated with increased risk of psychiatric disorders; however, the association with affective disorders has been insufficiently studied.

OBJECTIVE To examine whether cannabis use disorder (CUD) is associated with an increased risk of psychotic and nonpsychotic unipolar depression and bipolar disorder and to compare associations of CUD with psychotic and nonpsychotic subtypes of these diagnoses.

DESIGN, SETTING, AND PARTICIPANTS This prospective, population-based cohort study using Danish nationwide registers included all individuals born in Denmark before December 31, 2005, who were alive, aged at least 16 years, and living in Denmark between January 1, 1995, and December 31, 2021.

EXPOSURE Register-based diagnosis of CUD.

MAIN OUTCOME AND MEASURES The main outcome was register-based diagnosis of psychotic or nonpsychotic unipolar depression or bipolar disorder. Associations between CUD and subsequent affective disorders were estimated as hazard ratios (HRs) using Cox proportional hazards regression with time-varying information on CUD, adjusting for sex; alcohol use disorder; substance use disorder; having been born in Denmark; calendar year; parental educational level (highest attained); parental cannabis, alcohol, or substance use disorders; and parental affective disorders.

RESULTS A total of 6 651765 individuals (50.3% female) were followed up for 119 526 786 person-years. Cannabis use disorder was associated with an increased risk of unipolar depression (HR, 1.84; 95% CI, 1.78-1.90), psychotic unipolar depression (HR, 1.97; 95% CI, 1.73-2.25), and nonpsychotic unipolar depression (HR, 1.83; 95% CI, 1.77-1.89). Cannabis use was associated with an increased risk of bipolar disorder in men (HR, 2.96; 95% CI, 2.73-3.21) and women (HR, 2.54; 95% CI, 2.31-2.80), psychotic bipolar disorder (HR, 4.05; 95% CI, 3.52-4.65), and nonpsychotic bipolar disorder in men (HR, 2.96; 95% CI, 2.73-3.21) and women (HR, 2.60; 95% CI, 2.36-2.85). Cannabis use disorder was associated with higher risk for psychotic than nonpsychotic subtypes of bipolar disorder (relative HR, 1.48; 95% CI, 1.21-1.81) but not unipolar depression (relative HR, 1.08; 95% CI, 0.92-1.27).

CONCLUSIONS AND RELEVANCE This population-based cohort study found that CUD was associated with an increased risk of psychotic and nonpsychotic bipolar disorder and unipolar depression. These findings may inform policies regarding the legal status and control of cannabis use.

Multimedia

Supplemental content

Author Affiliations: Author affiliations are listed at the end of this article.

Corresponding Author: Oskar Hougaard Jefsen, MD, Psychosis Research Unit, Aarhus University Hospital-Psychiatry, Palle Juul-Jensens Boulevard 175, 8200 Aarhus N, Denmark (oskar.jefsen@rm.dk).

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annabis is one of the most widely used psychoactive drugs in the world, and an increasing number of countries are legalizing its production and sale for medicinal and recreational use.² Over the past decades, both the use and the average potency of cannabis have increased.^{3,4} Use of cannabis may, however, lead to addiction (ie, cannabis use disorder [CUD]).5 Cannabis use disorder is frequent among individuals with affective disorders⁶ and, in this group, is associated with increased symptom severity,^{7,8} suicidality,⁹ and mortality.10 Although disputed, evidence suggests that use of cannabis may be associated with increased risk of developing psychiatric disorders¹¹; however, the association could also be reversed (ie, premorbid illness leading to cannabis use) or attributable to confounding (ie, common genetic liability for cannabis use and psychiatric disorders¹²). Mendelian randomization studies, which use genetic variants as instrumental variables to infer causal relationships, suggest a causal effect of cannabis use on schizophrenia¹³ but not on bipolar disorder¹⁴ or major depressive disorder, although this may be due to lack of statistical power. 15,16 The accumulating epidemiologic evidence, which supports an association between cannabis use and psychosis, 17,18 includes dose-response relationships 19 and a positive association between cannabis potency (Δ9tetrahydrocannabinol concentration) and risk of psychosis.²⁰ When taking the increased use and potency of cannabis into consideration, an increased incidence of schizophrenia may be expected. The incidence of schizophrenia²¹ and the population-attributable risk fraction (PARF) of CUD for schizophrenia²² have increased over recent years. Based on the existing evidence, it is possible that cannabis use may be associated with risks of other mental disorders, such as affective disorders.

Evidence regarding the association between use of cannabis and affective disorders is limited. Self-reported cannabis use was not found to be associated with unipolar depression or bipolar disorder after adjustment for confounders in a sample of Swedish military conscripts aged 18 to 20 years, ²³ although a dose-dependent association with the risk of schizophrenia was identified. 24 Similarly, no association was found between cannabis use and subsequent risk of affective disorders in a nationally representative sample of US adults. 5 However, a positive association between cannabis use and subsequent depression, 25 bipolar disorder, 26 and manic symptoms 27 has been demonstrated in other longitudinal studies. Risk estimates may be smaller for the association between cannabis use and affective disorders than estimates for the association between cannabis use and schizophrenia. ^{23,24,28-30} It is possible that the effects of cannabis might primarily be psychotogenic, in which case, higher risk of psychotic (vs nonpsychotic) subtypes of affective disorders would be expected. Still, this hypothesis remains to be tested.

The aim of the current study was to analyze whether CUD was associated with a subsequent diagnosis of unipolar depression or bipolar disorder. To assess whether an association was primarily psychotogenic, we conducted separate analyses with respect to psychotic and nonpsychotic subtypes of these affective disorders. These questions were studied using longitudinal data from nationwide Danish health registers.

Key Points

Question Is cannabis use disorder associated with an increased risk of psychotic and nonpsychotic unipolar depression and bipolar disorder?

Findings In this cohort study of 6 651765 individuals in Demark, cannabis use disorder was associated with an increased risk of both psychotic and nonpsychotic unipolar depression and bipolar disorder.

Meaning The findings suggest that cannabis use disorder is independently associated with bipolar disorder and unipolar depression.

Methods

Study Design, Data Sources, and Study Population

We conducted a register-based prospective cohort study by linking nationwide Danish register data. Since 1968, the Danish Civil Registration System³¹ has provided all permanent residents in Denmark with a unique identification number, which allows for individual-level linkage of data from different registers. The Civil Registration System³¹ also contains information on date of birth, birthplace, and vital status. Data on contacts with psychiatric and somatic hospitals, including information on diagnoses, were obtained from the Psychiatric Central Research Register since 1969³² and the National Patient Register since 1977,33 respectively. Finally, we obtained data on treatments provided for substance use from the municipal Register of Substance Abusers in Treatment.³⁴ Information on redeemed prescriptions was derived from the National Prescription Registry³⁵ since 1995. We included all individuals born no later than December 31, 2005, who were alive, aged at least 16 years, and living in Denmark at some point between January 1, 1995, and December 31, 2021. The study was approved by the Danish Data Protection Agency. Register-based studies do not require informed consent according to Danish law. The present analyses were conducted using encrypted personal identification numbers on servers at Statistics Denmark. The study followed the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guideline.

Exposures

Cannabis use disorder was defined as a recorded diagnosis during a hospital contact in either the Psychiatric Central Research Register or the National Patient Register or a record of treatment for CUD provided by a municipality. Diagnoses of CUD were recorded using International Classification of Diseases, Eighth Revision (ICD-8) code 304.5 and International Statistical Classification of Diseases and Related Health Problems, Tenth Revision (ICD-10) code F12.X. The ICD-8 was used in Denmark until 1994, when it was replaced by the ICD-10. The International Classification of Diseases, Ninth Revision was never implemented in Denmark. Information on CUD was also obtained from the municipal register of substance users seeking treatment, with information as to whether cannabis was the person's primary misused substance.

Covariates

We obtained information on alcohol use disorder (AUD) and substance use disorders (SUDs) using psychiatric diagnostic codes

(eTable 1 in Supplement 1) and registered treatment in the municipal register of substance users seeking treatment. Finally, redeeming a prescription of naltrexone counted toward AUD, while buprenorphine or methadone counted toward SUD.

In addition to the aforementioned data, we obtained information on sex; date of birth; country of birth; parental CUD, AUD, and SUD; parental affective disorders; and highest level of parental education. In addition, history of other psychiatric disorders (*ICD-8*: 290-309; *ICD-10*: any code in the F chapter except for those already part of other variables) was included.

Outcomes

Information on affective disorders was obtained from the Psychiatric Central Research Register³² and the psychiatric segment of the National Patient Registry.³⁶ To distinguish affective disorders with and without psychotic features, we restricted the study period to the years when *ICD-10* codes were used. The following *ICD-10* codes were used for the outcome categories: unipolar depression (F32.X or F33.X), unipolar depression with psychotic features (F32.3 or F33.3), unipolar depression without psychotic features (F32 and F33, excluding F32.3 and F33.3), bipolar disorder (F31.X), bipolar disorder with psychotic features (F31.2 or F31.5), and bipolar disorder without psychotic features (F31, excluding F31.2 and F31.5).

Statistical Analysis

We plotted cumulative probabilities for affective disorders using Kaplan-Meier curves and applied Cox proportional hazards regression to estimate hazard ratios (HRs) to compare the risk of affective disorders depending on the exposure (CUD vs no CUD). Individuals were entered into the analysis on their 16th birthday or January 1, 1995, whichever came last. Individuals were followed up until development of an affective disorder; censoring due to development of schizophrenia, death, or emigration; or the end of data collection on May 3, 2022. We included CUD, AUD, and SUD as time-varying covariates in all models. Men and women were examined separately if an interaction between sex and CUD was detected in crude preliminary analyses. In the adjusted analyses, we included sex (if not stratified by sex); AUD and SUD; born in Denmark (yes, no); calendar year; parental educational level (highest attained); parental CUD, AUD, and SUD; and parental affective disorders. Age was used as the underlying time scale in all analyses. When calculating risk of unipolar depression, individuals were censored at the date of diagnosis of bipolar disorder as this diagnosis would preclude a later unipolar depression diagnosis. Individuals who had been diagnosed with an affective disorder (ICD-8: 296.X) prior to 1995 were not considered to have incident cases of unipolar depression or bipolar disorder and were thus censored before inclusion in the analyses. We estimated relative HRs for associations between CUD and the psychotic and nonpsychotic subtypes by dividing the 2 HRs. The SE for this metric was estimated by summing the nonexponentially transformed SEs of the 2 estimated HRs, and this was then used to estimate a 95% CI around the relative HR. We conducted 2 sensitivity analyses to address potential confounding by other psychiatric disorders; we adjusted for the presence of other psychiatric disorders (1) prior to CUD diagnosis and (2) over the entire follow-up period. We estimated PARFs from the adjusted HRs as previously reported.²²

Table 1. Characteristics of the Population

Characteristic	Individuals, No. (%) (N = 6 651 765)	
Sex		
Female	3 347 142 (50.3)	
Male	3 304 623 (49.7)	
Born in Denmark	521 840 (7.9)	
Parental CUD, AUD, and/or SUD	666 427 (10.0)	
Parental affective disorder	313 305 (4.7)	
Parental educational level		
Primary or lower secondary	1 145 564 (17.2)	
Upper secondary	1 095 498 (16.5)	
Short-cycle tertiary	75 650 (1.1)	
Bachelor's degree	297 728 (4.5)	
Master's degree or higher	107 305 (1.6)	
Not registered	3 930 020 (59.1)	
CUD	55 968 (0.8)	
AUD	399 086 (6.0)	
SUD	214 110 (3.2)	

Abbreviations: AUD, alcohol use disorder; CUD, cannabis use disorder; SUD. substance use disorder.

All analyses were conducted using STATA/MP, version 17.0 (StataCorp LLC). Two-sided *P* < .05 was considered significant.

Results

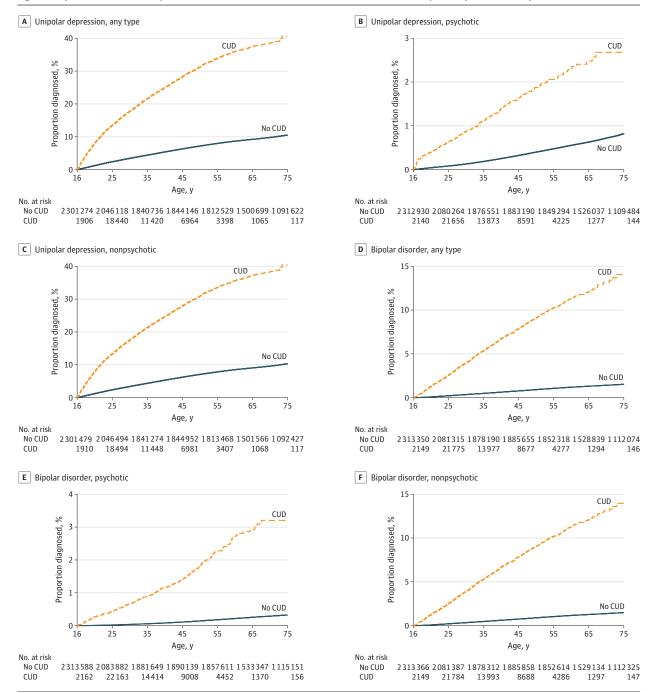
A total of 6 651 765 individuals were included and followed up over 119 526 786 person-years (50.3% female; 49.7% male). Table 1 presents study population characteristics, all of which were significantly associated with the outcomes. The study population had a broad age distribution, as shown in the eFigure in Supplement 1. In all, 60 696 individuals (0.9% of the study population) received a diagnosis of CUD during followup, and 260 746 (3.9%) developed an affective disorder.

CUD and Unipolar Depression

All analyses regarding unipolar depression were conducted jointly for men and women as no interaction between sex and CUD was observed (any type of unipolar depression: χ_1^2 , 1.01; P=.03; psychotic unipolar depression: χ_1^2 , 0.43; P=.51; non-psychotic unipolar depression: χ_1^2 , 1.37; P=.24). Altogether, 40.7% of individuals with CUD received a diagnosis of unipolar depression, as shown in the Kaplan-Meier plot (**Figure 1**). The majority of these individuals (96.1%) were diagnosed with nonpsychotic unipolar depression, while 3.9% were diagnosed with psychotic unipolar depression.

When adjusting for sex; AUD and SUD; having been born in Denmark; calendar year; parental CUD, AUD, and SUD; and parental affective disorders, individuals with CUD had a higher risk of any type of unipolar depression (HR, 1.84; 95% CI, 1.78-1.90) compared with individuals with no records of a CUD (Table 2). Elevated risks were also found with respect to psychotic depression (HR, 1.97; 95% CI, 1.73-2.25) and nonpsychotic depression (HR, 1.83; 95% CI, 1.77-1.89). We found no statistically significant difference in the associations between CUD and the psychotic vs nonpsychotic type of unipolar depression (relative HR, 1.08; 95%

Figure 1. Kaplan-Meier Plots for Exposure to Cannabis Use Disorder (CUD) and the Outcomes of Unipolar Depression and Bipolar Disorder



CI, 0.92-1.27). The PARFs for unipolar depression associated with CUD ranged from 0.71% (95% CI, 0.69%-0.73%) to 0.85% (95% CI, 0.84%-0.87%). The HRs for the associations between AUD or SUD and any type of unipolar depression were found to be nominally greater than those for the association between CUD and any type of unipolar depression (eTable 2 in Supplement 1).

When assessing risks with respect to the time between the first diagnosis of CUD and subsequent unipolar depression, the highest risk was found within the first 6 months of being diagnosed (HR, 6.84; 95% CI, 6.34-7.38) compared with no

diagnosis of CUD (**Figure 2**). However, the excess risk of unipolar depression among those with CUD remained significant up to 10 years after the initial diagnosis.

In the sensitivity analyses adjusting for other psychiatric disorders prior to CUD, associations remained between CUD and unipolar depression (HR, 1.72; 95% CI, 1.67-1.77) and the psychotic (HR, 1.87; 95% CI, 1.65-2.13) and nonpsychotic (HR, 1.71; 95% CI, 1.65-1.76) subtypes, but the HRs were smaller than in the main analysis. After adjusting for other psychiatric disorders over the entire follow-up period (both before and after

Table 2. Associations of CUD With Unipolar Depression and Bipolar Disorder and PARFs

Outcome	Incident cases, No.	HR (95% CI)		
		Unadjusted	Adjusted ^a	PARF, % (95% CI) ^b
Unipolar depression				
Any type	240 347	4.89 (4.75-5.04)	1.84 (1.78-1.90)	0.85 (0.84-0.86)
Psychotic	17 906	4.72 (4.05-5.51)	1.97 (1.73-2.25)	0.71 (0.69-0.73)
Nonpsychotic	235 157	4.91 (4.77-5.06)	1.83 (1.77-1.89)	0.85 (0.84-0.87)
Bipolar disorder				
Any type ^c				
Males	12 545	11.36 (10.60-12.17)	2.96 (2.73-3.21)	4.72 (4.58-4.86)
Females	19 330	11.94 (10.96-13.02)	2.54 (2.31-2.80)	1.68 (1.63-1.73)
Psychotic	6567	12.26 (10.86-13.84)	4.05 (3.52-4.65)	3.22 (3.03-3.41)
Nonpsychotic ^c				
Males	12 198	11.51 (10.74-12.35)	2.96 (2.73-3.21)	4.79 (4.64-4.93)
Females	18 907	12.32 (11.30-13.43)	2.60 (2.36-2.85)	1.76 (1.71-1.82)

Abbreviations: CUD, cannabis use disorder; HR, hazard ratio; PARF, population-attributable risk fraction.

CUD), HRs were even smaller for associations with unipolar depression (HR, 1.08; 95% CI, 1.04-1.11) and the nonpsychotic subtype (HR, 1.07; 95% CI, 1.04-1.10), and there was no association with the psychotic subtype (HR, 1.05; 95% CI, 0.92-1.19).

Cannabis Use Disorder and Bipolar Disorder

We found an interaction between sex and CUD for any type of bipolar disorder (χ_1^2 , 5.02; P = .03) and nonpsychotic bipolar disorder (χ_1^2 , 6.62; P = .01) but not for psychotic bipolar disorder (χ_1^2 , 0.43; P = .51). Analyses for the first 2 outcomes were thus stratified by sex.

The Kaplan-Meier curves revealed that 14.1% of individuals with CUD eventually received a diagnosis of bipolar disorder (Figure 1). The majority of these individuals (90.2%) were diagnosed with nonpsychotic bipolar disorder, while 9.8% were diagnosed with psychotic bipolar disorder.

Cannabis use disorder was associated with a higher risk of any type of bipolar disorder among both men (HR, 2.96; 95% CI, 2.73-3.21) and women (HR, 2.54; 95% CI, 2.31-2.80) compared with nonexposed individuals in the adjusted analysis (Table 2). Likewise, CUD was associated with psychotic bipolar disorder (HR, 4.05; 95% CI, 3.52-4.65) and with nonpsychotic bipolar disorder in both men (HR, 2.96; 95% CI, 2.73-3.21) and women (HR, 2.60; 95% CI, 2.36-2.85). Cannabis use disorder was associated with a higher risk for the psychotic type than the nonpsychotic type of bipolar disorder (relative HR, 1.48; 95% CI, 1.21-1.81). The PARF for bipolar disorder varied from 1.68% (95% CI, 1.63%-1.73%) for any type of bipolar disorder in women to 4.79% (95% CI, 4.64-4.93) for nonpsychotic bipolar disorder in men (Table 2). Alcohol use disorder was associated with a nominally greater risk of bipolar disorder compared with CUD (eTable 3 in Supplement 1).

When assessing risks with respect to the time between first diagnosis of CUD and subsequent bipolar disorder, the highest risk was found within the first 6 months of diagnosis (HR, 16.45; 95% CI, 13.97-19.38) compared with no diagnosis of CUD

(Figure 2). However, the risk of bipolar disorder among those with CUD remained elevated even after 10 or more years (Figure 2).

In the sensitivity analyses, after adjustment for other psychiatric disorders prior to CUD, associations remained between CUD and bipolar disorder in both men (HR, 2.79; 95% CI, 2.59-3.02) and women (HR, 2.46; 95% CI, 2.24-2.69), but HRs were smaller. The HR was similar for the association between CUD and psychotic bipolar disorder (HR, 4.04; 95% CI, 3.54-4.60). For nonpsychotic bipolar disorder, after adjustment for other psychiatric disorders prior to CUD, the associations remained for both men (HR, 2.79; 95% CI, 2.58-3.01) and women (HR, 2.50; 95% CI, 2.28-2.40), but the HRs were smaller. After adjusting for other psychiatric disorders over the entire follow-up period (both before and after CUD), HRs were even smaller for the associations between CUD and bipolar disorder in men (HR, 1.49; 95% CI, 1.38-1.61) and women (HR, 1.53; 95% CI, 1.39-1.67), between CUD and psychotic bipolar disorder (HR, 2.07, 95% CI, 1.82-2.36), and between CUD and nonpsychotic bipolar disorder in men (HR, 1.48; 95% CI, 1.37-1.60) and women (HR, 1.55; 95% CI, 1.41-1.69).

Discussion

In this nationwide cohort study of 6 651 765 individuals, CUD was found to be associated with an increased risk of unipolar depression and bipolar disorder when adjusting for relevant confounders. Although excess risks of unipolar depression and bipolar disorder were highest immediately after diagnosis of CUD, they remained significantly elevated up to 5 to 10 years after CUD.

Our findings add support to previous large-scale studies showing an association between CUD and affective disorders. ^{23,26,27} Two previous studies found significant associations between cannabis use and unipolar depression but not bipolar disorder. ^{23,26} We found significant associations between CUD and both bipolar disorder and unipolar depression, but

^a Adjusted for sex (if not stratified by sex); alcohol use disorder; substance use disorder; born in Denmark (yes, no); calendar year; parental educational level (highest attained); parental CUD, alcohol use disorder, and substance use disorder; and parental affective disorders. The associations were conditioned on age since age was used as the underlying time scale in all analyses.

^b Calculated from the estimates of the adjusted HRs.

^c Estimates for the associations are reported for males and females separately as there was a significant interaction between CUD and sex for these outcomes. There was no interaction between sex and CUD for psychotic bipolar disorder or for any of the outcomes under unipolar depression.

