

DEA TOX DRUG ENFORCEMENT ADMINISTRATION

TOXICOLOGY TESTING PROGRAM

2022 ANNUAL REPORT



U.S. Department of Justice Drug Enforcement Administration Diversion Control Division Drug and Chemical Evaluation Section

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Introduction

The Drug Enforcement Administration's Toxicology Testing Program (DEA TOX) began in May 2019 as a surveillance program aimed at detecting new psychoactive substances within the United States. In response to the ongoing synthetic drug epidemic, the Drug Enforcement Administration (DEA) awarded a contract with the University of California at San Francisco (UCSF) to analyze biological samples generated from overdose victims of synthetic drugs.

In many cases, it can be difficult to ascertain the specific substance responsible for the overdose. The goal of DEA TOX is to connect symptom causation to the abuse of newly emerging synthetic drugs (e.g. synthetic cannabinoids, synthetic cathinones, synthetic opioids, other hallucinogens, etc.).

DEA has reached out to local health departments, law enforcement partners, poison centers, drug court laboratories, hospitals, and other medical facilities to offer testing of leftover or previously collected samples for analysis of synthetic drugs. DEA TOX is interested in patients thought to have ingested a synthetic drug, where the traditional drug screen has produced little or no viable options to explain the symptoms exhibited by the patient (alcohol and delta-9 THC are exempted). DEA TOX may approve leftover unused biological samples (or biological samples) or occasionally non-biological samples for testing from a medical facility or law enforcement partner only.

Once DEA TOX is contacted (DEATOX@DEA.GOV) and upon approval by DEA of the request for testing of specific samples, the originating laboratory is invited to send their samples to the Clinical Toxicology and Environmental Biomonitoring (CTEB) Laboratory at UCSF. DEA covers the full cost of analysis for each sample approved for testing. Using liquid chromatography quadrupole time-of-flight mass spectrometry, synthetic drugs identified within the samples are confirmed and quantified. Levels denoted in the tables below with a defined range represent the low and high concentrations reported when the frequency of detection is greater than one. The CTEB laboratory currently maintains a comprehensive drug library consisting of the following:

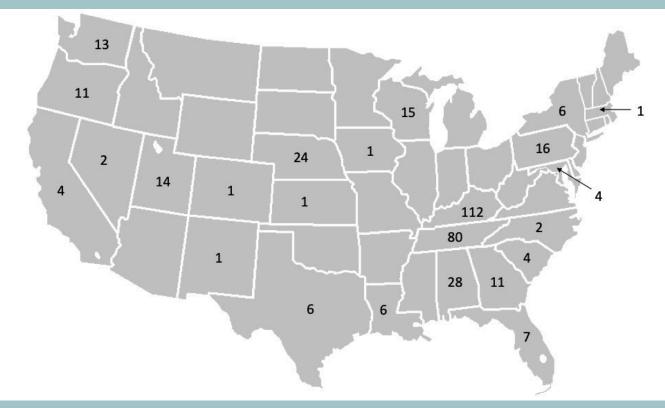
912 new psychoactive substances (NPS);
161 traditional illicit drugs (TID);
93 prescription or over-the-counter (OTC) drugs;
15 dietary supplement stimulants (DSS); and
Multiple precursor chemicals, additives or impurities (P/A/I)

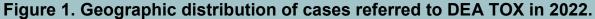
Each sample is analyzed by targeted and suspect screening through liquid chromatography- quadrupole time-of-flight mass spectrometry (LC-QTOF/MS) using non-targeted data acquisition. Confirmed drugs are quantified through targeted LC-QTOF/MS testing using the isotope dilution method (See Appendix 1 for method details).

This publication presents the results of cases analyzed and completed by the CTEB laboratory from January 1, 2022 through December 31, 2022.

Summary

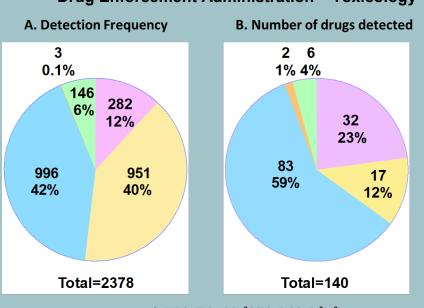
Between January 1, 2022 through December 31, 2022, 356 biological samples, 19 drug products, and 1 drug paraphernalia from 370 cases originating in 24 states (Fig.1) were submitted to DEA TOX. These samples were analyzed for NPS, TID, prescription or OTC drugs, DSS, and P/A/I. The biological samples submitted consisted of 41 serum, 38 plasma, 151 whole blood, 1 liver tissue, 1 muscle tissue and 124 urine samples. The drug products are further described on page 19.





DEA TOX identified and confirmed a total of 2,378 drugs and metabolites that consisted of 282 NPS detections, 951 TID detections, 996 prescription or OTC drug detections, 3 DSS, and 146 P/A/I detections during this reporting period (Fig. 2A). While some drugs identified could be placed in more than one category, for purposes of this report and for consistency, DEA TOX placed such substances in a single category only. Many prescription drugs that are commonly abused and encountered are listed as TID. Substances that are not approved by the Food and Drug Administration for medical use within the U.S. are considered NPS or adulterants.