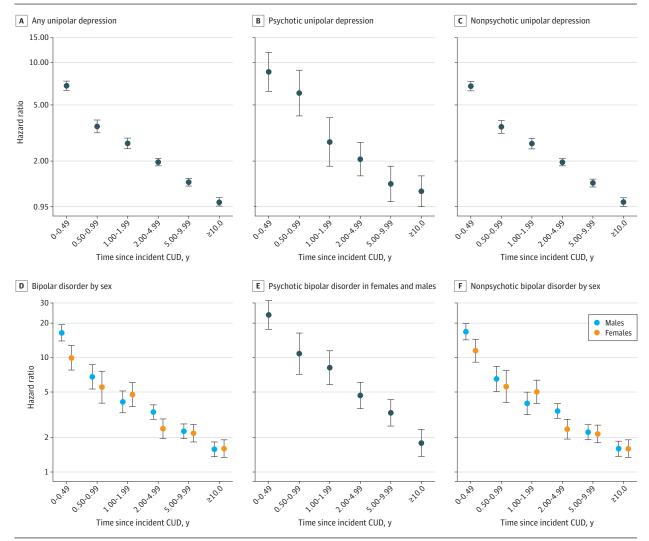


Figure 2. Risk of Unipolar Depression and Bipolar Disorder Among Individuals With vs Without Cannabis Use Disorder (CUD)

Whiskers indicate 95% Cls. D and F, Due to a significant interaction between sex and CUD for any type of bipolar disorder and psychotic bipolar disorder, these analyses were performed for males and females separately.

the risk of bipolar disorder was nominally higher. Importantly, differences in the information on cannabis use (self-reported use vs nationwide health records) and analytical strategies may explain some of these discrepancies. Specifically, some studies adjusted for baseline depressive or manic symptoms²⁷ or baseline psychiatric disorders.^{23,26} When we adjusted for other psychiatric disorders prior to CUD, the associations with mood disorders remained. After adjustment for other psychiatric disorders over the entire follow-up period to reduce potential residual confounding, associations remained with the exception of the association between CUD and psychotic unipolar depression. Adjustment for psychiatric disorders diagnosed after CUD may, however, induce collider stratification bias by conditioning on mediators between the exposure and the outcome; thus, the latter analysis may be overadjusted.³⁷

Implications

Our findings lend support to the notion that cannabis use may represent an independent factor associated with unipolar depression

and bipolar disorder. The risk of psychiatric disorders appears to be higher for schizophrenia 18,22 than for affective disorders 38 and higher for psychotic bipolar disorder than for nonpsychotic bipolar disorder, potentially pointing to a primarily psychotogenic effect of cannabis. $\Delta 9$ -Tetrahydrocannabinol, the main psychoactive constituent of cannabis, acts on cannabinoid (CBI) receptors and is suggested to increase the risk of psychosis by altering striatal dopaminergic function 39,40 or by disrupting normal endocannabinoid modulation of cortical development and function. 41,42 In addition to its links with psychosis, the dopaminergic system is intricately linked with neurocognitive processes relevant for affective disorders, such as reward processing. $^{43-45}$ However, a coherent model for how cannabis may influence the development of affective disorders is lacking. Future studies may further elucidate these effects in a transdiagnostic framework.

Based on our findings and the evidence regarding cannabis and schizophrenia, interventions to reduce cannabis use through both public education and more targeted interventions may be advisable. Direct evidence that cannabis cessation can reduce

the risk of affective disorders is, however, lacking, and although several interventions appear to be associated with reducing cannabis use in adolescents ⁴⁶ and healthy adults, ⁴⁷ they may be less effective in individuals with mental disorders. ⁴⁸ Although some trials have demonstrated significant improvements in depressive symptoms after a psychosocial intervention to reduce cannabis use, ⁴⁹ these improvements may be mediated by broader effects of the psychosocial interventions, providing little evidence for the beneficial effects of cannabis cessation itself. ⁵⁰ Targeted interventions for at-risk individuals are currently hindered by sparse knowledge on factors associated with transition from cannabis use (disorder) to psychiatric disorders, ^{51,52} calling for further studies.

Strengths and Limitations

A strength of this study is the large sample size, which makes it, to our knowledge, the largest investigation of the association between CUD and affective disorders to date. Data were collected prospectively and uniformly for all studied groups, eliminating recall bias and reducing selection bias. The availability of sociodemographic and historic psychiatric information on individuals and their parents enabled us to adjust for relevant confounders.

Important limitations should be mentioned. First, while individuals registered with a CUD diagnosis are likely to have CUD (ie, high positive predictive value), individuals without a register-based diagnosis of CUD may still have CUD (ie, suboptimal negative predictive value). This misclassification could bias our find-

ings toward the null if the misclassification was random or could confound our findings if individuals with a diagnosis of CUD were not representative of (heavy) cannabis users. ^{52,53} Second, the validity of the register-based diagnosis of affective disorders is evaluated as good in Denmark, ⁵⁴ but individuals with mild to moderate depression might be seen only in primary care and thus were not detected in our study. ⁵⁵ Third, detection bias is possible. Receiving a diagnosis and clinical care for CUD may imply that clinicians divert more attention to these individuals and, hence, are more likely to detect psychiatric disorders that might otherwise go undetected among nonexposed individuals. This could be an explanatory factor for the increased risk during the first year(s) after diagnosis of CUD. However, the sustained increased risk observed up to 10 years after the initial CUD diagnosis supports the notion of an association beyond the putative detection bias.

Conclusions

The results of this cohort study suggest that cannabis use is associated with an increased risk of psychotic and nonpsychotic bipolar disorder and unipolar depression. These findings have implications regarding the legalization and control of cannabis use. Importantly, there appears to be a need for improved knowledge on the dose-dependent effects of cannabis use on brain, cognition, and behavior; identification of risk factors for transition from cannabis use (disorder) to psychiatric disorders; and the effects of cannabis cessation on long-term psychiatric risk.

ARTICLE INFORMATION

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Author Affiliations: Psychosis Research Unit, Aarhus University Hospital-Psychiatry, Aarhus, Denmark (Jefsen); Center of Functionally Integrative Neuroscience, Department of Clinical Medicine, Aarhus University, Aarhus, Denmark (Jefsen); Copenhagen Research Center for Mental Health-CORE, Mental Health Centre Copenhagen, Copenhagen University Hospital, Copenhagen, Denmark (Erlangsen, Nordentoft, Hiorthøi): Danish Research Institute for Suicide Prevention, Mental Health Centre Copenhagen, Copenhagen, Denmark (Erlangsen); Department of Mental Health, Johns Hopkins Bloomberg School of Public Health, Baltimore, Maryland (Erlangsen): Centre for Mental Health Research, Research School of Population Health, The Australian National University, Canberra. Australia (Erlangsen); Faculty of Health and Medical Sciences, Department of Clinical Medicine, University of Copenhagen, Copenhagen, Denmark (Nordentoft); Section of Epidemiology, Department of Public Health, University of Copenhagen, Copenhagen, Denmark (Hjorthøj)

Author Contributions: Dr Hjorthøj had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. *Concept and design:* Jefsen, Nordentoft, Hjorthøj. *Acquisition, analysis, or interpretation of data:* All authors.

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Critical revision of the manuscript for important intellectual content: Jefsen, Nordentoft, Hjorthøj. Statistical analysis: Nordentoft, Hjorthøj. Obtained funding: Nordentoft. Supervision: Nordentoft.

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REFERENCES

- 1. United Nations Office on Drugs Crime. World drug report 2019. Accessed January 10, 2023. https://www.un-ilibrary.org/content/publication/a4dd519a-en
- 2. Hall W, Stjepanović D, Caulkins J, et al. Public health implications of legalising the production and sale of cannabis for medicinal and recreational use. *Lancet*. 2019;394(10208):1580-1590. doi:10.1016/S0140-6736(19)31789-1
- 3. Chandra S, Radwan MM, Majumdar CG, Church JC, Freeman TP, ElSohly MA. New trends in cannabis potency in USA and Europe during the last decade (2008-2017). Eur Arch Psychiatry Clin Neurosci. 2019; 269(1):5-15. doi:10.1007/s00406-019-00983-5
- 4. Rømer Thomsen K, Lindholst C, Thylstrup B, et al. Changes in the composition of cannabis from 2000-2017 in Denmark: analysis of confiscated samples of cannabis resin. *Exp Clin Psychopharmacol.* 2019;27(4):402-411. doi:10.1037/pha0000303
- **5**. Blanco C, Hasin DS, Wall MM, et al. Cannabis use and risk of psychiatric disorders: prospective evidence from a US national longitudinal study. *JAMA Psychiatry*. 2016;73(4):388-395. doi:10.1001/jamapsychiatry.2015.3229
- **6**. Toftdahl NG, Nordentoft M, Hjorthøj C. Prevalence of substance use disorders in psychiatric

- patients: a nationwide Danish population-based study. *Soc Psychiatry Psychiatr Epidemiol*. 2016;51 (1):129-140. doi:10.1007/s00127-015-1104-4
- 7. Gibbs M, Winsper C, Marwaha S, Gilbert E, Broome M, Singh SP. Cannabis use and mania symptoms: a systematic review and meta-analysis. *J Affect Disord*. 2015;171:39-47. doi:10.1016/j.jad.2014.09.016
- 8. Schoeler T, Petros N, Di Forti M, et al. Effects of continuation, frequency, and type of cannabis use on relapse in the first 2 years after onset of psychosis: an observational study. *Lancet Psychiatry*. 2016;3(10):947-953. doi:10.1016/S2215-0366(16) 30188-2
- Østergaard MLD, Nordentoft M, Hjorthøj C. Associations between substance use disorders and suicide or suicide attempts in people with mental illness: a Danish nation-wide, prospective, register-based study of patients diagnosed with schizophrenia, bipolar disorder, unipolar depression or personality disorder. Addiction. 2017;112(7):1250-1259. doi:10.1111/add.13788
- 10. Hjorthøj C, Østergaard MLD, Benros ME, et al. Association between alcohol and substance use disorders and all-cause and cause-specific mortality in schizophrenia, bipolar disorder, and unipolar depression: a nationwide, prospective, register-based study. *Lancet Psychiatry*. 2015;2(9): 801-808. doi:10.1016/S2215-0366(15)00207-2
- 11. Ganesh S, D'Souza DC. Cannabis and psychosis: recent epidemiological findings continuing the "causality debate." *Am J Psychiatry*. 2022;179(1):8-10. doi:10.1176/appi.ajp.2021.21111126
- 12. Johnson EC, Demontis D, Thorgeirsson TE, et al; Psychiatric Genomics Consortium Substance Use Disorders Workgroup. A large-scale genome-wide association study meta-analysis of cannabis use

- disorder. *Lancet Psychiatry*. 2020;7(12):1032-1045. doi:10.1016/S2215-0366(20)30339-4
- **13.** Gage SH, Jones HJ, Burgess S, et al. Assessing causality in associations between cannabis use and schizophrenia risk: a two-sample Mendelian randomization study. *Psychol Med.* 2017;47(5):971-980. doi:10.1017/S0033291716003172
- **14.** Jefsen OH, Speed M, Speed D, Østergaard SD. Bipolar disorder and cannabis use: a bidirectional two-sample Mendelian randomization study. *Addict Biol.* 2021;26(6):e13030. doi:10.1111/adb.13030
- 15. Hodgson K, Coleman JRI, Hagenaars SP, et al; Major Depressive Disorder Working Group of the Psychiatric Genomics Consortium. Cannabis use, depression and self-harm: phenotypic and genetic relationships. *Addiction*. 2020;115(3):482-492. doi:10.1111/add.14845
- **16.** Treur JL, Demontis D, Smith GD, et al. Investigating causality between liability to ADHD and substance use, and liability to substance use and ADHD risk, using Mendelian randomization. *Addict Biol.* 2021;26(1):e12849. doi:10.1111/adb.12849
- **17.** Gage SH, Hickman M, Zammit S. Association between cannabis and psychosis: epidemiologic evidence. *Biol Psychiatry*. 2016;79(7):549-556. doi:10.1016/j.biopsych.2015.08.001
- **18.** Nielsen SM, Toftdahl NG, Nordentoft M, Hjorthøj C. Association between alcohol, cannabis, and other illicit substance abuse and risk of developing schizophrenia: a nationwide population based register study. *Psychol Med*. 2017;47(9): 1668-1677. doi:10.1017/S0033291717000162
- 19. Marconi A, Di Forti M, Lewis CM, Murray RM, Vassos E. Meta-analysis of the association between the level of cannabis use and risk of psychosis. *Schizophr Bull.* 2016;42(5):1262-1269. doi:10.1093/schbul/sbw003
- 20. Di Forti M, Sallis H, Allegri F, et al. Daily use, especially of high-potency cannabis, drives the earlier onset of psychosis in cannabis users. Schizophr Bull. 2014;40(6):1509-1517. doi:10.1093/schbul/sbt181
- **21.** Kühl JOG, Laursen TM, Thorup A, Nordentoft M. The incidence of schizophrenia and schizophrenia spectrum disorders in Denmark in the period 2000-2012: a register-based study. *Schizophr Res.* 2016;176(2-3):533-539. doi:10.1016/j.schres.2016. 06.023
- **22.** Hjorthøj C, Posselt CM, Nordentoft M. Development over time of the population-attributable risk fraction for cannabis use disorder in schizophrenia in Denmark. *JAMA Psychiatry*. 2021;78(9):1013-1019. doi:10.1001/jamapsychiatry. 2021;471
- 23. Manrique-Garcia E, Zammit S, Dalman C, Hemmingsson T, Allebeck P. Cannabis use and depression: a longitudinal study of a national cohort of Swedish conscripts. *BMC Psychiatry*. 2012;12(1): 112. doi:10.1186/1471-244X-12-112
- 24. Manrique-Garcia E, Zammit S, Dalman C, Hemmingsson T, Andreasson S, Allebeck P. Cannabis, schizophrenia and other non-affective psychoses: 35 years of follow-up of a population-based cohort. *Psychol Med*. 2012;42(6): 1321-1328. doi:10.101/S0033291711002078
- 25. Lev-Ran S, Roerecke M, Le Foll B, George TP, McKenzie K, Rehm J. The association between cannabis use and depression: a systematic review and meta-analysis of longitudinal studies. *Psychol Med*. 2014;44(4):797-810. doi:10.1017/S0033291713001438

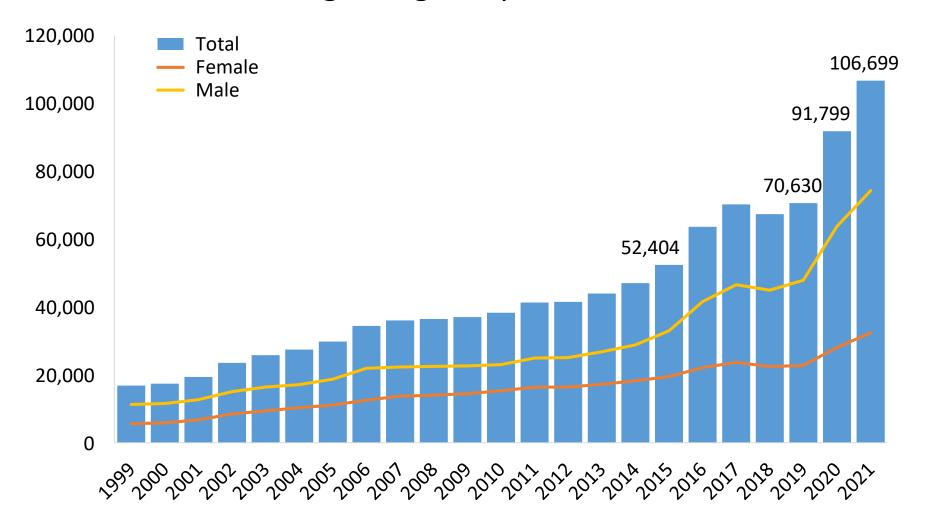
- **26**. Feingold D, Weiser M, Rehm J, Lev-Ran S. The association between cannabis use and mood disorders: a longitudinal study. *J Affect Disord*. 2015;172:211-218. doi:10.1016/j.jad.2014.10.006
- **27**. Henquet C, Krabbendam L, de Graaf R, ten Have M, van Os J. Cannabis use and expression of mania in the general population. *J Affect Disord*. 2006;95(1-3):103-110. doi:10.1016/j.jad.2006.05.002
- **28**. Arseneault L, Cannon M, Poulton R, Murray R, Caspi A, Moffitt TE. Cannabis use in adolescence and risk for adult psychosis: longitudinal prospective study. *BMJ*. 2002;325(7374):1212-1213. doi:10.1136/bmj.325.7374.1212
- 29. McGee R, Williams S, Poulton R, Moffitt T. A longitudinal study of cannabis use and mental health from adolescence to early adulthood. *Addiction*. 2000;95(4):491-503. doi:10.1046/j.1360-0443.2000.9544912.x
- **30**. Fergusson DM, Horwood LJ, Swain-Campbell N. Cannabis use and psychosocial adjustment in adolescence and young adulthood. *Addiction*. 2002; 97(9):1123-1135. doi:10.1046/j.1360-0443.2002. 00103.x
- **31**. Pedersen CB. The Danish Civil Registration System. *Scand J Public Health*. 2011;39(7)(suppl): 22-25. doi:10.1177/1403494810387965
- **32.** Mors O, Perto GP, Mortensen PB. The Danish Psychiatric Central Research Register. *Scand J Public Health*. 2011;39(7)(suppl):54-57. doi:10.1177/1403494810395825
- 33. Lynge E, Sandegaard JL, Rebolj M. The Danish National Patient Register. *Scand J Public Health*. 2011;39(7)(suppl):30-33. doi:10.1177/ 1403494811401482
- **34.** Ministry of Children and Social Affairs. Bekendtgørelse om dataindberetninger på socialområdet. Executive order on data reporting on social services. 2018. Accessed April 12, 2023. https://www.retsinformation.dk/eli/lta/2021/2351
- **35.** Kildemoes HW, Sørensen HT, Hallas J. The Danish National Prescription Registry. *Scand J Public Health*. 2011;39(7)(suppl):38-41. doi:10.1177/1403494810394717
- **36.** Schmidt M, Schmidt SAJ, Sandegaard JL, Ehrenstein V, Pedersen L, Sørensen HT. The Danish National Patient Registry: a review of content, data quality, and research potential. *Clin Epidemiol*. 2015;7:449-490. doi:10.2147/CLEP.S91125
- **37**. VanderWeele TJ. Principles of confounder selection. *Eur J Epidemiol*. 2019;34(3):211-219. doi:10.1007/s10654-019-00494-6
- **38**. Moore TH, Zammit S, Lingford-Hughes A, et al. Cannabis use and risk of psychotic or affective mental health outcomes: a systematic review. *Lancet*. 2007;370(9584):319-328. doi:10.1016/S0140-6736 (07)(618-3)
- **39**. Murray RM, Mehta M, Di Forti M. Different dopaminergic abnormalities underlie cannabis dependence and cannabis-induced psychosis. *Biol Psychiatry*. 2014;75(6):430-431. doi:10.1016/j. biopsych.2014.01.011
- **40**. Colizzi M, Weltens N, McGuire P, et al. Delta-9-tetrahydrocannabinol increases striatal glutamate levels in healthy individuals: implications for psychosis. *Mol Psychiatry*. 2020;25(12):3231-3240. doi:10.1038/s41380-019-0374-8
- **41**. Bara A, Ferland JN, Rompala G, Szutorisz H, Hurd YL. Cannabis and synaptic reprogramming of the developing brain. *Nat Rev Neurosci*. 2021;22(7): 423-438. doi:10.1038/s41583-021-00465-5

- **42**. Volk DW, Lewis DA. The role of endocannabinoid signaling in cortical inhibitory neuron dysfunction in schizophrenia. *Biol Psychiatry*. 2016;79(7):595-603. doi:10.1016/j.biopsych.2015.06.015
- **43**. Heller AS, Johnstone T, Shackman AJ, et al. Reduced capacity to sustain positive emotion in major depression reflects diminished maintenance of fronto-striatal brain activation. *Proc Natl Acad Sci U S A*. 2009;106(52):22445-22450. doi:10.1073/pnas.0910651106
- **44**. Kasanova Z, Ceccarini J, Frank MJ, et al. Striatal dopaminergic modulation of reinforcement learning predicts reward-oriented behavior in daily life. *Biol Psychol.* 2017;127:1-9. doi:10.1016/j. biopsycho.2017.04.014
- **45**. Parsons LH, Hurd YL. Endocannabinoid signalling in reward and addiction. *Nat Rev Neurosci*. 2015;16(10):579-594. doi:10.1038/nrn4004
- **46**. Porath-Waller AJ, Beasley E, Beirness DJ. A meta-analytic review of school-based prevention for cannabis use. *Health Educ Behav*. 2010;37(5): 709-723. doi:10.1177/1090198110361315
- **47**. Chatters R, Cooper K, Day E, et al. Psychological and psychosocial interventions for cannabis cessation in adults: a systematic review. *Addict Res Theory*. 2016;24(2):93-110. doi:10.3109/16066359.2015.1073719
- **48**. Hjorthøj C, Fohlmann A, Nordentoft M. Treatment of cannabis use disorders in people with schizophrenia spectrum disorders—a systematic review. *Addict Behav.* 2009;34(6-7):520-525. doi:10.1016/j.addbeh.2009.02.001
- **49**. González-Ortega I, Echeburúa E, Alberich S, et al. Cognitive behavioral therapy program for cannabis use cessation in first-episode psychosis patients: a 1-year randomized controlled trial. *Int J Environ Res Public Health*. 2022;19(12):7325. doi:10.3390/ijerph19127325
- **50.** Arias AJ, Hammond CJ, Burleson JA, et al. Temporal dynamics of the relationship between change in depressive symptoms and cannabis use in adolescents receiving psychosocial treatment for cannabis use disorder. *J Subst Abuse Treat*. 2020; 117:108087. doi:10.1016/j.jsat.2020.108087
- **51.** Valmaggia LR, Day FL, Jones C, et al. Cannabis use and transition to psychosis in people at ultra-high risk. *Psychol Med*. 2014;44(12):2503-2512. doi:10.1017/S0033291714000117
- **52.** Van der Pol P, Liebregts N, de Graaf R, Korf DJ, van den Brink W, van Laar M. Predicting the transition from frequent cannabis use to cannabis dependence: a three-year prospective study. *Drug Alcohol Depend*. 2013;133(2):352-359. doi:10.1016/j.drugalcdep.2013.06.009
- **53.** Feingold D, Livne O, Rehm J, Lev-Ran S. Probability and correlates of transition from cannabis use to *DSM-5* cannabis use disorder: results from a large-scale nationally representative study. *Drug Alcohol Rev.* 2020;39(2):142-151. doi:10.1111/dar.13031
- **54**. Kessing L. Validity of diagnoses and other clinical register data in patients with affective disorder. *Eur Psychiatry*. 1998;13(8):392-398. doi:10.1016/S0924-9338(99)80685-3
- **55.** Bock C, Bukh JD, Vinberg M, Gether U, Kessing LV. Validity of the diagnosis of a single depressive episode in a case register. *Clin Pract Epidemiol Ment Health*. 2009;5(1):4. doi:10.1186/1745-0179-5-4