A breakdown of the 2,378 total drug and metabolite confirmations demonstrated 140 different drugs, which consisted of 32 NPS, 17 TID, 83 prescription and OTC drugs, 2 DSS and 6 P/A/I (Fig. 2B).



Legend: NPS, TID, PD/OTC, DSS, P/A/I

Figure 2. Detection frequency and number of drugs detected in each drug category *Case Types*

The cases referred in 2022 can be classified into four categories: deaths (147), overdoses (186), urinalysis (19), and drug products (18). One case submitted only drug paraphernalia. The general drug classes found in overall, overdose, and death cases are presented in Figure 3. Overall, NPS accounted for 41% (153 of 370) of the cases. This increased to 67% (98 of 147) in death cases. In cases where only TID's were found, fentanyl was the predominant substance identified, especially in non-fatal submissions.

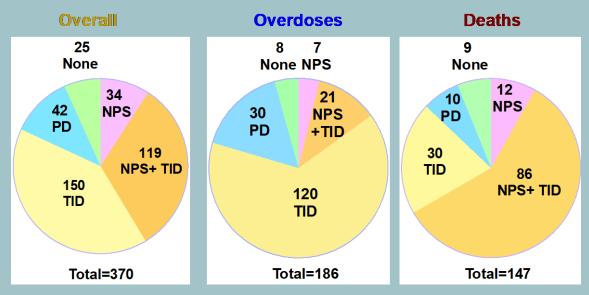


Figure 3. Drug class distribution in all, overdose and death cases

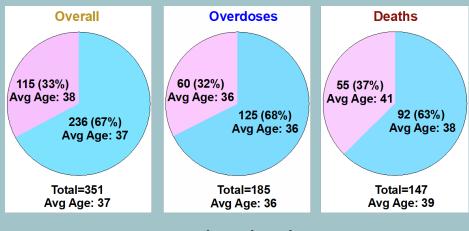
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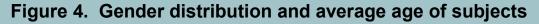
Drug Enforcement Administration – Toxicology Testing Program

Drug Enforcement Administration – Toxicology Testing Program Gender and Age of Subjects

Information on gender and/or age was available for 352 cases, of which 236 are males and 115 females (one case was not identified) (Fig. 4). The average age for males is 37 (range: 1.3-67) while the average for females is 38 (range: 1.9-70). The overall average age is 37 (range: 1.3-70).



Legend: Female, Male



Polydrug use

The number of NPS and TID detected were tallied for each of the 352 cases with known gender and/or age. Two or more substances were confirmed in 219 (62%) cases. The highest number found in death cases was nine, while the highest number in overdose cases was seven. The distribution of number of drugs detected in the cases is summarized in Fig 5.

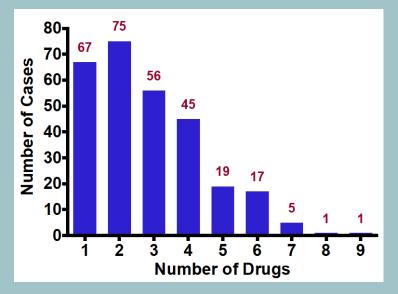


Figure 5. Distribution of the number of NPS and TID in cases

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Drug Enforcement Administration – Toxicology Testing Program New Psychoactive Substances

DEA TOX confirmed 282 detections comprising of 32 NPS[§] from seven different classes of drugs (Fig. 6) in 2022. The total encounters for each NPS class are summarized in Figure 7. The quantitative levels found in the cases are presented for each NPS pharmacological class in Tables 1-4. Phenibut is outside these classes. There were 10 cases confirmed from Alabama and one each from Maryland and Washington. The levels quantified were 92,000 (serum) and 28.4 (plasma) ng/mL. No quantitation was requested in the 10 urine samples referred.

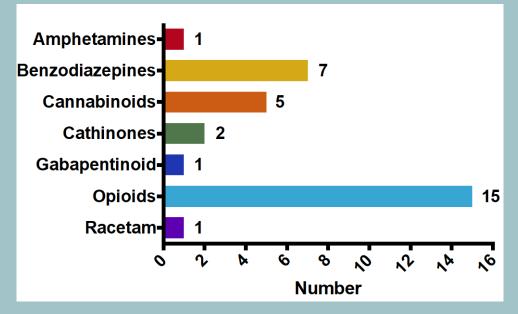


Figure 6. Number of drugs in each NPS class detected in cases

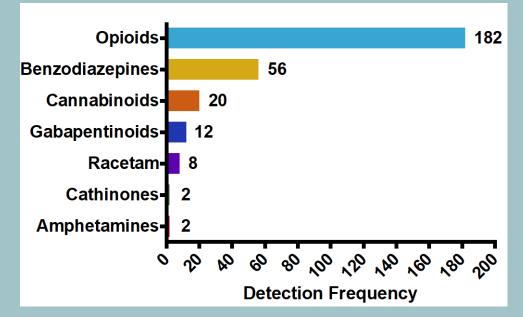


Figure 7. Detection frequency of each NPS class detected in cases

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Drug Enforcement Administration – Toxicology Testing Program Table 1. NPS stimulants detected in 2022

	Drug	Erog	States	Confir	med Le	vels (ng/	'mL)**
Drug Class	Drug	Freq.	Freq. Found*	S	Р	WB	U
Amphetamine	MDEA	2	GA			0.8- 1.2	
Cathinanaa	Eutylone	1	FL				
Cathinones	Pentylone	1	TN			1.1	
Racetam	Phenylpiracetam	8	AL(6), TN, WA	36400			

Table 2. NPS cannabinoids detected in 2022

Drug	Erog	States	Confi	rmed Le	vels (ng/	mL)**
Diug	Drug Freq. Found*		S	Ρ	WB	U
4F-ABUTINACA	1	WA				
4F-MDMB-BUTICA	5	FL				
11-nor-9-carboxy-delta-8-THC	4	KY(3), TX	46.9	178		107- 3230
ADB-BUTINACA	6	FL(5), UT(1)		6.1		
Delta-8-THC	3	KY(3)	5.7			128
Delta-10-THC	1	KY				

Table 3. NPS benzodiazepines detected in 2022

Drug	Eroa	States Found*	Confir	ned Le	vels (ng/	mL)**
Drug	Freq.	Freq. States Found*		Р	WB	U
8-Aminoclonazolam	20	GA(3), KS, KY, LA, MD, NE, TN(11), UT	0.4- 7.4	0.4	0.4- 47.2	471
Bromazolam	21	GA(2), KY(2), NM, TN(15), UT,	17.5- 46.6	3	0.1- 62.9	
Clonazolam	1	UT		1.2		
Deschloroetizolam	1	KY			6.8	
Etizolam	3	IA, TN, WA			0.7- 18.5	
Flualprazolam	4	GA(2), TN, UT	1.8- 3.3	2.9	5.1	
Flubromazepam	3	TN(3)			1.1- 20.7	
Flubromazolam	3	IA, WA(2)			0.7	

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Drug Enforcement Administration – Toxicology Testing Program Table 4. NPS opioids detected in 2022

Dimen	F		Confir	med Le	evels (ng	/mL)**
Drug	Freq.	States Found*	S	Ρ	WB	U
2-Methyl-AP-237	2	KS, WA			313- 379	
5-Amino Isotonitazene	1	TN			1.2	
7-OH Mitragynine	5	GA, KY(2), LA, TN	15.1		2.9	1370- 1960
Acetyl Fentanyl	10	KY, WA(9)				
Despropionyl <i>para-</i> Fluorofentanyl	36	AL, CA, GA(3), KY(3), NE(4), TN(20), WA(3), WI	0.6		0.1- 3.3	62.5- 276
Isotonitazene	1	TN			0.9	
Metonitazene	15	KY, TN(14)			1.1- 13.8	
Mitragynine	16	GA(2), KS, KY(4), LA(2), PA, TN(4), UT, WA	15.3- 139	4.4	0.7- 113	12.1- 1070
N-Methyl Norfentanyl	2	KY, WA				98
<i>N</i> -Piperidinyl Etonitazene	3	TN(3)			1.6-12	
<i>N</i> -Pyrrolidino Etonitazene	1	LA			1.1	
<i>para</i> -Bromofentanyl	1	WA				
para-Chlorofentanyl	1	WA				
<i>para</i> -Fluoro acetylfentanyl	3	CA, TN, WA			0.4- 1.0	
<i>para</i> -Fluorofentanyl	74	AL, CA(3), GA(6), KY(7), NE(7), TN(44), WA(4), WI(2)	1.2- 17.7		0.2- 111	1.9- 1120
Protonitazene	1	TN				4
Remifentanil Acid	3	KY, LA(2)			3.4- 7.5	353
Tianeptine	7	AL, TN			9.9	135- 7880

* AL – Alabama; CA – California; GA – Georgia; KS – Kansas; KY – Kentucky; LA – Louisiana;
 NE – Nebraska, PA – Pennsylvania; TN – Tennessee; UT – Utah; WA – Washington; WI – Wisconsin;
 **S – Serum; P – Plasma; WB – Whole Blood; U – Urine

§ - Parent drugs or metabolites are only counted once for the number of drugs detected in Tables 1-4. If only a metabolite is encountered in the absence of a parent drug, it will still be counted as a unique drug. Both parent drugs and metabolites are counted as detections.

Drug Enforcement Administration – Toxicology Testing Program Traditional Illicit Drugs

DEA TOX confirmed 951 detections comprising of 17 TIDs[§] (Table 5) in 2022.