Drug Overdose Death Rate 1999-2021

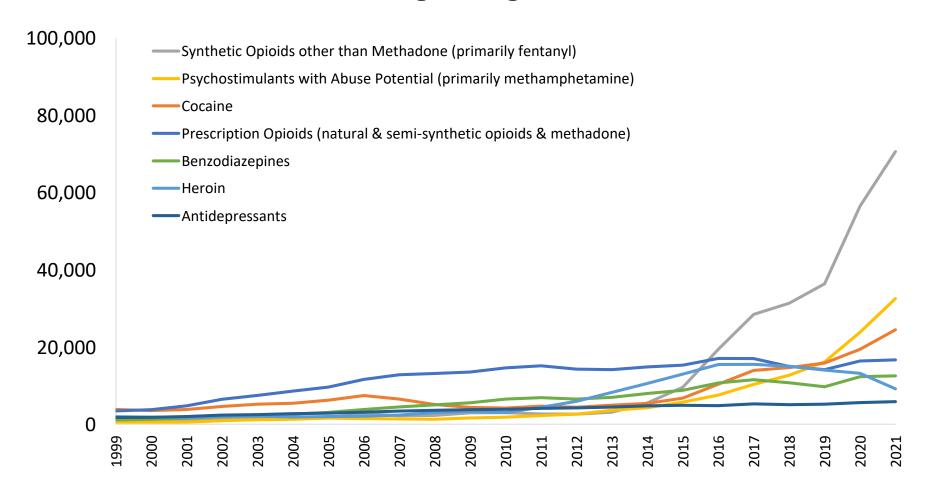
National Center for Health Statistics (NCHS)

Figure 1. National Drug-Involved Overdose Deaths*, Number Among All Ages, by Gender, 1999-2021



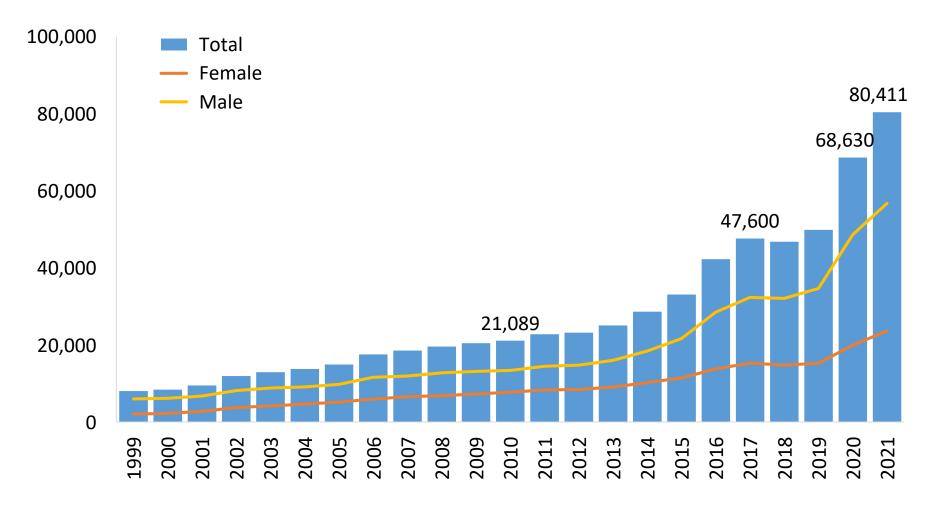
^{*}Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug poisoning (X60–X64), homicide drug poisoning (X85), or drug poisoning of undetermined intent (Y10–Y14), as coded in the International Classification of Diseases, 10th Revision. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 2. National Drug-Involved Overdose Deaths*, Number Among All Ages, 1999-2021



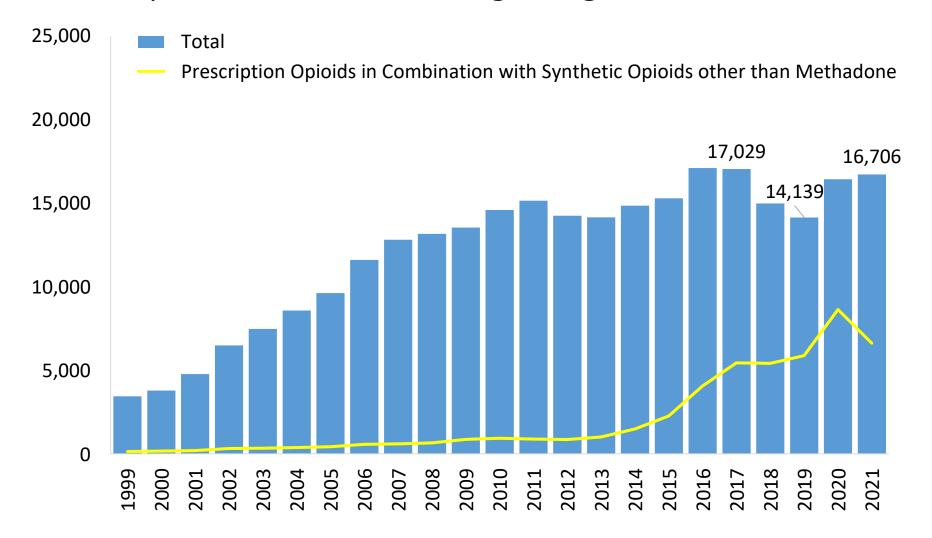
^{*}Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug poisoning (X60–X64), homicide drug poisoning (X85), or drug poisoning of undetermined intent (Y10–Y14), as coded in the International Classification of Diseases, 10th Revision. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 3. National Overdose Deaths Involving Any Opioid*, Number Among All Ages, by Gender, 1999-2021



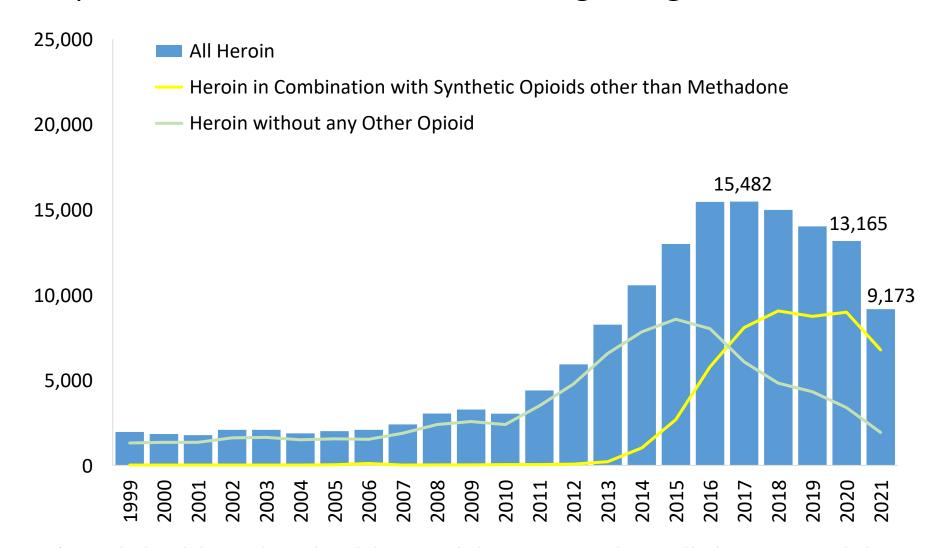
^{*}Among deaths with drug overdose as the underlying cause, the "any opioid" subcategory was determined by the following ICD-10 multiple cause-of-death codes: natural and semi-synthetic opioids (T40.2), methadone (T40.3), other synthetic opioids (other than methadone) (T40.4), or heroin (T40.1). Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 4. National Overdose Deaths Involving Prescription Opioids*, Number Among All Ages, 1999-2021



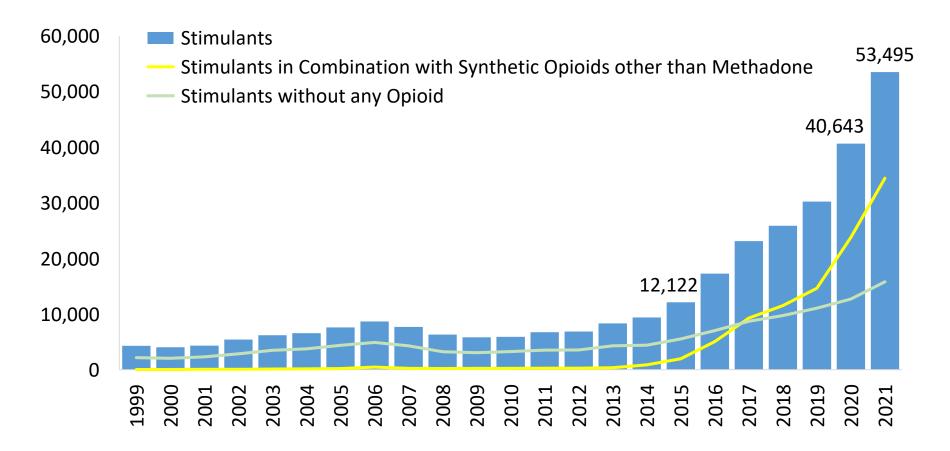
^{*}Among deaths with drug overdose as the underlying cause, the prescription opioid subcategory was determined by the following ICD-10 multiple cause-of-death codes: natural and semi-synthetic opioids (T40.2) or methadone (T40.3). Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 5. National Overdose Deaths Involving Heroin*, by other Opioid Involvement, Number Among All Ages, 1999-2021



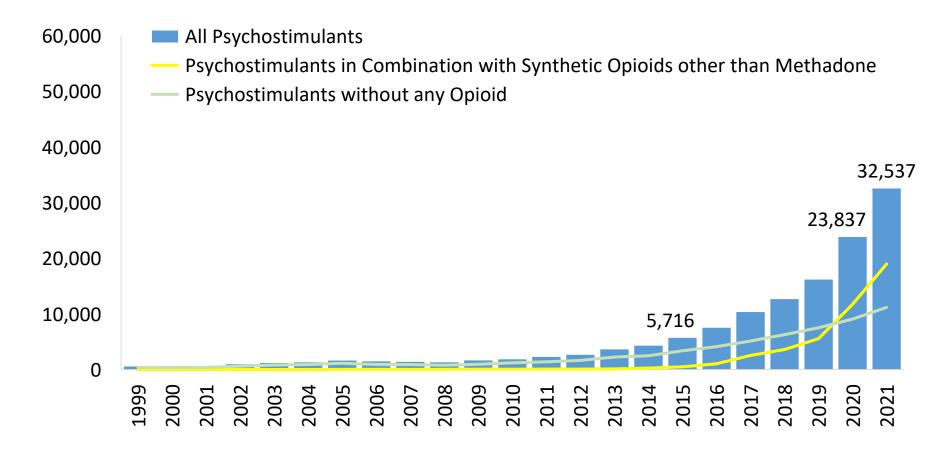
^{*}Among deaths with drug overdose as the underlying cause, the heroin category was determined by the T40.1 ICD-10 multiple cause-of-death code. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 6. National Overdose Deaths Involving Stimulants (Cocaine and Psychostimulants*), by Opioid Involvement, Number Among All Ages, 1999-2021



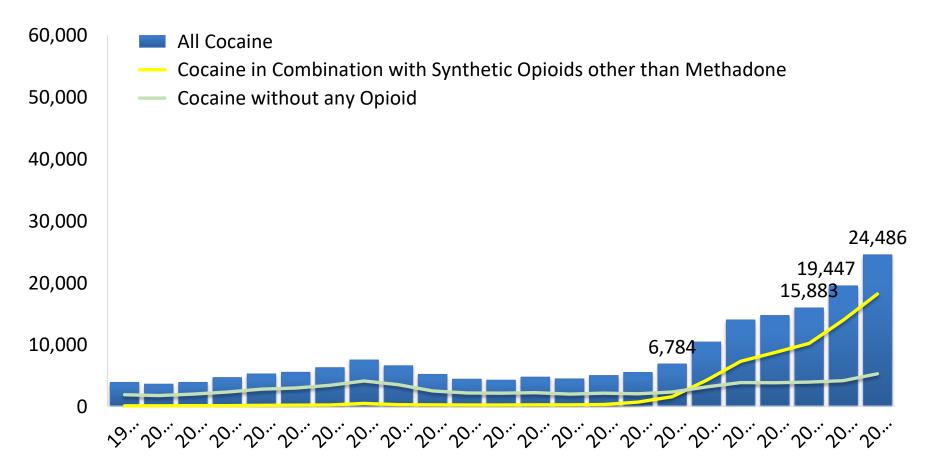
^{*}Among deaths with drug overdose as the underlying cause, the psychostimulants with abuse potential (primarily methamphetamine) category was determined by the T43.6 ICD-10 multiple cause-of-death code. Abbreviated to *psychostimulants* in the bar chart above. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 7. National Overdose Deaths Involving Psychostimulants with Abuse Potential (Primarily Methamphetamine)*, by Opioid Involvement, Number Among All Ages, 1999-2021



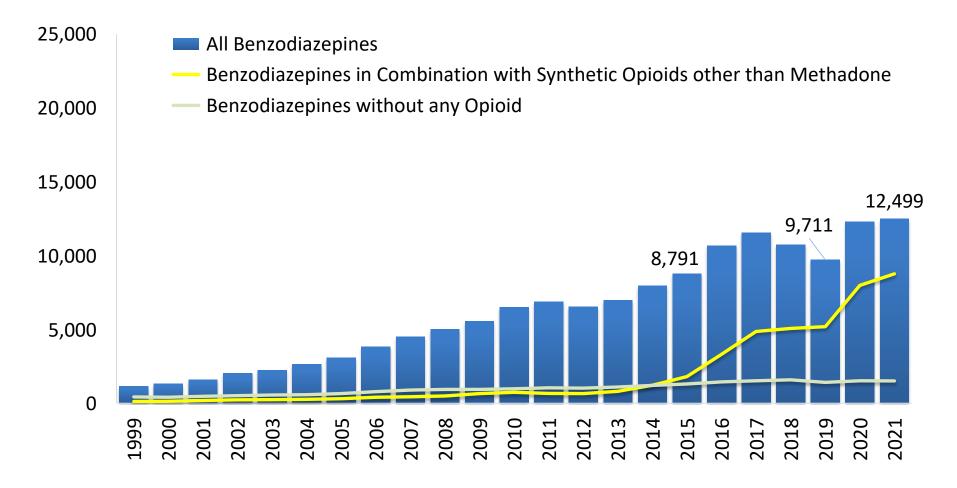
^{*}Among deaths with drug overdose as the underlying cause, the psychostimulants with abuse potential (primarily methamphetamine) category was determined by the T43.6 ICD-10 multiple cause-of-death code. Abbreviated to *psychostimulants* in the bar chart above. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 8. National Drug Overdose Deaths Involving Cocaine*, by Opioid Involvement, Number Among All Ages, 1999-2021



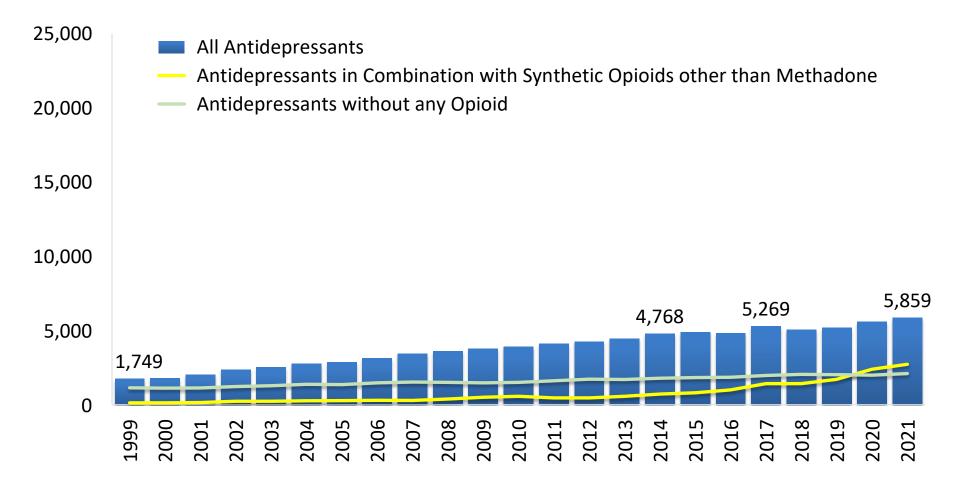
^{*}Among deaths with drug overdose as the underlying cause, the cocaine category was determined by the T40.5 ICD-10 multiple cause-of-death code. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 9. National Drug Overdose Deaths Involving Benzodiazepines*, by Opioid Involvement, Number Among All Ages, 1999-2021



^{*}Among deaths with drug overdose as the underlying cause, the benzodiazepine category was determined by the T42.4 ICD-10 multiple cause-of-death code. Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Figure 10. National Drug Overdose Deaths Involving Antidepressants*, by Opioid Involvement, Number Among All Ages, 1999-2021



^{*}Among deaths with drug overdose as the underlying cause, the antidepressant subcategory was determined by the following ICD-10 multiple cause-of-death codes: Tricyclic and tetracyclic antidepressants (T43.0), monoamine-oxidase-inhibitor antidepressants (T43.1), and other unspecified antidepressants (T43.2). Source: Centers for Disease Control and Prevention, National Center for Health Statistics. Multiple Cause of Death 1999-2021 on CDC WONDER Online Database, released 1/2023.

Descriptions of Figures

- The figures above are bar charts showing the number of U.S. overdose deaths involving select illicit or prescription drugs from 1999 through 2021. The bars are overlaid by lines representing gender or concurrent opioid involvement. Please note the y-axis scale varies by bar chart and caution should be applied when comparing graphs side-by-side.
- There were 106,699 drug-involved overdose deaths reported in the U.S. in 2021 (Figure 1); 69% of cases occurred among males (yellow line). Synthetic opioids other than methadone (primarily fentanyl) were the main driver of drug overdose deaths with a nearly 7.5-fold increase from 2015 to 2021 (Figure 2).
- Drug overdose deaths involving any opioid—prescription opioids (including natural and semi-synthetic opioids and methadone), other synthetic opioids other than methadone (primarily fentanyl), and heroin—continued to rise through 2021 with 80,411 deaths. More than 70% of deaths occurred among males (Figure 3). From 2020 to 2021, the number of deaths involving prescription opioids remained steady (Figure 4).
- Overdose deaths involving heroin have trended down since 2016 with 9,173 deaths reported in 2021 (Figure 5). Nearly 75% of overdose deaths in 2021 involving heroin also involved synthetic opioids other than methadone (primarily fentanyl).
- Drug overdose deaths involving stimulants, cocaine, or psychostimulants with abuse potential (primarily methamphetamine) have significantly increased since 2015 from 12,122 to 53,495 in 2021 (Figure 6).
- Since 2015, the number of deaths involving psychostimulants with abuse potential (primarily methamphetamine) has risen significantly each year—with 32,537 deaths in 2021 (Figure 7). The number of deaths involving cocaine has also increased steadily since 2015 with 24,486 deaths reported in 2021 (Figure 8).
- The final two charts show the number of overdose deaths involving benzodiazepines (Figure 8) or antidepressants (Figure 9). Benzodiazepines were involved in 12,499 deaths in 2021—steadily increasing since 2015. The proportion of deaths involving synthetic opioids other than methadone (primarily fentanyl) has increased significantly since 2015. Of the 8,791 deaths involving benzodiazepines in 2015, 20% also involved fentanyl. In 2021, this proportion increased to 70% of all deaths involving benzodiazepines. Antidepressant-involved deaths have also risen steadily, driven by fentanyl, with 5,859 deaths reported in 2021.