Table 5. TID detected in 2022

	Drug	rug Erog States		Confir	med Le	vels (ng/	'mL)**
Drug Class	Drug	Freq.	Freq. Found*		Ρ	WB	U
	4-Hydoxymeth- amphetamine	19	KY(16), NE, TN(2)			1.3- 19.8	12.1- 8075
	Amphetamine	46	GA(2), KY(16), NE(6), NM, OR, TN(16), TX, UT(2), WI	13.7- 978	45.6	4.3- 435	722- 15400
	HMMA [‡]	3	KY				3.5- 835
Amphetamines	MDA	1	KY			14.5	
(4)	MDMA	2	KY, UT		88.6	660	
	Methamphetamine	122	AL(3), CA, GA(4), KY(45), NC, NE(9), NM, OR(3), PA(3), SC, TN(39), TX, UT(7), WA(2), WI(2)	2.5- 5940	7.9- 1430	1.1- 7870	1.5- 204000
Arylcyclohexyl- amines (1)	Ketamine	18	GA, KY(15), PA, WI	9.1- 398			99.6- 50700
Cannabinoids (2)	11-nor-9- carboxy-delta-9- THC	44	CO, FL, GA, IA, KS, KY(21), MA, NE, OR(3), PA(3), SC, TN(2), TX(2), UT(3), WI(2)	60.6- 232	35.5- 342	37.6- 141	60.6- 38800
	11-OH-delta-9- THC	1	KY				685
	Cannabidiol	1	KY				NQ

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Drug Enforcement Administration – Toxicology Testing Progra							
Drug Class	Drug	Freq.	States	Confir	med Le	vels (ng/	mL)**
Brug Glass	Brug	TTCq.	Found*	S	Р	WB	U
	Delta-9-THC	4	KS, TN, UT(2)		23	34.0- 56.8	99.5
	Benzoylecgonine	99	AL(2), FL(2), GA(8), KY(32), MA, NE(13), OR(3), PA(12), TN(17), UT(2), WI(7)	12.3- 1700	0.9- 998	1.5- 1880	4.0- 129000
	Cocaethylene	27	AL, FL(2), GA(4), KY(7), NE(8), TN(4), WI	NQ		NQ	NQ
Cocaine (1)	Cocaine	59	AL, FL(2), GA(5), KY(14), MD, NE(10), PA(6), TN(13), WA(2), WI(5)	0.4- 94.2	1.0- 30.8	0.2- 996	2.4- 257000
	Ecgonine methyl ester	62	AL(2), FL(2), GA(6), KY(19), MD, NE(6), OR(3), PA(12), TN(6), WI(5)	NQ	NQ	NQ	NQ
	6-Acetyl Morphine	1	WI				261
Opioids (9)	Beta- Hydroxyfentanyl	41	CO, FL, KY(11), MD, NE(4), NM, TN(19), WI(3)		0.3- 1.4	0.5-2.9	4.3- 410
	Codeine	6	TN(5), UT			0.2-2.3	
	Desmethyl-cis- Tramadol	7	KS, KY(4), TN(2)			0.5- 49.9	5.2- 58.3

	Drug Enfor	cement /	Administration	· · · · · · · · · · · · · · · · · · ·			
Drug Class	Drug	Freq.	States		-	vels (ng/	-
		•	Found*	S	Р	WB	U
	Fentanyl	174	AL, CO, GA(8), FL(2), IA, KY(44), MA, MD, NC, NE(13), NM, NV(2), OR(3), PA(11), SC, TN(59), TX(4), UT(3), WA(11), WI(6)	0.8- 104	0.8- 71.0	0.5- 106	4.9- 6160
	Hydrocodone	10	KY(4), TN(6)			0.2- 19.2	51.2- 177
	Hydromorphone	13	KY(7), TN(6)	20.6		1.8- 2270	19.2- 247
	Morphine	31	IA, KY, NE(2), TN(24), UT(2), WI		7.3	0.5- 113	1520 - 8160
	Norfentanyl	122	AL, CO, FL(2), GA, IA, KY(43), MA, MD, NE(13), NM, OR(3), PA(11), SC, TN(31), TX, UT, WA(3), WI(6)	0.1- 9.6	0.3- 10.5	0.3- 20.6	1.7- 42800
	Oxycodone	22	KY(6), LA, NE, OR, TN(9), UT ,WI(3)		3.4- 20.2	0.3- 281	14.7- 157000
	Oxymorphone	1	KY				NQ
	Tramadol	15	KY(6), TN(6), UT, WA(2)		3.4	0.2- 1020	7.0- 80.8

*AL – Alabama; CA – California; CO – Colorado; FL – Florida; GA – Georgia; IA – Iowa; KS – Kansas; KY – Kentucky; LA – Louisiana; MA – Massachusetts; MD – Maryland; NC – North Carolina; NE – Nebraska; NM – New Mexico; NV – Nevada; OR – Oregon; PA – Pennsylvania; SC – South Carolina; TN – Tennessee; TX – Texas; UT – Utah; WA – Washington; WI – Wisconsin

**S – Serum; P – Plasma; WB – Whole Blood; U – Urine; NQ – Not Quantified [‡] HHMA (3,4-Dihyroxymethamphetamine is a metabolite of MDMA)

§ - Parent drugs or metabolites are only counted once for the number of drugs detected in Table 5. If only a metabolite is encountered in the absence of a parent drug, it will still be counted as a unique drug. Both parent drugs and metabolites are counted as detections.

Prescription and Over the Counter Drugs

DEA TOX confirmed 996 detections comprising of 83 prescription or OTC drugs[§] (Table 6) in 2022. Drugs for the prescription/OTC drugs panel are not typically quantitated unless specifically requested thus "Confirmed Levels" are not provided.

Table 6. Prescription or OTC drugs detected in 2022

Drug Class	Drug	Freq.	States Found*
Anesthetic	Lidocaine	35	AL, CA(2), GA, KY(14), MD, NC, NE(3), PA(2), SC, TN(7), UT, WA
Antibiotio	Linezolid	2	GA, TN
Antibiotic	Sulfomethoxazole	3	KY, TN(2)
	Carbamazepine	3	KY(2), TN
	Gabapentin	54	AL, GA, KY(15), LA, NE(5), OR, TN(29), WI
Anticonvulsant	Lamotrigine	9	GA, KY(6), TN, UT
	Levetiracetam	7	GA, KY(3), TN(3)
	Oxcarbazepine	5	KY(3), TN(2)
	Topiramate	1	NE
	Amitriptyline	7	KY(6), TN
	Citalopram	19	KY(6), NC, NE(3), OR, TN(4), TX, UT(2), WA
	Doxepin	1	TN
	Duloxetine	4	KY, OH, TN(2)
	Fluoxetine	16	AL, KY(10), LA, NE, TN(2), WA
	mCPP [‡]	18	KS, KY(8), LA, NE, TN(5), UT, WI
	Nordoxepin	1	КҮ
Antidepressant	Norfluoxetine	14	KS, KY(7), LA, NE, NY, TN(2), WA
	Nortriptyline	7	KY(6), TN
	Paroxetine	4	AL, TN(2), WI
	Pipradol	1	WI
	Protriptyline	2	KY(2)
	Sertraline	11	GA, KY(3), OR, NE, TN(4), WI
	Trazodone	23	KS, KY(8), LA, NE(2), TN(9), UT, WI
	Venlafaxine	4	KY(3), UT
Antidiabetic	Metformin	4	KY(2), OR, TN

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	Drug Enforcement Administration – Toxicology Testing Progra							
Drug Class	Drug	Freq.	States Found*					
Antidiarrheal	Loperamide	6	GA, KY, TN(2), WI(2)					
	Chlorpheniramine	6	AL, KY(2), NE, TN(2)					
	Cimetidine	2	KY, WA					
	Diphenhydramine	75	GA, KS, KY(25), NC, NE(2), NY, OR(2), PA, TN(37), TX, UT(2), WI					
Antihistamine	Doxylamine	5	AL, NE, TN(2), WI					
	Hydroxyzine	21	CA, GA, IA, KY(7), NE(2), OR, TN(3), UT, WI(4)					
	Promethazine	6	KY(2), NE, TN(2), WI					
	Pseudoephedrine	2	CA, NE					
	Aripiprazole	5	GA(2), TN(3)					
	Clozapine	1	КҮ					
Antipsychotic	Haloperidol	5	OR, TN(2), UT(2)					
	Olanzapine	8	GA, KY(4), OR, UT(2)					
	Ziprasidone	3	OR, TN(2), UT(2)					
A antine tree vised	Emtricitabine	1	WI					
Antiretroviral	Tenofovir	1	WI					
Anxiolytic	Buspirone	6	NE, TN(4), WI					
	7-amino Clonazepam	17	GA(2), KS, KY(6), LA, NE, PA, TN(2), UT, WI(2)					
	7-amino Nitrazepam	1	GA					
	Alpha-hydroxy Alprazolam	11	IA(2), KY(3), NE, TN(2), UT, WI(2)					
	Alprazolam	23	AL, GA, IA. KY(2), NE(3), NY, TN(11), UT, WI(2)					
	Chlordiazepate	2	KY, WI					
	Clobazam	3	KY (3)					
Benzodiazepine	Clonazepam	5	KY(2), NE, LA, WI					
	Diazepam	15	GA, KY(3), LA, NE(2), TN(6), TX, UT					
	Lorazepam	28	CO, GA, KY(18), OR(2), TN(3), UT(2), WI					
	Midazolam	49	AL, CO, FL, GA, KY(32), LA, NC, NE, OR(3), PA, TN(2), UT(2), WA, WI					
	Mirtazapine	3	AL, KY, PA					
	Nitrazepam	1	КҮ					
	Nordiazepam	19	GA(2), KY(5), LA, NE(2), TN(6), TX, UT, WI					