National Drug Overdos

Source: CDC WONDER, Multiple For information about this data go to

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	В		

All Ages

Number Drug OD Deaths
Rate Drug OD Deaths

Ages 15-24 Years

Number Drug OD, 15-24 Years

Rate Drug OD, 15-24 Years

Demographics

Rate OD Deaths, by Demographic

se (OD) Deaths, 1999-2021

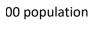
Cause of Death (Detailed Mortality)

https://wonder.cdc.gov/mcd.html

Number of National Drug Overdose Deaths Involving Select Prescription and Illicit Drugs
Rate of National Overdose Deaths Involving Select Prescription and Illicit Drugs, Rates are Age-Adjusted per 100,00

Number of National Drug Overdose Deaths Involving Select Prescription and Illicit Drugs, Ages 15-24 Years Old Rate of National Drug Overdose Deaths Involving Select Prescription and Illicit Drugs, Ages 15-24 Years Old, Rates are per 100,000 population

Rate of National Drug Overdose Deaths, by Demographic, Rates are Age-Adjusted per 100,000 population





Number of National Drug Overdose Deaths* Involving Select Prescription and Illicit Drugs Source: National Center on Health Statistics, CDC WONDER

Total Overdose Deaths	1999 16.849	2000 17.415	2001 19.394	2002	2003	2004	29,813	2006 34.425	2007 36,010	2008 36.450	2009 37,004	2010 38.329	2011 41.340	2012 41.502	2013	2014 47,055	2015 52.404	2016 63,632	2017 70,237	2018 67,367	2019 70,630	2020 91,799	2021 106,699
emale Male	5,591 11,258	17,415 5,852 11,563	6,736 12.658	8,490 15,028	9,386 16.399	10,304 17,120	29,813 11,089 18.724	12,532 21.893	13,712 22,298	13,982 22,468	14,411 22,593	15,323 23.006	41,340 16,352 24,988	16,390 25,112	43,982 17,183 26,799	18,243 28,812	19,447 32,957	22,074 41,558	23,685 46.552	67,367 22,426 44,941	70,630 22,749 47,881	28,071 63.728	32,398 74,301
e Opioid¹ ale	8,050	8,407	9,496	11,920	12,940	13,756	14,918	17,545	18,516	19,582	20,422	21,089	22,784	23,166	25,052	28,647	33,091	42,249 13,751	47,600	46,802 14,724	49,860 15,225	68,630 19,970	80,411 23,654
<u> </u>	2,057 5,993	2,264 6,143	2,767 6,729	3,760 8,160	4,138 8,802	4,643 9,113	5,161 9,757	5,945 11,600	6,581 11,935	6,819 12,763	7,287 13,135	7,734 13,355	8,325 14,459	8,432 14,734	9,055 15,997	10,227 18,420	11,420 21,671	28,498	15,263 32,337	32,078	34,635	48,660	56,757
ect Opioids ^{1a} AND Synthetic Opioids other than Methadone scription Opioids ²	167 3,442	205 3,785	225 4,770	358 6,483	383 7,461	428 8,577	485 9,612	781 11,589	642 12,796	703 13,149	923 13,523	1,002 14,583	961 15,140	944 14,240	1,219 14,145	2,493 14,838	4,806 15,281	9,299 17,087	12,556 17,029	13,491 14,975	13,596 14,139	16,465 16,416	15,444 16,706
nale le	1,022 2,420	1,236 2,549	1,608 3,162	2,304 4,179	2,681 4,780	3,144 5,433	3,572 6,040	4,274 7,315	4,863 7,933	4,959 8,190	5,212 8,311	5,644 8,939	6,082 9,058	5,995 8,245	6,049 8,096	6,506 8,332	6,664 8,617	7,109 9,978	7,156 9,873	6,252 8,723	5,755 8,384	6,441 9,975	6,623 10,083
escription Opioids AND Synthetic Opioids other than Methadone	142 65	167	199	322 157	344 151	384 184	426 207	573 246	601 286	655 309	872 444	939 453	889 426	861 445	1,015	1,489 661	2,263 898	4,055	5,444 1,859	5,417 1,872	5,876 1,949	8,626 2,798	9,644 6,393
ale escription Opioids WITHOUT Synthetic Opioids other than Methadone	77 3,300	91 3,618	113 4,571	165 6,161	193 7,117	200 8,193	219 9,186	327 11,016	315 12,195	346 12,494	428 12,651	486 13,644	463 14,251	416 13,379	527 13,130	828 13,349	1,365 13,018	2,661 13,032	3,585 11,585	3,545 9,558	3,927 8,263	5,828 7,790	3,251
emale Aale	957	1,160	1,522	2,147	2,530	2,960	3,365	4,028	4,577	4,650	4,768	5,191	5,656	5,550	5,561	5,845	5,766	5,715	5,297	4,380	3,806	3,643	230
are nthetic Opioids other than Methadone (primarily fentanyl) ³	2,343 730	2,458 782	3,049 957	4,014 1,295	4,587 1,400	5,233 1,664	5,821 1,742	6,988 2,707	7,618 2,213	7,844 2,306	7,883 2,946	8,453 3,007	8,595 2,666	7,829 2,628	7,569 3,105	7,504 5,544	7,252 9,580	7,317 19,413	28,466	5,178 31,335	4,457 36,359	4,147 56,51 6	6,832 70,601
emale Fale	330 400	374 408	447 510	614 681	643 757	798 866	823 919	1,030 1,677	1,053 1,160	1,083 1,223	1,445 1,501	1,440 1,567	1,247 1,419	1,195 1,433	1,431 1,674	2,079 3,465	3,020 6,560	5,578 13,835	7,942 20,524	8,807 22,528	10,076 26,283	15,250 41,266	19,571 51,030
roin ⁴ emale	1,960	1,842 279	1,779 313	2,089	2,080 358	1,878	2,009	2,088	2,399	3,041 551	3,278	3,036	4,397 878	5,925 1,213	8,257 1,732	10,574 2,414	12,989 3,108	15,469 3,717	15,482	14,996 3,705	14,019 3,520	13,165 3,284	9,173 2,372
1ale	1,654	1,563 18	1,466	1,730	1,722 16	1,537	1,620 34	1,744 113	2,000 13	2,490 28	2,701 29	2,452	3,519 44	4,712	6,525 209	8,160 1,027	9,881 2,685	11,752 5,781	11,596 8,091	11,291 9,068	10,499 8,746	9,881 8,990	6,801 6,783
eroin AND Synthetic Opioids other than Methadone emale	15	7	15 4	5	3	6	9	25	3	13	10	8	11	19	58	275	670	1,430	2,035	2,267	2,256	2,294	1,791
Male Proin WITHOUT Synthetic Opioids other than Methadone	11 1,945	11 1,824	11 1,764	2,074	2,064	7 1,865	25 1,975	88 1,975	2,386	3, 013	3,249	37 2,991	4, 353	50 5,856	151 8,048	752 9,547	2,015 10,304	4,351 9,688	6,056 7,391	6,801 5,928	6,490 5,273	6,696 4,175	4,992 2,390
emale 1ale	302 1,643	272 1,552	309 1,455	354 1,720	355 1,709	335 1,530	380 1,595	319 1,656	396 1,990	538 2,475	567 2,682	576 2,415	867 3,486	1,194 4,662	1,674 6,374	2,139 7,408	2,438 7,866	2,287 7,401	1,851 5,540	1,438 4,490	1,264 4,009	990 3,185	581 1,809
mulants ^{5a} emale	4,271 980	4,017 980	4,308 1,083	5,423 1,400	6,215 1,626	6,591 1,767	7,606 2,001	8,668 2,214	7,697 2,028	6,320 1,667	5,824 1,586	5,914 1,683	6,765 1,955	6,879 2,026	8,338 2,412	9,395 2,720	12,122 3,527	17,258 4,895	23,139 6,645	25,877 7,529	30,231 8,532	40,643 11,338	53,495 15,087
nulants AND Any Opioid	3,291 2,101	3,037 1,972	3,225 1,996	4,023 2,578	4,589 2,732	4,824 2,850	5,605	6,454 3,764	5,669 3,394	4,653 3,085	4,238 2,766	4,231 2,662	4,810 3,255	4,853 3,340	5,926 4,037	6,675 4,999	8,595 6 594	12,363 10,222	16,494 14,455	18,348 16,165	21,699 19,192	29,305 27,966	38,408 37,682
male	433	422	480	658	692	743	3,215 853	979	902	829	752	803	1,024	1,057	1,240	1,519	2,000	2,987	4,292	4,870	5,561	7,919	10,898
ale mulants WITHOUT Any Opioid	1,668 2,170	1,550 2,045	1,516 2,312	1,920 2,845	2,040 3,483	2,107 3,741	2,362 4,391	2,785 4,904	2,492 4,303	2,256 3,235	2,014 3,058	1,859 3,252	2,231 3,510	2,283 3,539	2,797 4,301	3,480 4,396	4,594 5,528	7,235 7,036	10,163 8,684	11,295 9,712	13,631 11,039	20,047 12,677	26,784 15,813
male ale	547 1,623	558 1,487	603 1,709	742 2,103	934 2,549	1,024 2,717	1,148 3,243	1,235 3,669	1,126 3,177	838 2,397	834 2,224	880 2,372	931 2,579	969 2,570	1,172 3,129	1,201 3,195	1,527 4,001	1,908 5,128	2,353 6,331	2,659 7,053	2,971 8,068	3,419 9,258	4,189 11,624
nulants AND Synthetic Opioids other than Methadone male	58 16	51	80	83	135 57	157 52	203 75	463 120	246 76	227 78	240 83	235	274 122	261 91	373 149	869 283	1,969 577	5,029 1,428	9,262 2,612	11,516 3,405	14,627 4,123	23,782 6,631	34,429 9,799
le nulants WITHOUT Synthetic Opioids other than Methadone	42 4,213	36 3,966	52 4,228	53 5,340	78 6,080	105 6,434	128 7,403	343 8,205	170 7,451	149 6,093	157 5,584	136 5,679	152 6,491	170 6,618	224 7,965	586 8,526	1,392 10,153	3,601 12,229	6,650 13,877	8,111 14,361	10,504 15,604	17,151 16,861	24,630 19,066
male	964	965	1,055	1,370	1,569	1,715	1,926	2,094	1,952	1,589	1,503	1,584	1,833	1,935	2,263	2,437	2,950	3,467	4,033	4,124	4,409	4,707	5,288
ile aine ⁵	3,249 3,822	3,001 3,544	3,173 3,833	3,970 4,599	4,511 5,199	4,719 5,443	5,477 6,208	6,111 7,448	5,499 6,512	4,504 5,129	4,081 4,350	4,095 4,183	4,658 4,681	4,683 4,404	5,702 4,944	6,089 5,415	7,203 6,784	8,762 10,375	9,844 13,942	10,237 14,666	11,195 15,883	12,154 19,447	13,778 24,486
nale Ile	850 2,972	843 2,701	957 2,876	1,143 3,456	1,322 3,877	1,405 4,038	1,620 4,588	1,860 5,588	1,665 4,847	1,322 3,807	1,141 3,209	1,132 3,051	1,314 3,367	1,262 3,142	1,376 3,568	1,535 3,880	1,899 4,885	2,882 7,493	3,921 10,021	4,228 10,438	4,336 11,547	5,245 14,202	6,858 17,628
aine AND Any Opioid	1,964 399	1,834 387	1,886 453	2,318 560	2,456 603	2,522 634	2,842 737	3,372 845	3,027	2,656 695	2,210 574	2,086 572	2,505 746	2,448 720	2,831	3,414 973	4,506 1,261	7,263 2,048	10,131 2,898	10,887 3,189	11,998 3,308	15,338 4,193	19,250 5,418
le	1,565	1,447	1,433	1,758	1,853	1,888	2,105	2,527	2,243	1,961	1,636	1,514	1,759	1,728	2,028	2,441	3,245	5,215	7,233	7,698	8,690	11,145	13,832
raine WITHOUT Any Opioid male	1,858 451	1,710 456	1,947 504	2,281 583	2,743 719	2,921 771	3,366 883	4,076 1,015	3,485 881	2,473 627	2,140 567	2,097 560	2,176 568	1,956 542	2,113 573	2,001 562	2,278 638	3,112 834	3,811 1,023	3,779 1,039	3,885 1,028	4,109 1,052	5,236 1,440
ale caine AND Synthetic Opioids other than Methadone	1,407 47	1,254 46	1,443 75	1,698 65	2,024 109	2,150 130	2,483 174	3,061 432	2,604 219	1,846 182	1,573 176	1,537 167	1,608 189	1,414 182	1,540 245	1,439 628	1,640 1,542	2,278 4,184	2,788 7,241	2,740 8,659	2,857 10,139	3,057 13,903	3,796 18,153
male ale	13 34	13 33	25 50	19 46	44 65	41 89	62 112	109 323	63 156	59 123	61 115	63 104	87 102	59 123	87 158	187 441	425 1,117	1,171 3,013	2,023 5,218	2,496 6,163	2,738 7,401	3,752 10,151	5,071 13,082
caine WITHOUT Synthetic Opioids other than Methadone	3,775 837	3,498 830	3,758 932	4,534 1,124	5,090 1,278	5,313 1,364	6,034 1,558	7,016 1,751	6,293 1,602	4,947 1,263	4,174 1,080	4,016 1,069	4,492 1,227	4,222 1,203	4,699 1,289	4,787 1,348	5,242 1,474	6,191 1,711	6,701 1,898	6,007 1,732	5,744 1,598	5,544 1,493	6,333 1,787
ale	2,938	2,668	2,826	3,410 941	3,812	3,949	4,476 1.608	5,265	4,691	3,684 1,302	3,094 1.632	2,947	3,265 2,266	3,019 2.635	3,410 3,627	3,439 4,298	3,768 5,716	4,480 7,542	4,803 10,333	4,275 12,676	4,146 16,167	4,051 23.837	4,546 32,537
ychostimulants With Abuse Potential (primarily methamphetamine) ⁶	547 158	578 164	563 152	285	1,179 353	1,305 393	438	1,462 411	1,378 409	375	489	1,854 592	693	816	1,106	1,278	1,745	2,194	3,093	3,775	4,733	6,890	9,218
lale ychostimulants With Abuse Potential AND Any Opioid	389 187	414 202	411 164	656 325	826 359	912 407	1,170 476	1,051 526	969 473	927 495	1,143 654	1,262 640	1,573 876	1,819 993	2,521 1,354	3,020 1,806	3,971 2,345	5,348 3,416	7,240 5,203	8,901 6,405	11,434 8,642	16,947 14,777	23,319 21,371
emale ale	46 141	49 153	42 122	114 211	119 240	128 279	145 331	170 356	150 323	151 344	207 447	255 385	316 560	378 615	491 863	610 1,196	819 1,526	1,072 2,344	1,683 3,520	2,058 4,347	2,696 5,946	4,402 10,375	6,317 15,054
ychostimulants With Abuse Potential WITHOUT Any Opioid	360 112	376	399 110	616 171	820 234	898 265	1,132 293	936 241	905 259	807 224	978 282	1,214 337	1,390 377	1,642 438	2,273 615	2,492 668	3,371 926	4,126 1,122	5,130 1,410	6,271 1,717	7,525 2,037	9,060 2,488	11,166 2,901
ale	248	261	289	445	586	633	839	695	646	583	696	877	1,013	1,204	1,658	1,824	2,445	3,004	3,720	4,554	5,488	6,572	8,265
chostimulants With Abuse Potential AND Synthetic Opioids other than thadone	11	7	6	19	28	29	33	37	35	47	69	73	93	91	142	276	494	1,042	2,546	3,613	5,564	11,717	18,986
emale ale	3 8	3 4	3	12 7	14 14	12 17	13 20	16 21	17 18	19 28	25 44	37 36	40 53	40 51	67 75	106 170	174 320	322 720	766 1,780	1,163 2,450	1,711 3,853	3,461 8,256	5,498 13,488
ychostimulants With Abuse Potential WITHOUT Synthetic Opioids other than ethadone	536	571	557	922	1,151	1,276	1,575	1,425	1,343	1,255	1,563	1,781	2,173	2,544	3,485	4,022	5,222	6,500	7,787	9,063	10,603	12,120	13,551
emale Nale	155 381	161 410	149 408	273 649	339 812	381 895	425 1.150	395 1.030	392 951	356 899	464 1.099	555 1.226	653 1,520	776 1.768	1,039 2.446	1,172 2.850	1,571 3.651	1,872 4.628	2,327 5.460	2,612 6.451	3,022 7.581	3,429 8.691	3,720 9.831
nzodiazepines ⁷	1,135	1,298	1,594	2,022	2,248	2,627	3,084	3,835	4,500	5,010	5,567	6,497	6,872	6,524	6,973	7,945	8,791	10,684	11,537	10,724	9,711	12,290	12,499
nale ile	420 715	480 818	614 980	763 1,259	885 1,363	1,079 1,548	1,209 1,875	1,472 2,363	1,894 2,606	2,046 2,964	2,281 3,286	2,579 3,918	2,902 3,970	2,789 3,735	3,026 3,947	3,487 4,458	3,779 5,012	4,359 6,325	4,772 6,765	4,481 6,243	3,986 5,725	4,810 7,480	5,056 7,443
nzodiazepines AND Any Opioid emale	701 233	892 310	1,121 411	1,511 553	1,692 620	2,049 797	2,430 922	3,045 1,137	3,605 1,473	4,070 1,618	4,633 1,853	5,517 2,125	5,826 2,408	5,500 2,283	5,869 2,485	6,733 2,876	7,485 3,137	9,233 3,723	10,010 4,031	9,140 3,699	8,301 3,303	10,771 4,108	10,992 4,307
ale nzodiazepines WITHOUT Any Opioid	468 434	582 406	710 473	958 511	1,072 556	1,252 578	1,508 654	1,908 790	2,132 895	2,452 940	2,780 934	3,392 980	3,418 1,046	3,217 1,024	3,384 1,104	3,857 1,212	4,348 1,306	5,510 1,451	5,979 1,527	5,441 1,584	4,998 1,410	6,663 1,519	6,685 1,507
male ale	187 247	170 236	203 270	210 301	265 291	282 296	287 367	335 455	421 474	428 512	428 506	454 526	494 552	506 518	541 563	611 601	642 664	636 815	741 786	782 802	683 727	702 817	749 758
nzodiazepines AND Synthetic Opioids other than Methadone	122	136	186	230	242	270	312	407	436	491	658	746	665	655	804	1,222	1,801	3,308	4,869	5,066	5,187	7,983	8,759
male ale	51 71	55 81	83 103	107 123	105 137	128 142	150 162	165 242	224 212	250 241	330 328	355 391	324 341	304 351	388 416	534 688	738 1,063	1,188 2,120	1,696 3,173	1,808 3,258	1,834 3,353	2,733 5,250	3,180 5,579
zodiazepines WITHOUT Synthetic Opioids other than Methadone male	1,013 369	1,162 425	1,408 531	1,792 656	2,006 780	2,357 951	2,772 1,059	3,428 1,307	4,064 1,670	4,519 1,796	4,909 1,951	5,751 2,224	6,207 2,578	5,869 2,485	6,169 2,638	6,723 2,953	6,990 3,041	7,376 3,171	6,668 3,076	5,658 2,673	4,524 2,152	4,307 2,077	3,740 1,876
le depressants ⁸	644 1,749	737 1,798	877 2.017	1,136 2.370	1,226 2,512	1,406 2,758	1,713 2,861	2,121 3,133	2,394 3,425	2,723 3.610	2,958 3,768	3,527 3.889	3,629 4,113	3,384 4.259	3,531 4,458	3,770 4,768	3,949 4.894	4,205 4.812	3,592 5,269	2,985 5,064	2,372 5,175	2,230 5,597	1,864 5,859
ale	926	984	1,009	1,318	1,384	1,549	1,575	1,819	1,958	2,047	2,133	2,204	2,422	2,469	2,673	2,857	2,909	2,835	3,112	2,993	3,048	3,122	3,343
depressants AND Any Opioid	823 611	814 679	1,008 890	1,052 1,148	1,128 1,234	1,209 1,379	1,286 1,508	1,314 1,662	1,467 1,901	1,563 2,111	1,635 2,292	1,685 2,389	1,691 2,501	1,790 2,536	1,785 2,763	1,911 2,983	1,985 3,062	1,977 2,960	2,157 3,301	2,071 3,019	2,127 3,151	2,475 3,611	2,516 3,767
nale le	312 299	387 292	426 464	634 514	692 542	762 617	806 702	946 716	1,094 807	1,199 912	1,308 984	1,364 1,025	1,468 1,033	1,470 1,066	1,664 1,099	1,767 1,216	1,758 1,304	1,719 1,241	1,871 1,430	1,715 1,304	1,773 1,378	1,931 1,680	2,022 1,745
idepressants WITHOUT Any Opioid male	1,138 614	1,119 597	1,127 583	1,222 684	1,278 692	1,379 787	1,353	1,471 873	1,524 864	1,499 848	1,476 825	1,500	1,612 954	1,723 999	1,695 1,009	1,785 1,090	1,832 1,151	1,852 1,116	1,968 1,241	2,045 1,278	2,024 1,275	1,986 1,191	2,092 1,321
ale	524	522	544	538	586	592	584	598	660	651	651	660	658	724	686	695	681	736	727	767	749	795	771
tidepressants AND Synthetic Opioids other than Methadone emale	122 61	123 77	147 79	238 154	230 139	264 164	278 159	300 176	292 203	384 248	505 337	568 358	463 283	464 273	571 348	723 437	808 463	1,002 529	1,414 692	1,423 704	1,710 887	2,387 1,149	2,721 1,347
ale tidepressants WITHOUT Synthetic Opioids other than Methadone	1,627	46 1,675	68 1,870	2,132	91 2,282	100 2,494	2,583	2,833	89 3,133	136 3,226	168 3,263	210 3,321	3,650	191 3,795	223 3,887	286 4,045	345 4,086	473 3,810	722 3,855	719 3,641	823 3,465	1,238 3,210	1,374 3,138
emale	865	907	930	1,164	1,245	1,385	1,416	1,643	1,755	1,799	1,796	1,846	2,139	2,196	2,325	2,420	2,446	2,306	2,420	2,289	2,161	1,973	1,996

*Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug poisoning (X60–X64), homicide drug poisoning of undetermined intent (Y10–Y14), as coded in the International Classification of Diseases, 10th

^See https://www.cdc.gov/nchs/nvss/vsrr/drug-overdose-data.htm for technical information.

¹ Any Opioid ICD-10 codes (T40.0-T40.4, T40.6)

^{1a} Opioids include ICD-10 codes (T40.0-T40.3 and T40.6)

² Prescription Opioids ICD-10 codes (T40.2-T40.3)

³Synthetic Opioids other than Methadone (primarily fentanyl) ICD-10 code (T40.4) This category is dominated by fentanyl related overdoses.

⁴Heroin ICD-10 codes (T40.1)
^{5a}Stimulants ICD-10 codes (T40.5 & T43.6)

⁵Cocaine ICD-10 codes (T40.5)

⁶Psychostimulants With Abuse Potential ICD-10 code (T43.6) This category is dominated by methamphetamine related overdoses.

⁷Benzodiazepines ICD-10 code(T42.4)

'Benzodiazepines ICD-10 code(T42.4)

8Antidepressants ICD-10 code(T43.0-T43.2)



Rate of National Drug Overdose Deaths* Involving Select Pres

Rates are Age-Adjusted per 100,000 population

Source: National Center on Health Statistics, CDC WONDER

	1999	2000
Total Overdose Deaths	6.1	6.2
Female	3.9	4.1
Male	8.2	8.3
Any Opioid ¹	2.9	3.0
Female	1.4	1.6
Male	4.3	4.4
Prescription Opioids ²	1.2	1.3
Female	0.7	0.9
Male	1.7	1.8
Prescription Opioids AND Synthetic Opioids other than Methadone	0.0	0.0
Female Male	0.1	0.1
Prescription Opioids WITHOUT Synthetic Opioids other than Methadone	0.1 1.2	0.1 1.3
Female	1.2	1.5
Male	1.6	1.7
Synthetic Opioids other than Methadone (primarily fentanyl) ³	0.3	0.3
Female	0.2	0.3
Male	0.2	0.3
Heroin⁴	0.7	0.7
Female	0.2	0.2
Male	1.2	1.1
Heroin AND Synthetic Opioids other than Methadone		
Female		
Male		
Heroin WITHOUT Synthetic Opioids other than Methadone		
Female		
Male		
Stimulants ^{5a}	1.5	1.4
Female	0.7	0.7
Male	2.3	2.2
Stimulants AND Any Opioid	0.7	0.7
Female	0.3	0.3
Male	1.2	1.1
Stimulants WITHOUT Any Opioid	0.8	0.7
Female Male	0.4	0.4
Stimulants AND Synthetic Opioids other than Methadone	1.1 0.0	1.1 0.0
	0.0	0.0
Female		_
Male	0.0	0.0
Stimulants WITHOUT Synthetic Opioids other than Methadone	1.5	1.4
Female Male	າ າ	2.2
iviale	2.3	2.2

Cocaine ⁵	1.4	1.3
Female	0.6	0.6
Male	2.1	1.9
Cocaine AND Any Opioid	0.7	0.6
Female	0.3	0.3
Male	1.1	1.0
Cocaine WITHOUT Any Opioid	0.7	0.7
Female	0.3	0.3
Male	1.0	0.9
Cocaine AND Synthetic Opioids other than Methadone		
Female		
Male		
Cocaine WITHOUT Synthetic Opioids other than Methadone		
Female Male		
	0.3	0.3
Psychostimulants With Abuse Potential (primarily methamphetamine) ⁶	0.2	0.2
Female	0.1	0.1
Male	0.3	0.3
Psychostimulants With Abuse Potential AND Any Opioid	0.1	0.1
Female	0.1	0.1
Male Resolventian Mith Abuse Petertial WITHOUT Any Original	0.1	0.1
Psychostimulants With Abuse Potential WITHOUT Any Opioid Female	0.1 0.1	0.1 0.1
Male	0.1	0.1
Psychostimulants With Abuse Potential AND Synthetic Opioids other than	0.2	0.2
Male Psychostimulants With Abuse Potential WITHOUT Synthetic Opioids other than Methadone Female Male		
Benzodiazepines ⁷	0.4	0.5
Female	0.3	0.3
Male	0.5	0.6
Benzodiazepines AND Any Opioid	0.2	0.3
Female	0.2	0.2
Male	0.3	0.4
Benzodiazepines WITHOUT Any Opioid	0.2	0.2
Female	0.1	0.1
Male	0.2	0.2
Benzodiazepines AND Synthetic Opioids other than Methadone		
Female		
Male		
Benzodiazepines WITHOUT Synthetic Opioids other than Methadone		
Female		
Male		
Antidepressants ⁸	0.6	0.6
Female	0.6	0.7
Male	0.6	0.6
Antidepressants AND Any Opioid	0.2	0.3
Female	0.2	0.3
Male	0.2	0.2
Antidepressants WITHOUT Any Opioid	0.4	0.3
Female	0.4	0.4

Male	0.4	0.4
Antidepressants AND Synthetic Opioids other than Methadone		
Female		0.1
Male		0.0
Antidepressants WITHOUT Synthetic Opioids other than Methadone		
Female		0.6
Male		0.6

^{*}Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug poison Blank fields designated by unreliable or suppressed data. For more information visit CDC WONDER.

¹ Any Opioid ICD-10 codes (T40.0-T40.4, T40.6)

² Prescription Opioids ICD-10 codes (T40.2-T40.3)

³Other Synthetic Narcotics (other than methadone) ICD-10 code (T40.4) This category is dominated by 1

⁴Heroin ICD-10 codes (T40.1)

^{5a}Stimulants ICD-10 codes (T40.5 & T43.6)

⁵Cocaine ICD-10 codes (T40.5)

⁶Psychostimulants With Abuse Potential ICD-10 code (T43.6) This category is dominated by methamphe

⁷Benzodiazepines ICD-10 code(T42.4)

⁸Antidepressants ICD-10 code(T43.0-T43.2)

cription and Illicit Drugs

2001	2002	2003	2004	2005	2006	2007
6.8	8.2	8.9	9.4	10.1	11.5	11.9
4.6	5.8	6.4	6.9	7.3	8.2	8.8
9.0	10.6	11.5	11.8	12.8	14.8	14.9
3.3	4.1	4.5	4.7	5.1	5.9	6.1
1.9	2.6	2.8	3.1	3.4	3.9	4.3
4.8	5.7	6.1	6.3	6.6	7.8	8.0
1.7	2.3	2.6	2.9	3.2	3.9	4.2
1.1	1.6	1.8	2.1	2.4	2.8	3.2
2.2	2.9	3.3	3.7	4.1	4.9	5.3
0.1	0.1	0.1	0.1	0.2	0.2	0.2
	0.1	0.1	0.1	0.1	0.2	0.2
0.1	0.1	0.1	0.1	0.1	0.2	0.2
1.6	2.2	2.5	2.8	3.0	3.7	4.0
	1.5	1.7	2.0	2.3	2.6	3.0
2.1	2.8	3.2	3.6	4.0	4.7	5.1
0.3	0.4	0.5	0.6	0.6	0.9	0.7
0.3	0.4	0.4	0.5	0.5	0.7	0.7
0.4	0.5	0.5	0.6	0.6	1.1	0.8
0.6	0.7	0.7	0.6	0.7	0.7	0.8
0.2	0.2	0.2	0.2	0.3	0.2	0.3
1.0	1.2	1.2	1.1	1.1	1.2	1.3

1.5	1.9	2.1	2.3	2.6	2.9	2.6
0.8	1.0	1.1	1.2	1.4	1.5	1.4
2.3	2.8	3.2	3.3	3.8	4.4	3.8
0.7	0.9	0.9	1.0	1.1	1.3	1.2
0.3	0.4	0.5	0.5	0.6	0.6	0.6
1.1	1.4	1.4	1.4	1.6	1.9	1.7
0.8	1.0	1.2	1.3	1.5	1.6	1.4
0.5	0.6	0.6	0.7	0.8	0.9	0.8
1.2	1.4	1.8	1.9	2.2	2.5	2.1
0.0	0.0	0.0	0.0	0.1	0.2	0.1
0.0	0.0	0.0	0.0	0.0	0.1	0.0
0.0	0.0	0.1	0.1	0.1	0.2	0.1
1.5	1.9	2.1	2.3	2.5	2.7	2.5
1.5	1.9	2.1	2.3	2.6	1.4	1.4
2.3	2.8	3.1	3.2	3.7	4.2	3.7

1.3	1.6	1.8	1.9	2.1	2.5	2.2
0.7	0.8	0.9	1.0	1.1	1.3	1.1
2.0	2.4	2.7	2.8	3.1	3.8	3.2
0.6	0.8	0.8	0.9	1.0	1.1	1.0
0.3	0.4	0.4	0.4	0.5	0.6	0.5
1.0	1.2	1.3	1.3	1.4	1.7	1.5
0.7	0.8	1.0	1.0	1.1	1.4	1.2
0.4	0.4	0.5	0.6	0.6	0.7	0.6
1.0	1.2	1.4	1.5	1.7	2.1	1.7
					0.2	0.1
					0.1	
		0.1	0.1	0.1	0.2	0.1
					2.3	2.1
					1.2	1.1
		2.6	2.7	3.0	3.6	3.1
0.2	0.3	0.4	0.4	0.5	0.5	0.4
0.1	0.2	0.2	0.3	0.3	0.3	0.3
0.3	0.5	0.6	0.6	0.8	0.7	0.7
0.1	0.1	0.1	0.1	0.2	0.2	0.2
	0.1	0.1	0.1	0.1	0.1	0.1
0.1	0.1	0.2	0.2	0.2	0.2	0.2
0.1	0.2	0.3	0.3	0.3	0.3	0.2
0.1	0.1	0.1	0.2	0.2	0.2	0.2
0.2	0.4	0.4	0.4	0.6	0.5	0.5

0.6	0.7	0.8	0.9	1.1	1.3	1.5
0.4	0.5	0.6	0.7	0.8	1.0	1.2
0.7	0.9	0.9	1.1	1.3	1.6	1.7
0.4	0.5	0.6	0.7	0.8	1.0	1.2
0.3	0.4	0.4	0.5	0.6	0.7	1.0
0.5	0.7	0.7	0.9	1.0	1.3	1.4
0.2	0.2	0.2	0.2	0.3	0.3	0.3
0.1	0.1	0.2	0.2	0.2	0.3	0.2
0.2	0.2	0.2	0.2	0.3	0.3	0.3
0.1	0.1	0.1	0.1	0.1	0.1	0.1
	0.1	0.1	0.1	0.1	0.1	0.2
0.1	0.1	0.1	0.1	0.1	0.2	0.1
0.5	0.6	0.7	0.8	1.0	1.2	1.4
	0.4	0.5	0.6	0.7	0.9	1.0
0.6	0.8	0.8	1.0	1.2	1.4	1.6
0.7	0.8	0.9	0.9	0.9	1.0	1.1
0.7	0.9	0.9	1.0	1.0	1.2	1.2
0.7	0.7	0.8	0.8	0.9	0.9	1.0
0.3	0.4	0.4	0.5	0.5	0.5	0.6
0.3	0.4	0.5	0.5	0.5	0.6	0.7
0.3	0.4	0.4	0.4	0.5	0.5	0.5
0.4	0.4	0.5	0.4	0.4	0.5	0.5
0.4	0.5	0.4	0.5	0.5	0.6	0.5

0.4	0.3	0.4	0.4	0.4	0.4	0.5
0.1	0.1	0.1	0.1	0.1	0.1	0.1
0.1	0.1	0.1	0.1	0.1	0.1	0.1
0.0	0.1	0.1	0.1	0.1	0.1	0.1
0.6	0.7	0.8	0.8	0.8	0.9	1.0
0.6	0.8	0.8	0.9	0.9	1.1	1.1
0.7	0.6	0.7	0.7	0.8	0.8	0.9

ing (X60–X64), homicide drug poisoning (X85), or drug poisoning of undetermined intent (Y10–Y14), as coded in

fentanyl related overdoses.

etamine related overdoses.

2008	2009	2010	2011	2012	2013	2014
11.9	11.9	12.3	13.2	13.1	13.8	14.7
8.9	9.1	9.6	10.2	10.2	10.6	11.1
14.9	14.8	15.0	16.1	16.1	17.0	18.3
6.4	6.6	6.8	7.3	7.4	7.9	9.0
4.4	4.6	4.9	5.2	5.3	5.6	6.3
8.4	8.7	8.7	9.4	9.5	10.2	11.7
4.3	4.4	4.7	4.9	4.5	4.4	4.6
3.2	3.3	3.6	3.8	3.7	3.7	3.9
5.4	5.5	5.8	5.9	5.3	5.1	5.2
0.2	0.3	0.3	0.3	0.3	0.3	0.5
0.2	0.3	0.3	0.3	0.3	0.3	0.4
0.2	0.3	0.3	0.3	0.3	0.3	0.5
4.1	4.1	4.4	4.6	4.2	4.1	4.1
3.0	3.0	3.3	3.5	3.4	3.4	3.5
5.2	5.2	5.5	5.6	5.0	4.8	4.7
0.8	1.0	1.0	0.8	0.8	1.0	1.8
0.7	0.9	0.9	0.8	0.7	0.9	1.3
0.8	1.0	1.0	0.9	0.9	1.1	2.2
1.0	1.1	1.0	1.4	1.9	2.7	3.4
0.4	0.4	0.4	0.6	0.8	1.2	1.6
1.6	1.8	1.6	2.3	3.1	4.2	5.2
					0.1	0.3
						0.2
					0.1	0.5
					2.6	3.1
					1.2	1.4
					4.1	4.7
2.1	1.9	1.9	2.2	2.2	2.6	3.0
1.1	1.0	1.1	1.3	1.3	1.5	1.7
3.1	2.8	2.8	3.1	3.1	3.7	4.2
1.0	0.9	0.9	1.1	1.1	1.3	1.6
0.5	0.5	0.5	0.7	0.7	0.8	1.0
1.5	1.3	1.2	1.5	1.5	1.8	2.3
1.1	1.0	1.0	1.1	1.1	1.3	1.4
0.6 1.6	0.5 1.5	0.6	0.6	0.6 1.6	0.7	0.7
0.1	0.1	1.6 0.1	1.6 0.1	0.1	1.9 0.1	1.9
						0.3
0.0	0.1	0.1	0.1	0.1	0.1	0.2
0.1	0.1	0.1	0.1	0.1	0.2	0.4
2.0	1.8	1.8	2.1	2.1	2.5	2.7
1.1	0.9	1.0	1.2	1.2	1.4	1.5
3.0	2.7	2.7	3.0	3.0	3.5	3.8

1.7						
	1.4	1.3	1.5	1.4	1.6	1.7
0.9	0.7	0.7	0.8	0.8	0.9	1.0
2.5	2.1	2.0	2.2	2.0	2.3	2.4
0.9	0.7	0.7	0.8	0.8	0.9	1.1
0.4	0.4	0.4	0.5	0.5	0.5	0.6
1.3	1.1	1.0	1.2	1.1	1.3	1.6
0.8	0.7	0.6	0.7	0.6	0.7	0.6
0.5	0.3	0.3	0.3	0.3	0.4	0.4
1.2	1.0	1.0	1.0	0.9	1.0	0.8
0.0	0.0	0.1	0.1	0.1	0.1	0.2
					0.1	0.1
0.1	0.1	0.1	0.1	0.1	0.1	0.3
1.7	1.4	1.2	1.4	1.3	1.5	1.5
0.9	0.7	0.7	0.8	0.8	0.8	0.9
2.4	2.0	1.9	2.1	1.9	2.2	2.1
0.4	0.5	0.6	0.7	0.8	1.2	1.4
0.2	0.3	0.4	0.4	0.5	0.7	0.8
0.6	0.8	0.8	1.0	1.2	1.6	1.9
0.2	0.2	0.2	0.3	0.3	0.4	0.6
0.1	0.1	0.2	0.2	0.2	0.3	0.4
0.2	0.3	0.2	0.4	0.4	0.6	0.8
0.2	0.3	0.4	0.4	0.5	0.8	0.8
0.1	0.2	0.2	0.2	0.3	0.4	0.4
0.4	0.5	0.6	0.6	0.8	1.0	1.1
					0.1	0.1
					0.0	0.1
					0.1	0.1 0.1
					0.1	0.1
					1.1	1.3
					0.7	0.7
					1.5	1.8
	1.8	2.1				1.0
16			22	2 1		2.5
1.6			1.8	2.1	2.2	2.5
1.3	1.5	1.6	1.8	1.7	2.2 1.9	2.1
1.3 2.0	1.5 2.2	1.6 2.6	1.8 2.6	1.7 2.4	2.2 1.9 2.5	2.1 2.8
1.3 2.0 1.3	1.5 2.2 1.5	1.6 2.6 1.8	1.8 2.6 1.9	1.7 2.4 1.8	2.2 1.9 2.5 1.8	2.1 2.8 2.1
1.3 2.0 1.3 1.0	1.5 2.2 1.5 1.2	1.6 2.6 1.8 1.3	1.8 2.6 1.9 1.5	1.7 2.4 1.8 1.4	2.2 1.9 2.5 1.8 1.5	2.1 2.8 2.1 1.8
1.3 2.0 1.3 1.0 1.6	1.5 2.2 1.5 1.2 1.8	1.6 2.6 1.8 1.3 2.2	1.8 2.6 1.9 1.5 2.2	1.7 2.4 1.8 1.4 2.1	2.2 1.9 2.5 1.8 1.5 2.2	2.1 2.8 2.1 1.8 2.5
1.3 2.0 1.3 1.0 1.6 0.3	1.5 2.2 1.5 1.2 1.8 0.3	1.6 2.6 1.8 1.3 2.2	1.8 2.6 1.9 1.5 2.2 0.3	1.7 2.4 1.8 1.4 2.1	2.2 1.9 2.5 1.8 1.5 2.2	2.1 2.8 2.1 1.8 2.5 0.4
1.3 2.0 1.3 1.0 1.6 0.3 0.3	1.5 2.2 1.5 1.2 1.8 0.3 0.3	1.6 2.6 1.8 1.3 2.2 0.3 0.3	1.8 2.6 1.9 1.5 2.2 0.3 0.3	1.7 2.4 1.8 1.4 2.1 0.3 0.3	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4	2.1 2.8 2.1 1.8 2.5 0.4 0.3
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4	1.5 2.2 1.5 1.2 1.8 0.3 0.3	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4	1.7 2.4 1.8 1.4 2.1 0.3 0.3	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4 0.3 0.4
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 0.2	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 0.2 1.9	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4 0.3 0.4 2.1
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 1.9 1.5	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4 2.1 1.8
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0	1.6 2.6 1.8 1.3 2.2 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 1.9 1.5 2.2	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.4 0.3 0.4 2.1 1.8 2.4
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 1.9 1.5 2.2 1.3	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4 2.1 1.8 2.4
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2 1.3	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2 1.4	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 1.9 1.5 2.2 1.3 1.5	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4 1.6	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.3 0.4 2.1 1.8 2.4 1.5
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1 1.8 1.1	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2 1.3 1.1	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2 1.4 1.1	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3 1.5 1.1	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 0.2 1.9 1.5 2.2 1.3 1.5 1.1	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4 1.6 1.1	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.4 0.3 0.4 2.1 1.8 2.4 1.5 1.7
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1 1.3 1.0 0.7	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2 1.3 1.1	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2 1.4 1.1 0.8	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3 1.5 1.1	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 0.2 1.9 1.5 2.2 1.3 1.5 1.1	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4 1.6 1.1 0.8	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.4 0.3 0.4 2.1 1.8 2.4 1.5 1.7 1.2 0.9
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1 1.3 1.0 0.7 0.8	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2 1.3 1.1 0.7 0.8	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2 1.4 1.1 0.8 0.8	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3 1.5 1.1 0.8 0.9	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 0.2 1.9 1.5 2.2 1.3 1.5 1.1 0.8 0.9	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4 1.6 1.1 0.8 1.0	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.4 0.3 0.4 2.1 1.8 2.4 1.5 1.7 1.2 0.9 1.1
1.3 2.0 1.3 1.0 1.6 0.3 0.3 0.4 0.2 0.2 0.2 1.4 1.1 1.8 1.1 1.3 1.0 0.7	1.5 2.2 1.5 1.2 1.8 0.3 0.3 0.4 0.2 0.2 0.2 1.6 1.3 2.0 1.2 1.3 1.1	1.6 2.6 1.8 1.3 2.2 0.3 0.3 0.4 0.2 0.2 0.3 1.9 1.4 2.3 1.2 1.4 1.1 0.8	1.8 2.6 1.9 1.5 2.2 0.3 0.3 0.4 0.2 0.2 2.0 1.6 2.4 1.3 1.5 1.1	1.7 2.4 1.8 1.4 2.1 0.3 0.3 0.3 0.2 0.2 0.2 1.9 1.5 2.2 1.3 1.5 1.1	2.2 1.9 2.5 1.8 1.5 2.2 0.4 0.4 0.3 0.2 0.2 0.3 2.0 1.7 2.2 1.4 1.6 1.1 0.8	2.1 2.8 2.1 1.8 2.5 0.4 0.3 0.4 0.3 0.4 2.1 1.8 2.4 1.5 1.7 1.2 0.9

0.4	0.5	0.5	0.4	0.4	0.4	0.4
0.1	0.2	0.2	0.1	0.1	0.2	0.2
0.2	0.2	0.2	0.2	0.2	0.2	0.3
0.1	0.1	0.1	0.1	0.1	0.2	0.2
1.0	1.0	1.0	1.2	1.2	1.2	1.3
1.1	1.1	1.2	1.3	1.3	1.4	1.4
0.9	1.0	1.0	1.0	1.0	0.9	1.0

the International Classification of Diseases, 10th Revision.