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Drug Enforcement Administration – Toxicology Testing Program						
Drug Class	Drug	Freq.	States Found*			
-	Oxazepam	11	KY(7), NC, NE, TN, WI			
	Temazepam	8	KY(2), NC, NE, TN(2), TX, WI			
	Zolpidem	1	КҮ			
	Amiodarone	14	AL, GA, KY(3), LA, NE(2), TN(4), TX, WA			
	Atenolol	4	CA, GA, KY(2)			
	Atorvastatin	5	IA, KY, LA, TN(2)			
	Atropine	12	AL, KY(6), NE, PA, TN(2), TX			
	Carvedilol	4	KY(2), LA, NE			
	Clonidine	7	KY(3), NE, TN(3)			
	Diltiazem	3	AL, TN, WI			
	Furosemide	2	KY(2)			
	Labetalol	2	KY, SC			
Cardiovascular	Lisinopril	7	IA, KY(5), TN			
Cardiovascular	Metoprolol	10	IA,KY, TN(6), WI(2)			
	Propanolol	2	KS, TN			
	Verapamil	1	WI			
	Warfarin	2	NE, WI			
Cough	Dextromethorphan	19	AL, GA, KY(8), OR, NE(3), PA(2), TN(3)			
Suppressant	Dextrorphan	15	AL, KY(8), NE(2), PA, TN(3)			
	Norpseudoephedrine	5	KY(3), TN(2)			
Decongestant	Phenylephrine	1	КҮ			
	Pseudoephedrine	9	KY(8), TN			
Diuretic	Furosemide	4	KY(3), WI			
	Baclofen	7	IA, KY(5), UT			
	Cyclobenzaprine	9	KY(5), TN(3), WI			
Muscle Relaxant	Methocarbamol	1	TN			
	Orphenadrine	1	NE			
	Buprenorphine	16	KY(10), OR, TN(2), UT, WI(2)			
	EDPP	9	GA(2), KY, NE(5), TN			
	Methadone	12	GA(2), KY(3), NE(5), TN(2)			
Opioid	Naloxone (Antagonist)	98	AL, FL(2), GA(3), KY(39), LA, MD, NE(12), PA(7), TN(23), TX(4), WI(5)			
	Naltrexone (Antagonist)	1	КҮ			
	Norbuprenorphine	10	KY(5), OR, TN, WI(3)			

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Drug Enforcement Administration – Toxicology Testing Program								
Drug Class	Drug	Freq.	States Found*					
Pain Reliever	Acetaminophen	86	FL(2), GA(3), AL(2), CO, KY(46), LA, NE, NM, OR, PA(3), TN(14), TX, UT(3), WA(6), WI					
	Naproxen	4	FL, KY, NE(2)					
Respiratory	Albuterol	3	KY(3)					
Stimulant	Methylphenidate	1	TN					
Tuberculostatic	Levofloxacin	5	GA, KY, TN(3)					

*AL – Alabama; CA – California; CO – Colorado; FL – Florida; GA – Georgia; IA – Iowa; KS – Kansas; KY – Kentucky; LA – Louisiana; MA – Massachusetts; MD – Maryland; NC – North Carolina; NE – Nebraska; NM – New Mexico; NV – Nevada; OR – Oregon; PA – Pennsylvania; SC – South Carolina; TN – Tennessee; TX – Texas; UT – Utah; WA – Washington; WI – Wisconsin

**S – Serum; P – Plasma; WB – Whole Blood; U – Urine; NQ – Not Quantified

^{‡-} - The mcPP (meta-Chlorophenylpiperazine) detected in these cases is the metabolite of trazodone.

§ - Parent drugs or metabolites are only counted once for the number of drugs detected in Table 6. If only a metabolite is encountered in the absence of a parent drug, it will still be counted as a unique drug. Both parent drugs and metabolites are counted as detections.

Dietary Supplement Stimulants

DEA TOX confirmed three detections comprising of two DSS (Table 7) in 2022.

Table 7. DSS detected in 2022

Drug	Erog	States Found*	Confirmed Levels (ng/mL)*		mL)**	
Drug	Freq.	States Found	S	Р	WB	U
Hordenine	2	WI(2)				NQ
Yohimbine	1	LA			NQ	

*LA – Louisiana; WI – Wisconsin

**S – Serum; P – Plasma; WB – Whole Blood; U – Urine; NQ- Not Quantified

Precursors/Additives/Impurities

DEA TOX confirmed 146 detections comprising of six P/A/I[§] (Table 8) in 2022.