2015	2016	2017	2018	2019	2020	2021
16.3	19.8	21.7	20.7	21.6	28.3	32.4
11.8	13.4	14.4	13.6	13.7	17.1	19.6
20.8	26.2	29.1	27.9	29.6	39.5	45.1
10.4	13.3	14.9	14.6	15.5	21.4	24.7
7.1	8.5	9.4	9.0	9.3	12.3	14.5
13.7	18.1	20.4	20.1	21.7	30.4	34.8
4.7	5.2	5.2	4.5	4.2	4.9	4.9
4.0	4.3	4.2	3.7	3.4	3.8	3.8
5.4	6.2	6.1	5.3	5.1	6.1	6.0
0.7	1.3	1.7	1.7	1.8	2.7	2.9
0.6	0.9	1.2	1.2	1.2	1.7	2.0
0.8	1.7	2.3	2.2	2.4	3.6	3.8
4.0	3.9	3.5	2.8	2.4	2.2	2.0
3.4	3.4	3.0	2.5	2.2	2.1	1.8
4.6	4.5	3.8	3.1	2.7	2.5	2.2
3.1	6.2	9.0	9.9	11.4	17.8	21.8
1.9	3.5	5.0	5.5	6.3	9.6	12.2
4.2	8.9	13.0	14.2	16.6	25.9	31.4
4.1	4.9	4.9	4.7	4.4	4.1	2.8
2.0	2.4	2.5	2.3	2.2	2.0	1.5
6.3	7.5	7.3	7.1	6.6	6.1	4.1
0.9	1.8	2.6	2.9	2.7	2.8	2.1
0.4	0.9	1.3	1.4	1.4	1.5	1.1
1.3	2.8	3.8	4.3	4.1	4.1	3.0
3.2	3.1	2.3	1.8	1.7	1.3	0.7
1.6	1.5	1.2	0.9	0.8	0.5	0.4
5.0	4.7	3.5	2.8	2.5	2.0	1.1
3.8	5.4	7.2	8.0	9.3	12.6	16.3
2.2	3.1	4.2	4.7	5.3	7.1	9.3
5.4	7.8	10.3	11.4	13.4	18.1	23.2
2.1	3.3	4.6	5.1	6.1	8.8	11.7
1.3	1.9	2.7 6.5	3.1	3.5	5.0	6.8
2.9	4.7 2.1	2.6	7.2 2.9	8.6	12.6 3.8	16.5 4.6
1.7 0.9	1.2	1.5	1.6	3.2 1.8	2.1	2.5
2.5	3.1	3.8	4.2	4.8	5.5	6.7
0.6	1.6	3.0	3.7	4.7	7.5	10.7
0.4	0.9	1.7	2.2	2.6	4.2	6.2
0.9	2.3	4.3	5.2	6.7	10.9	15.2
3.2	3.8	4.2	4.3	4.6	5.1	5.6
1.8	2.2	2.5	2.5	2.7	2.9	3.1
4.5	5.5	6.0	6.2	6.7	7.2	8.0

12 18 2.5 2.6 2.7 3.2 4.2 3.1 47 6.2 6.4 7.1 8.7 10.5 1.4 2.3 3.2 3.4 3.8 4.8 5.9 0.8 1.3 1.8 2.0 2.1 2.6 3.4 0.7 0.9 1.1 1.1 1.1 1.1 1.2 1.4 0.4 0.5 0.7 0.6 0.6 0.6 0.6 0.8 1.0 1.4 1.7 1.6 1.6 1.7 2.1 0.5 1.3 2.3 2.8 3.2 8.3.2 4.3 5.6 0.3 0.8 1.3 1.6 1.7 2.4 3.2 0.7 0.9 1.0 1.4 1.7 1.6 1.6 1.7 2.4 3.2 0.7 0.9 3.3 3.9 4.7 6.3 7.9 1.6 1.9 2.0 1.7 1.7 1.7 1.7 1.7 1.7 0.9 1.0 1.2 1.0 1.0 0.8 1.0 0.4 2.7 2.9 2.5 2.4 2.4 2.4 2.6 1.8 2.4 3.2 3.9 5.0 7.5 10.0 1.1 1.4 1.9 2.4 2.9 4.3 5.8 0.7 1.1 1.1 1.2 2.8 4.7 0.5 0.7 1.1 1.3 1.7 2.8 4.0 0.7 1.1 1.3 1.7 2.8 4.0 0.5 0.7 1.1 1.3 1.7 2.8 4.0 0.6 0.7 0.8 1.1 1.2 1.2 1.5 1.8 0.6 0.7 0.8 1.1 1.2 1.5 1.8 0.8 3.8 0.8 1.1 1.2 2.8 3.8 0.6 0.7 0.8 1.1 1.2 2.8 3.8 0.6 0.7 0.8 1.1 1.2 1.5 1.8 0.2 0.3 0.8 1.1 1.2 1.5 1.8 0.2 0.3 0.8 1.1 1.2 1.5 1.8 0.2 0.3 0.8 1.1 1.2 2.8 3.3 0.6 0.7 0.8 1.1 1.2 1.5 1.8 0.8 0.8 0.9 0.8 1.1 1.1 1.7 2.9 3.0 0.9 0.9 1.1 1.1 1.1 1.1 1.1 1.2 1.1 1.1 1.1 1.1	2.1	3.2	4.3	4.5	4.9	6.0	7.3
14							
1.4							
0.8							
0.7 0.9 1.1 1.1 1.1 1.2 1.4 0.4 0.5 0.7 0.6 0.6 0.6 0.8 1.0 1.4 1.7 1.6 1.6 1.7 2.1 0.5 1.3 2.3 2.8 3.2 4.3 5.6 0.3 0.8 1.3 1.6 1.7 2.4 3.2 0.7 2.0 3.3 3.9 4.7 6.3 7.9 1.6 1.9 2.0 1.7 1.7 1.7 1.7 1.7 0.9 1.0 1.2 1.0 1.0 0.8 1.0 2.4 2.7 2.9 2.5 2.4 2.4 2.2 2.4 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 1.0 6.7 1.0 1.1 1.7 2.1 2.8 4.0 1.2 2.8 4.0 0.5 0.7	0.8	1.3		2.0		2.6	3.4
0.4	2.1	3.3	4.5	4.8	5.5	7.0	8.4
10	0.7	0.9	1.1	1.1	1.1	1.2	1.4
0.5 1.3 2.3 2.8 3.2 4.3 5.6 0.3 0.8 1.3 1.6 1.7 2.4 3.2 0.7 2.0 3.3 3.9 4.7 6.3 7.9 1.6 1.9 2.0 1.7 1.1 1.7 2.4 2.9 4.3 5.8 2.6 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 1.0.6 14.3 5.5 7.1 1.0.6 14.3 3.7 4.0 1.0 1.5 2.2 2.8 3.3 3.8 6.6 9.4 1.1 1.2 1.8 <td>0.4</td> <td>0.5</td> <td>0.7</td> <td>0.6</td> <td>0.6</td> <td>0.6</td> <td>0.8</td>	0.4	0.5	0.7	0.6	0.6	0.6	0.8
0.3							
0.7 2.0 3.3 3.9 4.7 6.3 7.9 1.6 1.9 2.0 1.7 1.0 0.8 1.0 2.4 2.7 2.9 2.5 2.4 2.4 2.6 2.4 2.4 2.6 1.0 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 10.6 1.4 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.3 1.7 2.8 4.0 1.0 1.5 2.2 2.8 3.3 3.6 6.9 1.1 1.3 1.5 1.8 2.2 2.8 3.3							
1.6 1.9 2.0 1.7 1.7 1.7 1.7 0.9 1.0 1.2 1.0 1.0 0.8 1.0 2.4 2.7 2.9 2.5 2.4 2.4 2.6 1.8 2.4 3.2 3.9 5.0 7.5 10.0 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 10.6 14.3 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.3 1.7 2.8 4.0 1.0 1.5 2.2 2.8 3.8 6.6 9.4 1.1 1.3 1.5 1.8 2.2 2.8 3.3 0.6 0.7 0.8 1.1 1.2 1.5 1.8 1.5 1.9 2.3 2.7 3.3 4.0 4.9 0.2 0.3 <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>							
0.9 1.0 1.2 1.0 1.0 0.8 1.0 2.4 2.7 2.9 2.5 2.4 2.4 2.4 2.6 1.8 2.4 3.2 3.9 5.0 7.5 10.0 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 10.6 13.3 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.3 1.7 2.8 3.8 6.6 9.4 1.0 1.5 2.2 2.8 3.8 6.6 9.4 1.1 1.3 1.5 1.8 2.2 2.8 3.3 0.6 0.7 0.8 1.1 1.2 1.5 1.8 1.5 1.9 2.3 2.7 3.3 4.0 4.9 0.2 0.3 0.8 1.2 1.8 3.8 6.0 0.1 0.2 0.5 0.8 1.1 2.2 3.5 0.2 0.5 1.2 1.6 2.5 5.3 8.5 1.6 2.1 2.4 2.7 3.2 3.7 4.0 1.0 1.2 1.4 1.6 1.8 2.1 2.3 2.3 2.9 3.3 3.9 4.6 5.3 5.8 2.3 2.7 2.9 2.7 2.4 2.9 3.0 3.2 4.0 4.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 2.9 3.0 3.8 3.8 2.3 2.7 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 2.9 3.1 4.4 1.6 1.8 2.1 2.9 3.0 3.2 4.0 4.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 2.2 3.9 3.6 4.7 4.6 2.3 2.9 3.1 2.8 2.6 3.4 3.4 1.9 2.3 2.4 0.4 0.4 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5							
2.4 2.7 2.9 2.5 2.4 2.4 2.6 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 10.6 14.3 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.3 1.7 2.8 4.0 1.0 1.5 2.2 2.8 3.8 6.6 9.4 1.1 1.3 1.5 1.8 2.2 2.8 3.3 0.6 0.7 0.8 1.1 1.2 1.5 1.8 1.5 1.9 2.3 2.7 3.3 4.0 4.9 0.2 0.3 0.8 1.2 1.8 3.8 6.0 0.1 0.2 0.5 0.8 1.1 2.2 3.5 0.2 0.5 0.2 0.8 1.1 2.2 3.5 1.6 2.1							
1.8 2.4 3.2 3.9 5.0 7.5 10.0 1.1 1.4 1.9 2.4 2.9 4.3 5.8 2.5 3.4 4.5 5.5 7.1 10.6 14.3 0.7 1.1 1.7 2.1 2.8 4.7 6.7 0.5 0.7 1.1 1.3 1.7 2.8 4.0 1.0 1.5 2.2 2.8 3.8 6.6 9.4 1.1 1.3 1.5 1.8 2.2 2.8 3.3 0.6 0.7 0.8 1.1 1.2 1.5 1.8 1.5 1.9 2.3 2.7 3.3 4.0 4.9 0.2 0.3 0.8 1.2 1.8 3.8 6.0 0.1 0.2 0.5 0.8 1.1 2.2 3.5 0.2 0.5 0.8 1.1 2.2 3.5 0.2 0.5 0.8 <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>							
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1.7 1.7 1.8 1.7 1.7 1.8 1.9 1.2 1.2 1.3 1.2 1.3 1.5 1.5 0.9 0.9 1.0 0.9 0.9 1.1 1.1 1.1 1.1 1.0 1.1 1.0 1.0 1.2 1.2 0.8 0.8 0.9 0.8 0.8 1.0 1.0 0.6 0.5 0.6 0.6 0.6 0.6 0.6 0.6							
1.2 1.2 1.3 1.2 1.3 1.5 1.5 0.9 0.9 1.0 0.9 0.9 1.1 1.1 1.1 1.0 1.1 1.0 1.0 1.2 1.2 0.8 0.8 0.9 0.8 0.8 1.0 1.0 0.6 0.5 0.6 0.6 0.6 0.6 0.6							
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0.6 0.5 0.6 0.6 0.6 0.6 0.6							

0.4	0.4	0.4	0.4	0.5	0.5	0.5
0.3	0.3	0.4	0.4	0.5	0.7	0.8
0.3	0.3	0.4	0.4	0.5	0.7	0.8
0.2	0.3	0.5	0.4	0.5	0.8	0.8
1.2	1.1	1.2	1.1	1.0	1.0	0.9
1.4	1.4	1.4	1.3	1.2	1.1	1.1
1.0	0.9	0.8	0.8	0.8	0.7	0.7

2015-2021 Fold
Change
2.0
1.7
2.2
2.4
2.0
2.5
1.0
1.0
1.1
4.1
3.3 4.8
0.5
0.5
0.5
7.0
6.4
7.5
0.7
0.8
0.7
2.3
2.8
2.3
0.2
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0.2
4.3
4.2
4.3
5.6 5.2
5.7
2.7
2.8
2.7
17.8
15.5
16.9
1.8
1.7

1.8

3.5
3.5
3.4
4.2
4.3 4.0
2.0
2.0
2.1
11.2
10.7
11.3 1.1
1.1
1.1
5.6
5.3
5.7
9.6
8.0 9.4
3.0
3.0
3.3
30.0
35.0
42.5
2.5
2.3 2.5
1.4
1.3
1.4
1.5
1.4 1.5
1.0
1.0
1.0
4.5
4.0
5.0 0.5
0.6
0.4
1.1
1.1
1.3
1.2 1.1
1.3
1.0
1.2

1.3
2.7
2.7
4.0
0.8
0.8
0.7



Number of National Drug Overdose Deaths* Involving Select Pr Source: National Center on Health Statistics, CDC WONDER

	1999	2000
Total Overdose Deaths	1,240	1,435
Female	346	369
Male	894	1,066
Any Opioid ¹	621	728
Female	141	140
Male	480	588
Prescription Opioids ²	228	288
Female	61	69
Male	167	219
Prescription Opioids AND Synthetic Opioids other than Methadone (primarily fentanyl)		11
Female		
Male		
Synthetic Opioids other than Methadone (primarily fentanyl) ³	33	40
Female		13
Male	28	27
Heroin⁴	198	216
Female	38	36
Male	160	180
Heroin AND Synthetic Opioids other than Methadone (primarily fentanyl)		
Female		
Male		
Stimulants ^{5a}	322	352
Female	83	94
Male	239	258
Stimulants AND Any Opioid	160	181
Female	40	43
Male	120	138
Stimulants AND Synthetic Opioids other than Methadone		
Female		
Male		
Cocaine ⁵	267	276
Female	60	72
Male	207	204
Cocaine AND Any Opioid	147	164
Female	34	39

Male	113	125
Cocaine AND Synthetic Opioids other than Methadone (primarily fentanyl)		
Female		
Male		
Psychostimulants With Abuse Potential (primarily methamphetamine) ⁶	67	97
Female	26	31
Male	41	66
Psychostimulants With Abuse Potential AND Any Opioid	22	32
Female		
Male		
Psychostimulants With Abuse Potential AND Synthetic Opioids other than Methadone (primarily fentanyl)		
Female		
Male		
Benzodiazepines ⁷	53	90
Female	12	22
Male	41	68
Benzodiazepines AND Any Opioid	37	74
Female		17
Male	31	57
Benzodiazepines AND Synthetic Opioids other than Methadone (primarily		10
fentanyl)		10
Female		
Male		
Antidepressants ⁸	59	68
Female	27	36
Male	32	32
Antidepressants AND Any Opioid	13	20
Female		
Male		
Antidepressants AND Synthetic Opioids other than Methadone (primarily		
fentanyl)		
Female		
Male		

^{*}Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug poisoning Years for which data are not provided include unreliable data

Blank fields designated by unreliable or suppressed data. For more information visit CDC WONDER.

¹ Any Opioid ICD-10 codes (T40.0-T40.4, T40.6)

² Prescription Opioids ICD-10 codes (T40.2-T40.3)

³ Synthetic Opioids other than Methadone (primarily fentanyl) ICD-10 code (T40.4) This category is domii

⁴Heroin ICD-10 codes (T40.1)

^{5a}Stimulants ICD-10 codes (T40.5 & T43.6)

⁵Cocaine ICD-10 codes (T40.5)

⁶Psychostimulants With Abuse Potential ICD-10 code (T43.6) This category is dominated by methamphet

⁷Benzodiazepines ICD-10 code(T42.4)

⁸Antidepressants ICD-10 code(T43.0-T43.2)

rescription and Illicit Drugs, Ages 15-24 Years Old

2001	2002	2003	2004	2005	2006	2007
1,700	2,095	2,491	2,751	2,918	3,460	3,550
438	560	632	683	733	820	883
1,262	1,535	1,859	2,068	2,185	2,640	2,667
944	1,179	1,399	1,596	1,685	2,096	2,176
203	269	306	333	371	444	491
741	910	1,093	1,263	1,314	1,652	1,685
489	641	833	1,031	1,058	1,388	1,536
101	152	175	202	225	290	353
388	489	658	829	833	1,098	1,183
12	21	31	28	31	52	41
					14	12
10	12	23	19	27	38	29
54	84	107	121	130	278	175
15	22	34	26	25	66	47
39	62	73	95	105	212	128
212	241	259	263	279	313	359
42	56	55	57	65	57	61
170	185	204	206	214	256	298

14

264	400	co=	62.4	740	700	67.0
364	492	607	634	710	789	674
96	139	162	161	171	202	161
268	353	445	473	539	587	513
189	251	301	343	404	476	411
47	68	65	83	103	108	93
142	183	236	260	301	368	318
		17	15	23	61	24
					17	
		14	13	20	44	17
280	388	475	507	546	676	563
68	109	118	121	132	168	119
212	279	357	386	414	508	444
167	223	266	312	345	424	365
42	56	57	76	91	94	81

12	5 167	209	236	254	330	284
		14	12	19	54	23
					37	
10	1 124	165	142	196	153	142
3	6 34	51	43	47	46	51
6	5 90	114	99	149	107	91
3	0 39	52	39	77	72	67

133	178	212	271	322	442	515
26	42	53	60	83	95	145
107	136	159	211	239	347	370
104	146	170	237	281	387	452
19	32	37	52	68	78	128
85	114	133	185	213	309	324
		22	14	30	40	38
				22	36	25
88	102	115	147	152	126	150
30	52	56	65	77	69	66
58	50	59	82	75	57	84
38	44	57	69	79	52	88
		28	28	33	24	33
26	29	29	41	46	28	55

ng (X60–X64), homicide drug poisoning (X85), or drug poisoning of undetermined intent (Y10–Y14), as coded in the

nated by fentanyl related overdoses.

amine related overdoses.	

2014	2013	2012	2011	2010	2009	2008
3,798	3,664	3,518	3,762	3,571	3,377	3,487
1,075	1,028	945	990	988	862	848
2,723	2,636	2,573	2,772	2,583	2,515	2,639
2,700	2,486	2,344	2,545	2,387	2,229	2,294
71:	640	574	624	603	521	482
1,995	1,846	1,769	1,921	1,784	1,708	1,812
933	988	1,120	1,427	1,530	1,406	1,449
252	241	289	363	389	338	299
679	747	831	1,064	1,141	1,068	1,150
8:	62	42	62	65	58	55
27	19	10	23	26	21	11
54	43	32	39	39	37	44
514	237	172	220	229	203	190
138	68	35	60	77	53	48
376	169	137	160	152	150	142
1,452	1,263	963	809	537	510	497
373	322	241	183	118	115	96
1,079	941	722	626	419	395	401
131	20					
42	7					
89	13					
635	549	452	486	429	395	500
178	166	129	134	145	98	115
457	383	323	352	284	297	385
457	349	310	346	288	271	334
120	104	87	87	93	59	71
337	245	223	259	195	212	263
65	24	13	22	15	26	26
19						
106	46	18	10	16	18	20
334	273	270	310	265	269	384
84	78	70	82	80	58	89
250	195	200	228	185	211	295
280	222	227	242	209	208	280
69	64	56	63	63	42	62

218	166	146	179	171	158	211
18	14	10	12	12	12	45
						32
127	147	179	199	197	303	340
31	46	73	63	63	100	103
96	101	106	136	134	203	237
59	79	91	123	94	148	204
	21	36	32	34	48	57
	58	55	91	60	100	147
						24

572	507	511	614	658	571	550
167	133	137	158	172	147	133
405	374	374	456	486	424	417
514	452	479	569	605	517	486
147	117	128	149	157	130	114
367	335	351	420	448	387	372
89	56	51	57	59	58	40
25						
64	39	37	44	43	40	32
206	173	159	183	198	177	157
98	75	68	83	78	76	74
108	98	91	100	120	101	83
121	101	90	114	126	103	94
49	40	36	48	51	41	37
72	61	54	66	75	62	57
30	23					

he International Classification of Diseases, 10th Revision.

2021	2020	2019	2018	2017	2016	2015
7,426	7,095	4,777	4,633	5,455	5,376	4,235
2,311	1,990	1,459	1,481	1,663	1,483	1,258
5,115	5,105	3,318	3,152	3,792	3,893	2,977
6,312	5,986	3,725	3,618	4,094	4,027	3,082
1,866	1,584	1,058	1,091	1,209	1,041	871
4,446	4,402	2,667	2,527	2,885	2,986	2,211
655	812	672	790	1,050	1,146	886
206	220	198	242	322	294	267
449	592	474	548	728	852	619
414	480	332	352	364	289	171
136	139	100	117	122	88	58
278	341	232	235	242	201	113
5,936	5,393	3,040	2,640	2,655	1,958	999
1,755	1,428	865	799	778	524	281
4,181	3,965	2,175	1,841	1,877	1,434	718
360	711	876	1,160	1,454	1,728	1,649
126	230	280	339	423	453	477
234	481	596	821	1,031	1,275	1,172
245	474	525	623	701	535	305
88	160	179	178	199	160	96
157	314	346	445	502	375	209
2,472	2,199	1,609	1,497	1,587	1,275	826
852	679	501	517	529	375	281
1,620	1,520	1,108	980	1,058	900	545
2,067	1,827	1,223	1,170	1,208	936	589
709	568	373	400	419	265	198
1,358	1,259	850	770	789	671	391
1,945	1,599	971	843	774	438	158
660	496	295	284	262	135	52
1,285	650	1,103	676	559	512	303
1,157	1,089	850	859	924	757	442
384	299	236	265	291	204	139
773	790	614	594	633	553	303
1,039	988	743	751	792	641	385
345	277	203	223	259	171	122

694	711	540	528	533	470	263
999	901	646	590	564	354	121
330	251	179	182	188	114	40
669	650	467	408	376	240	81
1,502	1,316	909	749	780	571	416
541	444	313	294	281	183	157
961	872	596	455	499	388	259
1,194	1,022	613	513	516	334	227
428	351	210	215	198	98	85
766	671	403	298	318	236	142
1,103	856	428	319	266	97	46
389	293	148	128	97	23	
714	563	280	191	169	74	30
1,118	1,267	727	899	1,031	1,046	665
329	330	208	263	298	247	189
789	937	519	636	733	799	476
1,020	1,156	662	812	942	931	602
292	292	190	236	275	217	164
728	864	472	576	667	714	438
932	1,007	486	522	509	377	152
272	257	140	162	153	92	43
660	750	346	360	356	285	109
333	263	239	237	231	200	203
185	119	127	119	111	94	105
148	144	112	118	120	106	98
172	149	128	123	125	102	104
83	53	55	52	51	40	49
89	96	73	71	74	62	55
145	122	86	81	71	35	38
72	44	35	34	26		21
73	78	51	47	45	20	

Fold Change 2015 to 2021

1.8

1.8

1.7

2.0

2.1

2.0

0.7

0.8

0.7

2.4

2.3

2.5

5.9

6.2

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3.0

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3.0 **3.5**

3.6

3.5

12.3

12.7

4.2

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8.3
8.3
8.3
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5.0
5.4
24.0
23.8
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6.1
6.1 6.3
6.1 6.3 6.1
1.7 6.1 6.3 6.1 1.6
1.7 6.1 6.3 6.1 1.6 1.8
1.7 6.1 6.3 6.1 1.6
1.7 6.1 6.3 6.1 1.6 1.8 1.5
1.7 6.1 6.3 6.1 1.6 1.8 1.5
1.7 6.1 6.3 6.1 1.6 1.8 1.5 1.7
1.7 6.1 6.3 6.1 1.6 1.8 1.5 1.7 1.7



Rate of National Drug Overdose Deaths* Involving Select |

Source: National Center on Health Statistics, CDC WONDER

	1999	2000
Total Overdose Deaths	3.2	3.7
Female	1.8	1.9
Male	4.5	5.3
Any Opioid ¹	1.6	1.9
Female	0.7	0.7
Male	2.4	2.9
Prescription Opioids ²	0.6	0.7
Female	0.3	0.4
Male	0.8	1.1
Prescription Opioids AND Synthetic Opioids other than Methadone Female		
Male		
Synthetic Opioids other than Methadone (primarily fentanyl) ³	0.1	0.1
Female		
Male	0.1	0.1
Heroin ⁴	0.5	0.6
Female	0.2	0.2
Male	0.8	0.9
Heroin AND Synthetic Opioids other than Methadone		
Female		
Male		
Stimulants ^{5a}	0.8	0.9
Female	0.4	0.5
Male	1.2	1.3
Stimulants AND Any Opioid	0.4	0.5
Female	0.2	0.2
Male	0.6	0.7
Stimulants AND Synthetic Opioids other than Methadone		
Female		
Male		
Cocaine ⁵	0.7	0.7
Female	0.3	0.4
Male	1.0	1.0
Cocaine AND Any Opioid	0.4	0.4
Female	0.2	0.2
Male	0.6	0.6

Cocaine AND Synthetic Opioids other than Methadone		
Female		
Male		
Psychostimulants With Abuse Potential (primarily	0.2	0.:
methamphetamine) ⁶		
Female	0.1	0.2
Male	0.2	0
Psychostimulants With Abuse Potential AND Any Opioid	0.1	0.
Female		
Male		0.
Psychostimulants With Abuse Potential AND Synthetic Opioids other		
than Methadone		
Female		
Male		
Benzodiazepines ⁷	0.1	0.:
Female		0.
Male	0.2	0.3
Benzodiazepines AND Any Opioid	0.1	0.2
Female		
Male	0.2	0.3
Benzodiazepines AND Synthetic Opioids other than Methadone		
Female		
Male		
Antidepressants ⁸	0.2	0.3
Female	0.1	0
Male	0.2	0
Antidepressants AND Any Opioid		0.:
Female		
Male		
Antidepressants AND Synthetic Opioids other than Methadone		
Female		
Male		

^{*}Includes deaths with underlying causes of unintentional drug poisoning (X40–X44), suicide drug Revision.

Years for which data are not provided include unreliable data

Blank fields designated by unreliable or suppressed data. For more information visit CDC WONDER.

¹ Any Opioid ICD-10 codes (T40.0-T40.4, T40.6)

² Prescription Opioids ICD-10 codes (T40.2-T40.3)

³Other Synthetic Narcotics (other than methadone) ICD-10 code (T40.4) This category is domina

⁴Heroin ICD-10 codes (T40.1)

^{5a}Stimulants ICD-10 codes (T40.5 & T43.6)

⁵Cocaine ICD-10 codes (T40.5)

⁶Psychostimulants With Abuse Potential ICD-10 code (T43.6) This category is dominated by meth

⁷Benzodiazepines ICD-10 code(T42.4)

⁸Antidepressants ICD-10 code(T43.0-T43.2)

Prescription and Illicit Drugs, Ages 15-24 Years Old

2001	2002	2003	2004	2005	2006	2007
4.2	5.1	6.0	6.6	6.9	8.1	8.2
2.2	2.8	3.1	3.3	3.5	3.9	4.2
6.1	7.3	8.8	9.6	10.0	12.0	12.0
2.3	2.9	3.4	3.8	4.0	4.9	5.0
1.0	1.4	1.5	1.6	1.8	2.1	2.3
3.6	4.3	5.2	5.9	6.0	7.5	7.6
1.2	1.6	2.0	2.5	2.5	3.2	3.6
0.5	0.8	0.9	1.0	1.1	1.4	1.7
1.9	2.3	3.1	3.9	3.8	5.0	5.3
	0.1	0.1	0.1	0.1	0.1	0.1
		0.1		0.1	0.2	0.1
0.1	0.2	0.3	0.3	0.3	0.6	0.4
	0.1	0.2	0.1	0.1	0.3	0.2
0.2	0.3	0.3	0.4	0.5	1.0	0.6
0.5	0.6	0.6	0.6	0.7	0.7	0.8
0.2	0.3	0.3	0.3	0.3	0.3	0.3
0.8	0.9	1.0	1.0	1.0	1.2	1.3

0.9	1.2	1.5	1.5	1.7	1.8	1.6
0.5	0.7	0.8	0.8	0.8	1.0	0.8
1.3	1.7	2.1	2.2	2.5	2.7	2.3
0.5	0.6	0.7	0.8	1.0	1.1	1.0
0.2	0.3	0.3	0.4	0.5	0.5	0.4
0.7	0.9	1.1	1.2	1.4	1.7	1.4
				0.1	0.1	0.1
				0.1	0.2	
0.7	0.9	1.1	1.2	1.3	1.6	1.3
0.3	0.5	0.6	0.6	0.6	0.8	0.6
1.0	1.3	1.7	1.8	1.9	2.3	2.0
0.4	0.5	0.6	0.7	0.8	1.0	0.8
0.2	0.3	0.3	0.4	0.4	0.5	0.4
0.6	0.8	1.0	1.1	1.2	1.5	1.3

					0.1	0.1
					0.2	
0.3	0.3	0.4	0.3	0.5	0.4	0.3
0.2	0.2	0.3	0.2	0.2	0.2	0.2
0.3	0.4	0.5	0.5	0.7	0.5	0.4
0.1	0.1	0.1	0.1	0.2	0.2	0.2
0.1	0.1	0.2	0.1	0.3	0.2	0.2

0.3	0.4	0.5	0.6	0.8	1.0	1.2
0.1	0.2	0.3	0.3	0.4	0.5	0.7
0.5	0.6	0.7	1.0	1.1	1.6	1.7
0.3	0.4	0.4	0.6	0.7	0.9	1.0
	0.2	0.2	0.3	0.3	0.4	0.6
0.4	0.5	0.6	0.9	1.0	1.4	1.5
		0.1		0.1	0.1	0.1
				0.1	0.2	0.1
0.2	0.2	0.3	0.4	0.4	0.3	0.3
0.2	0.3	0.3	0.3	0.4	0.3	0.3
0.3	0.2	0.3	0.4	0.3	0.3	0.4
0.1	0.1	0.1	0.2	0.2	0.1	0.2
		0.1	0.1	0.2	0.1	0.2
0.1	0.1	0.1	0.2	0.2	0.1	0.2

poisoning (X60–X64), homicide drug poisoning (X85), or drug poisoning of undetermined intent (Y10–Y14), as co

ted by fentanyl related overdoses.

namphetamine related overdoses.

		_
		_
		_
		_
		_

2008	2009	2010	2011	2012	2013	2014
8.0	7.7	8.2	8.6	8.0	8.3	8.6
4.0	4.1	4.6	4.6	4.4	4.8	5.0
11.9	11.3	11.6	12.4	11.4	11.7	12.1
5.3	5.1	5.5	5.8	5.3	5.7	6.2
2.3	2.5	2.8	2.9	2.7	3.0	3.3
8.1	7.7	8.0	8.6	7.9	8.2	8.9
3.3	3.2	3.5	3.3	2.5	2.2	2.1
1.4	1.6	1.8	1.7	1.3	1.1	1.2
5.2	4.8	5.1	4.7	3.7	3.3	3.0
0.1	0.1	0.1	0.1	0.1	0.1	0.2
	0.1	0.1	0.1			0.1
0.2	0.2	0.2	0.2	0.1	0.2	0.2
0.4	0.5	0.5	0.5	0.4	0.5	1.2
0.2	0.2	0.4	0.3	0.2	0.3	0.6
0.6	0.7	0.7	0.7	0.6	0.8	1.7
1.1	1.2	1.2	1.8	2.2	2.9	3.3
0.5	0.5	0.6	0.9	1.1	1.5	1.7
1.8	1.8	1.9	2.8	3.2	4.2	4.8
					0.0	0.3
						0.2
						0.4
1.2	0.9	1.0	1.1	1.0	1.2	1.4
0.5	0.5	0.7	0.6	0.6	0.8	0.8
1.7	1.3	1.3	1.6	1.4	1.7	2.0
0.8	0.6	0.7	0.8	0.7	0.8	1.0
0.3	0.3	0.4	0.4	0.4	0.5	0.6
1.2	1.0	0.9	1.2	1.0	1.1	1.5
0.1	0.1		0.1		0.1	0.1
0.1					0.2	0.5
0.9	0.6	0.6	0.7	0.6	0.6	0.8
0.4	0.3	0.4	0.4	0.3	0.4	0.4
1.3	0.9	0.8	1.0	0.9	0.9	1.1
0.6	0.5	0.5	0.6	0.5	0.5	0.6
0.3	0.2	0.3	0.3	0.3	0.3	0.3
1.0	0.7	0.7	0.8	0.8	0.7	0.9

0.1

0.1

0.3	0.3	0.4	0.5	0.4	0.7	0.8
0.1	0.2	0.3	0.3	0.3	0.5	0.5
0.4	0.5	0.5	0.6	0.6	0.9	1.1
0.1	0.2	0.2	0.3	0.2	0.3	0.5
	0.1	0.2	0.1	0.2	0.2	0.3
0.2	0.3	0.2	0.4	0.3	0.4	0.7

0.1

1.3	1.3	1.5	1.4	1.2	1.2	1.3
0.6	0.7	0.8	0.7	0.6	0.6	0.8
1.9	1.9	2.2	2.0	1.7	1.7	1.8
1.1	1.2	1.4	1.3	1.1	1.0	1.2
0.5	0.6	0.7	0.7	0.6	0.5	0.7
1.7	1.7	2.0	1.9	1.6	1.5	1.6
0.1	0.1	0.1	0.1	0.1	0.1	0.2
						0.1
0.1	0.2	0.2	0.2	0.2	0.2	0.3
0.4	0.4	0.5	0.4	0.4	0.4	0.5
0.3	0.4	0.4	0.4	0.3	0.3	0.5
0.4	0.5	0.5	0.4	0.4	0.4	0.5
0.2	0.2	0.3	0.3	0.2	0.2	0.3
0.2	0.2	0.2	0.2	0.2	0.2	0.2
0.3	0.3	0.3	0.3	0.2	0.3	0.3
					0.1	0.1

oded in the International Classification of Diseases, 10th

2015	2016	2017	2018	2019	2020	2021
9.7	12.4	12.6	10.8	11.2	16.7	17.2
5.9	7.0	7.9	7.1	7.0	9.6	11.0
13.3	17.5	17.1	14.3	15.2	23.5	23.3
7.0	9.3	9.5	8.4	8.7	14.1	14.6
4.1	4.9	5.7	5.2	5.1	7.6	8.8
9.8	13.4	13.0	11.5	12.2	20.3	20.2
2.0	2.6	2.4	1.8	1.6	1.9	1.5
1.2	1.4	1.5	1.2	0.9	1.1	1.0
2.8	3.8	3.3	2.5	2.2	2.7	2.0
0.4	0.7	0.8	0.8	0.8	1.1	1.0
0.3	0.4	0.6	0.6	0.5	0.7	0.6
0.5	0.9	1.1	1.1	1.1	1.6	1.3
2.3	4.5	6.1	6.1	7.1	12.7	13.8
1.3	2.5	3.7	3.8	4.1	6.9	8.3
3.2	6.4	8.5	8.4	10.0	18.2	19.0
3.8	4.0	3.4	2.7	2.1	1.7	0.8
2.2	2.1	2.0	1.6	1.3	1.1	0.6
5.2	5.7	4.7	3.7	2.7	2.2	1.1
0.7	1.2	1.6	1.4	1.2	1.1	0.6
0.4	0.8	0.9	0.8	0.9	0.8	0.4
0.9	1.7	2.3	2.0	1.6	1.4	0.7
1.9	2.9	3.7	3.5	3.8	5.2	5.7
1.3	1.8	2.5	2.5	2.4	3.3	4.0
2.4	4.0	4.8	4.5	5.1	7.0	7.4
1.3	2.2	2.8	2.7	2.9	4.3	4.8
0.9	1.2	2.0	1.9	1.8	2.7	3.4
1.7	3.0	3.6	3.5	3.9	5.8	6.2
0.4	1.0	1.8	2.0	2.3	3.8	4.5
0.2	0.6	1.2	1.4	1.4	2.4	3.1
1.4	2.3	2.5	3.1	5.1	3.0	5.8
1.0	1.7	2.1	2.0	2.0	2.6	2.7
0.7	1.0	1.4	1.3	1.1	1.4	1.8
1.3	2.5	2.9	2.7	2.8	3.6	3.5
0.9	1.5	1.8	1.7	1.7	2.3	2.4
0.6	0.8	1.2	1.1	1.0	1.3	1.6
1.2	2.1	2.4	2.4	2.5	3.3	3.2

0.3	0.8	1.3	1.4	1.5	2.1	2.3
0.2	0.5	0.9	0.9	0.9	1.2	1.6
0.4	1.1	1.7	1.9	2.1	3.0	3.0
0.9	1.3	1.8	1.7	2.1	3.1	3.5
0.7	0.9	1.3	1.4	1.5	2.1	2.6
1.2	1.7	2.3	2.1	2.7	4.0	4.4
0.5	0.8	1.2	1.2	1.4	2.4	2.8
0.4	0.5	0.9	1.0	1.0	1.7	2.0
0.6	1.1	1.4	1.4	1.8	3.1	3.5
0.1	0.2	0.6	0.7	1.0	2.0	2.6
	0.1	0.5	0.6	0.7	1.4	1.8
0.1	0.3	0.8	0.9	1.3	2.6	3.2
1.5	2.4	2.4	2.1	1.7	3.0	2.6
0.9	1.2	1.4	1.3	1.0	1.6	1.6
2.1	3.6	3.3	2.9	2.4	4.3	3.6
1.4	2.1	2.2	1.9	1.6	2.7	2.4
0.8	1.0	1.3	1.1	0.9	1.4	1.4
1.9	3.2	3.0	2.6	2.2	4.0	3.3
0.3	0.9	1.2	1.2	1.1	2.4	2.2
0.2	0.4	0.7	0.8	0.7	1.2	1.3
0.5	1.3	1.6	1.6	1.6	3.5	3.0
0.5	0.5	0.5	0.6	0.6	0.6	0.8
0.5	0.4	0.5	0.6	0.6	0.6	0.9
0.4	0.5	0.5	0.5	0.5	0.7	0.7
0.2	0.2	0.3	0.3	0.3	0.4	0.4
0.2	0.2	0.2	0.2	0.3	0.3	0.4
0.2	0.3	0.3	0.3	0.3	0.4	0.4
0.1	0.1	0.2	0.2	0.2	0.3	0.3
0.1		0.1	0.2	0.2	0.2	0.3
	0.1	0.2	0.2	0.2	0.4	0.3

Fold Change 2015 to 2021 1.8 1.9 1.8 2.1 2.1 2.1 0.8 8.0 0.7 2.5 2.0 2.6 6.0 6.4 5.9 0.2 0.3 0.2 0.9 1.0 0.8 3.0 3.1 3.1 3.7 3.8 3.6 11.3 15.5 4.1 2.7 2.6 2.7

2.72.72.7

7.7
8.0
7.5
3.9
3.7
3.7
5.6
5.0
5.8
26.0
32.0
1.7
1.8
1.7
1.7
1.8
1.7
7.3
6.5
6.0
1.6
1.8
1.8
2.0
2.0
2.0
3.0
3.0



Rate of National Drug Overdose Deaths, by Demographic

Rates are Age-Adjusted per 100,000 population

Source: National Center on Health Statistics, CDC WONDER

	1999	2000	2001	2002	2003
Total Overdose Deaths	6.1	6.2	6.8	8.2	8.9
Female	3.9	4.1	4.6	5.8	6.4
Male	8.2	8.3	9.0	10.6	11.5
White (Non-Hispanic)	6.2	6.6	7.4	9.2	10.2
Female	4.3	4.5	5.3	6.8	7.5
Male	8.0	8.6	9.6	11.6	12.9
Black (Non-Hispanic)	7.5	7.3	7.6	8.2	8.2
Female	4.0	4.2	4.4	5.1	5.4
Male	11.5	10.9	11.2	11.7	11.6
Asian* (Non-Hispanic)					
Female					
Male					
Native Hawaiin or Other Pacific Islander* (Non-Hispanic)					
Female					
Male					
Hispanic	5.4	4.6	4.5	5.4	5.6
Female	2.2	2.0	2.2	2.7	2.9
Male	8.6	7.1	6.7	8.0	8.3
American Indian or Alaska Native (Non-Hispanic)	6.0	5.5	6.9	8.5	10.8
Female	5.2	4.3	6.5	7.1	9.4
Male	6.7	6.7	7.5	10.0	12.2
Any Opioid ¹	2.9	3.0	3.3	4.1	4.5
Female	1.4	1.6	1.9	2.6	2.8
Male	4.3	4.4	4.8	5.7	6.1
White (Non-Hispanic)	2.8	3.1	3.7	4.7	5.2
Black (Non-Hispanic)	3.5	3.5	3.3	3.6	3.5
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					
Hispanic	3.5	2.7	2.6	3.2	3.2
American Indian or Alaska Native (Non-Hispanic)	2.9	2.7	3.5	4.1	5.3
Prescription Opioids ²	1.2	1.3	1.7	2.3	2.6
Female	0.7	0.9	1.1	1.6	1.8
Male	1.7	1.8	2.2	2.9	3.3
White (Non-Hispanic)	1.3	1.6	2.0	2.8	3.2
Black (Non-Hispanic)	0.8	0.8	0.9	1.2	1.1
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					

Hispanic	1.6	1.1	1.1	1.4	1.5
American Indian or Alaska Native (Non-Hispanic)	1.3	1.3	2.0	2.6	3.3
ynthetic Opioids other than Methadone (primarily fentany	0.3	0.3	0.3	0.4	0.5
Female	0.2	0.3	0.3	0.4	0.4
Male	0.3	0.3	0.4	0.5	0.
White (Non-Hispanic)	0.3	0.3	0.4	0.6	0.
Black (Non-Hispanic)	0.1	0.1	0.2	0.2	0.
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					
Hispanic	0.1	0.1	0.1	0.2	0.
American Indian or Alaska Native (Non-Hispanic)					
leroin ⁴	0.7	0.7	0.6	0.7	0.
Female	0.2	0.2	0.2	0.2	0.
Male	1.2	1.1	1.0	1.2	1.
White (Non-Hispanic)	0.7	0.6	0.6	0.7	0.
Black (Non-Hispanic)	0.8	0.9	0.8	0.9	0.
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					
Hispanic	1.1	0.9	0.9	1.0	1.
American Indian or Alaska Native (Non-Hispanic)					
timulants ^{5a}	1.5	1.4	1.5	1.9	2.
Female	0.7	0.7	0.8	1.0	1.
Male	2.3	2.2	2.3	2.8	3.
White (Non-Hispanic)	1.2	1.2	1.3	1.7	2.
Black (Non-Hispanic)	3.7	3.4	3.6	4.0	4.
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					
Hispanic	1.8	1.4	1.5	1.8	1.
American Indian or Alaska Native (Non-Hispanic)	1.1	1.2	1.4	1.6	2.
ocaine ⁵	1.4	1.3	1.3	1.6	1.
Female	0.6	0.6	0.7	0.8	0.
Male	2.1	1.9	2.0	2.4	2.
White (Non-Hispanic)	1.0	1.0	1.0	1.3	1.
Black (Non-Hispanic)	3.7	3.3	3.6	4.0	4.
Asian* (Non-Hispanic)					
Native Hawaiian or Other Pacific Islander* (Non-Hispanic)					
Hispanic	1.7	1.3	1.3	1.5	1.
American Indian or Alaska Native (Non-Hispanic)	0.9	1.0	1.0	1.1	1.
sychostimulants With Abuse Potential (primarily	0.2	0.2	0.2	0.2	_
nethamphetamine) ⁶	0.2	0.2	0.2	0.3	0.
Female	0.1	0.1	0.1	0.2	0.
Male	0.3	0.3	0.3	0.5	0.
		0.2	0.2	0.4	0.
	0.2	0.2	0.2	0.4	
White (Non-Hispanic) Black (Non-Hispanic)	0.2	0.2	0.2	0.1	
White (Non-Hispanic)		0.2	0.2		0.

Hispanic 0.2 0.2 0.2 0.3 0.4

American Indian or Alaska Native (Non-Hispanic)

¹ Any Opioid ICD-10 codes: T40.0-T40.4, T40.6

² Prescription Opioids ICD-10 codes: T40.2-T40.3

³Synthetic Opioids other than Methadone (Primarily Fentanyl) ICD-10 Code: T40.4

⁴Heroin ICD-10 codes: T40.1

^{5a}Stimulants ICD-10 codes (T40.5 & T43.6)

⁵Cocaine ICD-10 codes (T40.5)

⁶Psychostimulants With Abuse Potential ICD-10 code (T43.6) This category is dominated by met

^{*} Prior to 2018, mortlity data for Asian and Pacific Islander populations were combined. See http Blank fields designated by unreliable or suppressed data. For more information visit CDC WONDER.