Table 8. P/A/I detected in 2022

Drug	Drug	Freq.	req. States		onfirme (ng/n	ed Level nL)**	S
Class		Found*		s	Р	WB	U
	Brodifacoum	4	FL				
	Phenacetin	3	KY, PA(2)		6.1- 94.9		
Adulterant (4)	Quinine	16	GA(2), KY(6), NE(3), TN(3), TX, WI	7.5- 13.5	1.5- 2.1	0.6- 25.4	111- 846
	Xylazine	22	AL, KY(6), NE, TN(14)			0.1- 35.4	15.4- 139
Impurity	N,N-dimethyl- amphetamine	17	GA(2), KY(5), NE(5), NM, TN(4)	146- 546		1.8- 26.7	4.7- 9570
Precursor	4-ANPP	84	GA(3), IA, KY(13), MD, NE(5), NM, OR, PA(2), TN(44), WA(11), WI(2)	1.1-8.9	0.9- 2.7	0.2- 164	3.9- 104

*AL – Alabama; FL – Florida; GA – Georgia; IA – Iowa; KY – Kentucky; MD – Maryland; NE – Nebraska; NM – New Mexico; OR – Oregon; PA – Pennsylvania; TN – Tennessee; TX – Texas; WA – Washington; WI – Wisconsin

**S – Serum; P – Plasma; WB – Whole Blood; U – Urine; NQ – Not Quantified § - Parent drugs or metabolites are only counted once for the number of drugs detected in Table 8. If only a metabolite is encountered in the absence of a parent drug, it will still be counted as a unique drug. Both parent drugs and metabolites are counted as detections.

Drug Paraphernalia

DEA TOX received 20 exhibits and confirmed 91 detections in 19 drug products and one drug paraphernalia in 2022 (Table 9).

Table 9. Drug Paraphernalia exhibits analysis in 2022

Product (Weight)	Drug	Drug Class	State	Confirmed Level: mg drug/ g drug product	Actual Amount within Drug Product
Green herbal mixture	4F-MDMB-BUTICA Brodifacoum ADB-BUTINACA	Synthetic cannabinoid Additive Synthetic cannabinoid	FL	7.9 1.4 0.38	Bulk material provided
Green herbal mixture	4F-MDMB-BUTICA Brodifacoum ADB-BUTINACA	Synthetic cannabinoid Additive Synthetic cannabinoid Synthetic	FL	3.9 1.7 0.32 0.14	Bulk material provided
Green herbal mixture	Eutylone 4F-MDMB-BUTICA ADB-BUTINACA	cathinone Synthetic cannabinoid Synthetic cannabinoid	FL	6.1 0.21	Bulk material provided
Green herbal mixture	Brodifacoum 4F-MDMB-BUTICA ADB-BUTINACA	Additive Synthetic cannabinoid Synthetic cannabinoid	FL	6.3 0.58 0.061	Bulk material provided
Green herbal mixture	4F-MDMB-BUTICA ADB-BUTINACA Brodifacoum	Synthetic cannabinoid Synthetic cannabinoid Additive	FL	3.1 1.1 0.19	Bulk material provided
Gray crystal (188.2 mg)	Fentanyl Cocaine <i>para</i> -Fluorofentanyl 4-ANPP Diphenhydramine Norfentanyl Methamphetamine Phenacetin Despropionyl- <i>para</i> - fluorofentanyl	Opioid Stimulant Opioid Precursor Antihistamine Precursor Stimulant Additive Precursor	KY	26.0 15.0 11 5.2 2.1 0.80 0.098 0.075 0.048	4.9 mg 2.8 mg 2.1 mg 0.98 mg 0.40 mg 0.15 mg 0.018 mg 0.014 mg 0.009 mg

	Lidocaine	Anesthetic		0.043	0.0081 mg
	Acetyl Fentanyl	Opioid		0.021	0.004 mg
	Tramadol	Opioid	-	0.021	0.004 mg
Blue	Acetaminophen	Pain Reliever		533	55.0 mg
"M30"	Fentanyl	Opioid	•	18	1.9 mg
Tablet	4-ANPP	Precursor	WA	11	2.1 mg
(103.2 mg)	Acetyl Fentanyl	Opioid		0.015	0.0015 mg
Dhua	Acetaminophen	Pain Reliever		966	108 mg
Blue "M30"	Fentanyl	Opioid		35	3.9 mg
Tablet	4-ANPP	Precursor	WA	3.2	0.36 mg
(112.0	<i>N</i> -Methyl Norfentanyl	Opioid	VVA	0.015	0.0017mg
mg)	Acetyl Fentanyl	Opioid		0.014	0.0016 mg
Green "S 90 3" Tablet	Flubromazolam	Benzodiazepine	WA	40	0.89 mg
White	4-ANPP	Precursor		49	11 mg
Rock	Fentanyl	Opioid	WA	46	11 mg
(230.0	Acetyl Fentanyl	Opioid	VVA	0.28	0.064 mg
mg)	Norfentanyl	Precursor		0.11	0.025 mg
	Fentanyl	Opioid		51	3.8 mg
White	4-ANPP	Precursor		8	0.6 mg
Powder	Acetyl Fentanyl	Opioid	WA	0.14	0.010 mg
(73.7 mg)	Cocaine	Cocaine		0.030	0.0022mg
	Norfentanyl	Precursor		0.029	0.0021 mg
Blue	Acetaminophen	Pain Reliever		390	42 mg
"M30"	Fentanyl	Opioid		14	1.5 mg
Tablet	4-ANPP	Precursor	WA	3.9	0.42 mg
(107.6 mg)	Acetyl Fentanyl	Opioid		0.013	0.0014 mg
Blue	Acetaminophen	Pain Reliever		480	54 mg
"M30"	Fentanyl	Opioid		21	2.4 mg
Tablet	4-ANPP	Precursor	WA	3.1	0.35 mg
(114.1 mg)	Acetyl Fentanyl	Opioid		0.015	0.0017 mg
White	Fentanyl	Opioid		99	5.9 mg
Powder	4-ANPP	Precursor	WA	13	0.78 mg
(60.0 mg)	Methamphetamine	Amphetamine	v v /	7.8	0.47 mg
(00.0 mg)	Acetyl Fentanyl	Opioid		0.26	0.016 mg
	Acetaminophen	Pain Reliever		75	23 mg
Blue	Fentanyl	Opioid		43	13 mg
Powder	Methamphetamine	Amphetamine	WA	19	5.8 mg
(303.1 mg)	4F-ABUTINACA	Synthetic Cannabinoid	vvA	3.7	1.1 mg
5,	Etizolam	Benzodiazepine		1.7	0.52 mg