2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
9.4	10.1	11.5	11.9	11.9	11.9	12.3	13.2	13.1	13.8	14.7	16.3	19.8	21.7
6.9	7.3	8.2	8.8	8.9	9.1	9.6	10.2	10.2	10.6	11.1	11.8	13.4	14.4
11.8	12.8	14.8	14.9	14.9	14.8	15.0	16.1	16.1	17.0	18.3	20.8	26.2	29.1
11.0	11.8	13.6	14.5	14.8	15.0	15.7	16.9	16.8	17.6	19.0	21.1	25.3	27.5
8.3	8.8	10.0	11.0	11.4	11.6	12.5	13.3	13.2	13.8	14.6	15.8	17.7	19.1
13.7	14.7	17.2	18.0	18.3	18.3	19.0	20.5	20.4	21.4	23.2	26.2	32.7	35.8
8.3	9.3	10.8	9.7	8.4	8.3	8.0	8.5	8.7	9.7	10.5	12.2	17.1	20.6
5.6	6.2	6.5	6.5	5.6	5.8	5.9	6.2	6.2	6.5	7.3	7.7	10.4	11.7
11.4	13.0	15.7	13.3	11.6	11.1	10.5	11.3	11.6	13.3	14.2	17.3	24.7	30.8
5.2	5.8	6.3	5.9	5.8	5.8	5.6	6.1	6.3	6.7	6.7	7.7	9.5	10.6
2.9	3.0	3.4	3.1	3.2	3.5	3.6	4.0	4.0	4.1	4.1	4.4	5.0	5.1
7.5	8.4	9.1	8.7	8.4	8.2	7.6	8.1	8.5	9.2	9.3	10.9	13.9	15.9
12.5	13.1	14.1	14.2	15.8	17.7	16.8	18.2	18.9	18.7	20.9	21.2	24.2	25.7
10.4	11.3	10.6	13.8	12.2	14.0	14.8	15.9	17.6	17.5	16.9	16.8	20.2	20.1
14.7	14.9	17.6	14.5	19.6	21.5	19.0	20.5	20.1	19.9	25.0	25.8	28.3	31.5
4.7	5.1	5.9	6.1	6.4	6.6	6.8	7.3	7.4	7.9	9.0	10.4	13.3	14.9
3.1	3.4	3.9	4.3	4.4	4.6	4.9	5.2	5.3	5.6	6.3	7.1	8.5	9.4
6.3	6.6	7.8	8.0	8.4	8.7	8.7	9.4	9.5	10.2	11.7	13.7	18.1	20.4
5.7	6.2	7.3	7.8	8.2	8.6	9.1	9.7	9.8	10.5	12.0	13.9	17.5	19.4
3.2	3.4	4.2	3.6	3.4	3.6	3.4	3.8	4.0	4.7	5.6	6.6	10.3	12.9
2.9	3.0	3.3	3.2	3.4	3.2	2.9	3.3	3.5	3.8	4.0	4.6	6.1	6.8
6.2	6.9	7.1	7.8	9.3	11.3	9.4	10.3	11.0	10.6	12.0	12.1	13.9	15.7
2.9	3.2	3.9	4.2	4.3	4.4	4.7	4.9	4.5	4.4	4.6	4.7	5.2	5.2
2.1	2.4	2.8	3.2	3.2	3.3	3.6	3.8	3.7	3.7	3.9	4.0	4.3	4.2
3.7	4.1	4.9	5.3	5.4	5.5	5.8	5.9	5.3	5.1	5.2	5.4	6.2	6.1
3.7	4.1	5.0	5.6	5.7	5.8	6.4	6.6	6.0	5.9	6.2	6.4	7.0	6.9
1.4	1.6	1.8	1.8	1.8	1.9	2.0	2.0	2.0	2.2	2.5	2.6	3.3	3.5

1.4	1.5	1.8	1.9	1.9	1.7	1.7	1.8	1.8	1.9	1.8	1.8	2.1	2.2
4.2	4.7	4.7	5.2	6.1	7.8	7.2	7.1	7.5	7.1	7.2	7.0	6.5	7.2
0.6	0.6	0.9	0.7	0.8	1.0	1.0	0.8	0.8	1.0	1.8	3.1	6.2	9.0
0.5	0.5	0.7	0.7	0.7	0.9	0.9	0.8	0.7	0.9	1.3	1.9	3.5	5.0
0.6	0.6	1.1	0.8	0.8	1.0	1.0	0.9	0.9	1.1	2.2	4.2	8.9	13.0
0.7	0.8	1.1	1.0	1.0	1.2	1.3	1.2	1.1	1.3	2.4	4.2	8.2	11.9
0.2	0.2	1.0	0.2	0.2	0.3	0.4	0.3	0.3	0.5	1.1	2.1	5.6	9.0
0.2	0.2	0.3	0.2	0.2	0.3	0.2	0.3	0.3	0.3	0.6	0.9	2.7	3.7
0.9	1.1	1.2	0.9	1.8	2.4	1.2	1.4	1.6	1.2	1.7	2.0	4.1	6.5
0.6	0.7	0.7	0.8	1.0	1.1	1.0	1.4	1.9	2.7	3.4	4.1	4.9	4.9
0.2	0.3	0.2	0.3	0.4	0.4	0.4	0.6	0.8	1.2	1.6	2.0	2.4	2.5
1.1	1.1	1.2	1.3	1.6	1.8	1.6	2.3	3.1	4.2	5.2	6.3	7.5	7.3
0.7	0.7	0.8	0.9	1.2	1.3	1.2	1.8	2.5	3.4	4.4	5.4	6.3	6.1
0.6	0.8	0.7	0.8	0.8	0.9	0.8	1.0	1.3	2.0	2.5	3.1	4.5	4.9
0.7	0.8	0.8	0.8	1.0	1.0	0.8	1.0	1.3	1.6	1.9	2.3	2.8	2.9
			0.9		1.3	1.1	1.8	1.7	2.7	3.7	4.4	5.0	5.2
2.3	2.6	2.9	2.6	2.1	1.9	1.9	2.2	2.2	2.6	3.0	3.8	5.4	7.2
1.2	1.4	1.5	1.4	1.1	1.0	1.1	1.3	1.3	1.5	1.7	2.2	3.1	4.2
3.3	3.8	4.4	3.8	3.1	2.8	2.8	3.1	3.1	3.7	4.2	5.4	7.8	10.3
2.2	2.4	2.7	2.5	2.1	1.9	1.9	2.2	2.3	2.8	3.3	4.3	6.1	8.3
4.4	5.0	5.8	4.8	3.5	3.4	3.3	3.3	3.2	3.7	3.8	4.7	7.1	9.5
1.8	2.2	2.4	1.9	1.6	1.5	1.4	1.6	1.6	2.0	1.9	2.5	3.4	4.3
2.7	3.3	2.8	3	2.5	2.8	3.3	3.6	3.9	4.7	5.9	6.7	8.7	10.7
1.9	2.1	2.5	2.2	1.7	1.4	1.3	1.5	1.4	1.6	1.7	2.1	3.2	4.3
1.0	1.1	1.3	1.1	0.9	0.7	0.7	0.8	0.8	0.9	1.0	1.2	1.8	2.5
2.8	3.1	3.8	3.2	2.5	2.1	2.0	2.2	2.0	2.3	2.4	3.1	4.7	6.2
1.7	1.9	2.3	2.0	1.6	1.3	1.2	1.4	1.3	1.5	1.7	2.2	3.4	4.6
4.3	4.9	5.6	4.7	3.4	3.2	3.1	3.1	3.0	3.4	3.4	4.0	6.1	8.3
1.4	4.7	1.0	1.6	4.2	4.4	0.0	4.4	1.0	4.4	1.0	4.2	2.0	2.5
1.4	1.7	1.9	1.6	1.3	1.1	0.9	1.1	1.0	1.1	1.0	1.3	2.0	2.5
1.6	2.3	1.6	2.1	1.5	1.8	1.9	1.6	1.5	1.3	1.4	1.6	2.1	2.5
0.4	0.5	0.5	0.4	0.4	0.5	0.6	0.7	0.8	1.2	1.4	1.8	2.4	3.2
0.3	0.3	0.3	0.3	0.2	0.3	0.4	0.4	0.5	0.7	0.8	1.1	1.4	1.9
0.6	0.8	0.7	0.7	0.6	0.8	0.8	1.0	1.2	1.6	1.9	2.5	3.4	4.5
0.5	0.6	0.6	0.6	0.5	0.6	0.7	0.9	1.0	1.4	1.7	2.2	3.0	4.2
0.1	0.2	0.2	0.2	0.2	0.1	0.2	0.2	0.3	0.4	0.5	0.8	1.2	1.6

0.4	0.5	0.5	0.4	0.4	0.4	0.5	0.5	0.6	0.9	1.0	1.4	1.5	2.0
1.3	1.5	1.2	0.9	1.1	1.2	1.5	2.4	2.6	3.5	4.5	5.4	6.9	8.5

:hamphetamine related overdoses. ps://wonder.cdc.gov/mcd-icd10.html

2018	2019	2020	2021	Fold Change
				2015 to 2021
20.7	21.6	28.3	32.4	2.0
13.6	13.7	17.1	19.6	1.7
27.9	29.6	39.5	45.1	2.2
25.9	26.2	33.1	36.8	1.7
18.0	17.6	21.5	23.8	1.5
33.8	34.5	44.5	49.4	1.9
21.3	24.8	35.8	44.2	3.6
11.6	13.4	18.9	23.5	3.1
32.4	37.7	54.8	67.3	3.9
3.0	3.3	4.6	4.7	
1.8	1.6	2.2	2.3	
4.4	5.2	7.2	7.2	
12.3	9.5	13.7	20.1	
177	12.4	6.6	9.9	
17.7	13.4	20.8	30.0	2.7
11.0	12.7	17.6	21.1	2.7
5.2	5.7	7.5	9.4	2.1
16.6	19.5	27.3	32.4	3.0
26.8	30.5	42.5	56.5	2.7
20.7	24.5	34.0	44.1	2.6
33.1 14.6	36.6	51.2	69.3	2.7
	15.5	21.4	24.7	2.4
9.0	9.3	12.3	14.5	2.0
20.1	21.7	30.4	34.8	2.5
18.8	19.2	25.5	28.4	2.0
14.1	17.3	26.6	33.5	5.1
1.3	1.5	2.6	2.6	
4.0	0.0	6.0	9.7	2.5
7.5	8.8	13.1	16.0	3.5
13.8	17.7	28.1	38.7	3.2
4.5	4.2	4.9	4.9	1.0
3.7	3.4	3.8	3.8	1.0
5.3	5.1	6.1	6.0	1.1
6.0	5.5	6.3	6.3	1.0
3.3	3.5	4.7	4.9	1.9
0.4	0.4	0.5	0.5	

2.0	2.0	2.5	2.5	1.4
4.7	5.4	6.5	7.4	1.1
9.9	11.4	17.8	21.8	7.0
5.5	6.3	9.6	12.2	6.4
14.2	16.6	25.9	31.4	7.5
12.7	14.0	20.9	24.8	5.9
11.2	14.4	24.1	31.4	15.0
0.9	1.1	2.2	2.2	13.0
0.5	1.1	4.5	8.9	
4.7	6.3	10.7	14.2	15.8
7.0	10.8	21.6	33.6	16.8
4.7	4.4	4.1	1.0	0.2
2.3	2.2	2.0	1.0	0.5
7.1	6.6	6.1	1.0	0.2
5.9	5.2	4.8	3.1	0.6
4.9	5.1	5.0	3.6	1.2
0.3	0.4	0.5	0.3	1.2
0.5	0.4	0.5	0.5	
3.1	3.1	3.0	2.3	1.0
4.8	5.3	5.4	4.6	1.0
8.0	9.3	12.6	16.3	4.3
4.7	5.3	7.1	9.3	4.2
11.4	13.4	18.1	23.2	4.3
9.1	10.5	13.9	17.6	4.1
10.8	13.0	18.1	25.0	5.3
1.3	1.6	2.2	2.3	3.3
8.4	7.3	9.4	13.6	
5.2	6.2	8.6	11.2	4.5
13.6	16.3	22.7	33.1	4.9
4.5	4.9	6.0	7.3	3.5
2.6	2.7	3.2	4.2	3.5
6.4	7.1	8.7	10.5	3.4
4.7	4.7	5.4	6.2	2.8
9.1	10.8	14.3	19.6	4.9
0.5	0.7	1.0	0.9	4.5
0.5	0.7	1.0	0.5	
3.0	3.4	4.4	5.4	4.2
3.0	3.9	5.6	7.0	4.4
3.9	5.0	7.5	10.0	5.6
2.4	2.9	4.3	5.8	5.3
5.5	7.1	10.6	14.3	5.7
5.0	6.5	9.4	12.6	5.7
2.2	2.9	5.1	7.2	9.0
0.9	1.0	1.4	1.5	
7.4	6.6	9.0	11.8	
			,	

 2.5
 3.1
 4.7
 6.4
 4.6

 11.1
 13.1
 17.9
 27.4
 5.1

	Number	Deaths per 100,000	
	Number	population	
1980	6,094	2.7	
1981	6,227	2.7	
1982	6,299	2.7	
1983	6,445	2.8	
1984	6,723	2.8	
1985	7,082	3.0	
1986	7,969	3.3	
1987	7,920	3.3	
1988	9,031	3.7	
1989	9,275	3.8	
1990	8,413	3.4	
1991	9,392	3.7	
1992	10,604	4.1	
1993	12,133	4.7	
1994	12,714	4.8	
1995	12,779	4.8	
1996	13,227	4.9	
1997	14,445	5.3	
1998	15,315	5.5	
1999	16,849	6.0	
2000	17,415	6.2	
2001	19,394	6.8	
2002	23,518	8.2	
2003	25,785	8.9	
2004	27,424	9.3	
2005	29,813	10.1	
2006	34,425	11.5	
2007	36,010	11.9	
2008	36,450	12.0	
2009	37,004		
2010	38,329		
2011	41,340		
2012	41,502		
2013	43,982		
2014	47,055		
2015	52,404		
2016	63,632		
2017	70,237		
2018	67,367		
2019	70,630		
	852,867		

NOTE: Deaths were classified using the International Classification of Diseases, Tenth Revision (ICD-10) in 1999–2008 and using the N X49, X60-X69, X85-X90, Y10-Y19, Y35.2, or

*U01(.6-.7) and ICD-9 UCODs are: E850.0–E869.9, E950.0–E952.9, E962(.0–.9), E980.0–E982.9, E972. Drug poisoning ICD-10 UCODs Motor vehicle traffic ICD-10 UCODs are: V30-V39 (.4-.9), V40-V49 (.4-.9), V50-V59 (.4-.9), V60-V69 (.4-.9), V70-V79 (.4-.9), V81.1 V8. V20-V28 (.3-.9),V29 (.4-.9),V12-V14 (.3-.9),V19 (.4-.6), V02-V04 (.1,.9),V09.2,V80 (.3-

.5),V87(.0-.8),V89.2 and ICD-9 UCODs are: E810.0–E819.9, E958.5, E988.5. When the ICD-10

replaced ICD-9 in 1999, approximately 5% fewer deaths were classified as motor vehicle deaths and 2% more deaths were classified

SOURCE: CDC/NCHS, National Vital Statistics System.

Data Brief 81: Drug Poisoning Deaths in the United States, 1980–2008

Data table for Figure 2. Age-adjusted poisoning death rates: Comparison of state and U.S. rates: United States, 2008

	Poisoning deaths
	Leading cause of injury
State	death
14 h 10 h	
United States	Poisoning
Alabama	Motor Vehicle Traffic
Alaska	Poisoning
Arizona	Poisoning
Arkansas	Motor Vehicle Traffic
California	Poisoning
Colorado	Poisoning
Connecticut	Poisoning
Delaware	Poisoning
District of Columbia	Firearm
Florida	Poisoning
Georgia§	§
Hawaii	Poisoning
Idaho	Motor Vehicle Traffic
Illinois	Poisoning
Indiana	Poisoning
lowa	Motor Vehicle Traffic
Kansas	Motor Vehicle Traffic
Kentucky	Poisoning
Louisiana	Motor Vehicle Traffic
Maine	Poisoning
Maryland	Poisoning
Massachusetts	Poisoning
Michigan	Poisoning
Minnesota	Poisoning
Mississippi	Motor Vehicle Traffic
Missouri	Motor Vehicle Traffic
Montana	Motor Vehicle Traffic
Nebraska	Motor Vehicle Traffic
Nevada	Poisoning
New Hampshire	Poisoning
New Jersey	Poisoning
New Mexico	Poisoning
New York	Poisoning
North Carolina	Motor Vehicle Traffic
North Dakota	Motor Vehicle Traffic
Ohio	Poisoning
Oklahoma	Motor Vehicle Traffic
Oregon	Poisoning
Pennsylvania	Poisoning
•	<u> </u>

Rhode Island	Poisoning
South Carolina	Motor Vehicle Traffic
South Dakota	Motor Vehicle Traffic
Tennessee	Motor Vehicle Traffic
Texas	Motor Vehicle Traffic
Utah	Poisoning
Vermont	Poisoning
Virginia	Motor Vehicle Traffic
Washington	Poisoning
West Virginia	Poisoning
Wisconsin	Poisoning
Wyoming	Motor Vehicle Traffic

NOTE: § The cause of death was inconclusive for a high percentage of deaths in Georgia at the close of the 2008 final mortality file. The manner of death was investigations, including poisoning deaths, are among causes that remain pending at the close of the file. Thus, the poisoning death rate for Georgia may not I X90, Y10-Y19, Y35.2, or *U01(.6-.7). Drug poisoning ICD-10 UCODs are: X40-X44, X60-X64, X85, Y10-Y14.

SOURCE: CDC/NCHS, National Vital Statistics System.

Data Brief 81: Drug Poisoning Deaths in the United States, 1980–2008

Data table for Figure 3. Number of drug poisoning deaths involving opioid analgesics and other drugs: United States, 1980–2008

Drugs involved in drug poisoning deaths

	Any opioid analgesic	Specified dr
		other than (
Year		
		Number o
1999	4,030	
2000	4,400	
2001	5,528	
2002	7,456	
2003	8,517	
2004	9,857	
2005	10,928	
2006	13,723	
2007	14,408	
2008	14,800	

NOTE: Drug categories are mutually exclusive

include natural and semi-synthetic opioid analgesics (e.g. morphine, oxycodone, hydrocodone) and synthetic opioid an may involve opioid analgesics.

Drug poisoning ICD—10 underlying cause of death codes are: X40- X44, X60-X64, X85, Y10-Y14. Among deaths with drug any opioid analgesic (any of the codes T40.2- T40.4); specified drug(s) other than opioid analgesic (any of the codes T30- T40.4).

SOURCE: CDC/NCHS, National Vital Statistics System.

Data Brief 81: Drug Poisoning Deaths in the United States, 1980–2008

Data table for Figure 4. Number of drug poisoning deaths involving opioid analgesic by opioid analgesic category: Unite

		Opioid analgesic involved in drug poisoning deaths	
I	Natural and semi-synthetic opioid analgesic		Year
Numbe			
	2,749		1999
	2,917		2000
	3,479		2001
	4,416		2002
	4,867		2003
	5,231		2004
	5,774		2005
	7,017		2006
	8,158		2007
	9,119		2008

NOTE: Opioid analgesic categories are not mutually exclusive. Deaths involving more than one opioid analgesic categor oxycodone and hydrocodone and synthetic opioid analgesics include fentanyl. Drug poisoning deaths ICD—10 underlyin underlying cause, the following ICD-10 codes indicate the type of drug(s) involved: natural and semi-synthetic opioid ar

SOURCE: CDC/NCHS, National Vital Statistics System.

Data Brief 81: Drug Poisoning Deaths in the United States, 1980–2008

Data table for Figure 5. Drug poisoning death rates by age: United States, 1999–2008

|--|

Age (years)

	Under 15		0.1	0.1
	15-24		3.2	3.7
	25-34		8.1	7.9
	35-44		14.0	14.3
	45-54		11.1	11.6
	55-64		4.2	4.2
65 and		over	2.7	2.4

NOTE: Drug poisoning ICD–10 underlying cause of death codes are X40-X44, X60-X64, X85, Y10-Y14. SOURCE: CDC/NCHS, National Vital Statistics System.

Ninth Revision of the ICD (ICD-9) in 1980–1998. Poisoning ICD–10 underlying cause of death codes (UCOD) are: X40-

are: X40-X44, X60-X64, X85, Y10-Y14 and ICD-9 UCODs are: E850-E858, E950.0-E950.5, E962.0, E980.0-E980.5. 2.1,V83-V86 (.0-.3),

as poisoning deaths.

	Age-adjusted rate per 100,000 population	Comparison to US age-adjusted poisoning rate	
Number	pe. 100,000 pepalation		oo age aajastea poisoning rate
41,080		13.4	
674		14.5	Similar
166		24.2	
951		15.0	_
410		14.9	•
4,334		11.6	Lower
900		17.5	Higher
438		12.2	Similar
141		16.3	Similar
66		10.8	Similar
3,266		18.0	Higher
1,049		§ 10.7	Difference not tested [§]
159		12.1	Similar
184		12.3	Similar
1,526		11.8	Lower
951		15.1	Higher
279		9.5	
259		9.4	Lower
860		20.2	8
695		16.1	•
199		14.9	
748		12.8	
867		12.9	
1,399		13.7	
518		9.6	
347		12.2	
867		14.8	•
164		17.3	<u> </u>
132		7.6	
556 153		21.0 11.2	•
829		9.4	
590		30.8	
1,910		9.5	
1,296 57		14.0 9.5	
1,924		16.7	•
644 538		18.1 13.8	=
2,031		16.4	
2,031		10.4	Higher

212		19.9	Higher
618		13.8	Similar
	72	9.6	Lower
1,039		16.4	Higher
2,248		9.4	Lower
526		20.8	Higher
	83	12.3	Similar
802		10.1	Lower
1,122		16.4	Higher
492		27.6	Higher
691		12.0	Lower
	98	18.6	Higher

399-2008

ug(s)		
bioiqc	Only non- analgesic	specified drug(s)

of deaths		
	0.353	2.566
	9,253	3,566
	9,073	3,942
	9,446	4,420
	10,774	5,288
	11,358	5,910
	11,314	6,253
	12,063	6,822
	12,738	7,964
	12,746	8,856
	12,408	9,242

e. Opioid analgesics

algesics (e.g. methadone, fentanyl). Some deaths in which the drug was poorly specified or unspecified g poisoning as the underlying cause, the following ICD-10 codes indicate the type of drug(s) involved: 6–T50.8 other than T40.2–T40.4); only nonspecified drug(s) (only T50.9).

d States, 1999-2008

Methadone	Synthetic opioid analgesic, excluding methadone
r of deaths	
784	730
986	782
1,456	957
2,358	1,295
2,972	1,400
3,845	1,664
4,460	1,742
5,406	2,707
5,518	2,213
4,924	2,306

y are counted multiple times. Natural and semi-synthetic opioid analgesics include morphine, ig cause of death codes are: X40-X44, X60-X64, X85, Y10-Y14. Among deaths with drug poisoning as the halgesic (T40.2); methadone (T40.3); synthetic opioid analgesic, excluding methadone (T40.4).

2001	2002	2003	2004	2005	2006	2007	2008

Death	s per	100,00	0 population				
0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2
4.3	5.2	6.0	6.6	6.9	8.2	8.4	8.2
8.6	10.3	11.3	11.7	13.3	15.7	16.4	16.5
15.5	18.0	18.8	19.1	19.4	21.5	21.2	20.9
13.1	16.1	18.0	19.3	21.1	24.1	25.1	25.3
4.7	6.0	7.0	7.9	9.1	10.6	12.4	13.0
2.6	3.0	3.0	3.0	3.3	3.5	3.8	4.1