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	Drug En	forcement Admir	nistratio	on – Toxicology T	esting Program
	4-ANPP	Precursor		0.69	0.21 mg
	para-Fluorofentanyl	Opioid		0.66	0.20 mg
	Flubromazolam	Benzodiazepine		0.24	0.073 mg
	Lidocaine	Anesthetic		0.16	0.048 mg
	Tramadol	Opioid		0.076	0.023 mg
	Acetyl Fentanyl	Opioid		0.061	0.018 mg
	Cocaine	Cocaine		0.038	0.012 mg
	Despropionyl- <i>para-</i> Fluorofentanyl	Precursor		0.007	0.002 mg
Blue	Acetaminophen	Pain Reliever		430	46 mg
"M30"	Fentanyl	Opioid		16	1.7 mg
Tablet	4-ANPP	Precursor	WA	4.3	0.46 mg
(106.7 mg)	<i>para</i> -Fluorofentanyl	Opioid		0.053	0.0057 mg
	<i>para</i> -Fluorofentanyl	New Synthetic Opioid		12	1.2 mg
	Fentanyl	Opioid		3.0	0.29 mg
White Powder	Despropionyl- <i>para</i> - Fluorofentanyl	Precursor	WA	2.1	0.20 mg
(97.0 mg)	<i>para-</i> Fluoroacetylfentanyl	Opioid		0.11	0.011 mg
	4-ANPP	Precursor		0.078	0.0076 mg
	para-Chlorofentanyl	Opioid		0.038	0.0037 mg
	Fentanyl	Opioid		151	14.3 mg
	4-ANPP	Precursor		18	1.7 mg
	<i>para</i> -Fluorofentanyl	Opioid		2.1	0.20 mg
White	Tramadol	Opioid		0.97	0.092 mg
Rock	Norfentanyl	Opioid	WA	0.17	0.016 mg
(94.9 mg)	Despropionyl- <i>para-</i> Fluorofentanyl	Precursor		0.16	0.015 mg
	Acetyl Fentanyl	Opioid		0.10	0.0095 mg
	<i>para</i> -Bromofentanyl	Opioid		0.030	0.0028 mg
Vape	Delta-8 THC	Cannabinoid		600	155 mg
Liquid (259.1 mg)	Delta-10 THC	Cannabinoid	KY	212	55 mg

Drug Enforcement Administration – Toxicology Testing Program Selected Drug Trends in Death and Overdose Cases

Fentanyl

DEA TOX confirmed fentanyl in 184 samples from 174 cases (47%) of total cases) submitted by 20 states (Fig. 8). The samples consisted of 87 whole blood, 21 plasma, 20 serum, 44 urine and 12 drug products.

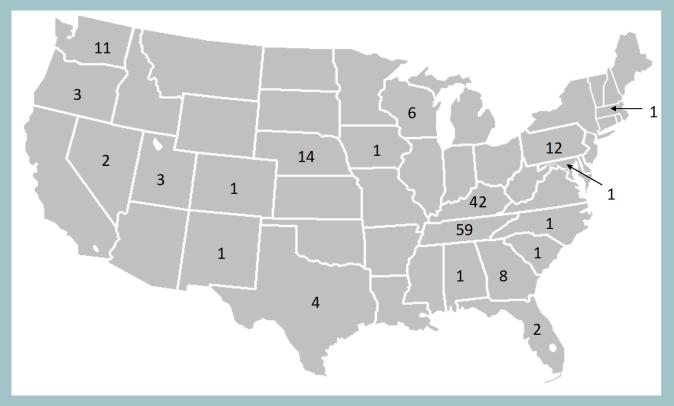


Figure 8. Geographic distribution of cases with confirmed fentanyl

There were 87 deaths and 76 overdoses. The gender distribution and average age in these cases are presented in Figure 9A. There were 87 and 37 blood samples in death and overdose cases, respectively. The mean fentanyl concentration in death cases (17.6 ng/mL) is 2.35x higher than in overdose cases (7.5 ng/mL) but significant overlap in concentrations was observed (Fig. 9B).

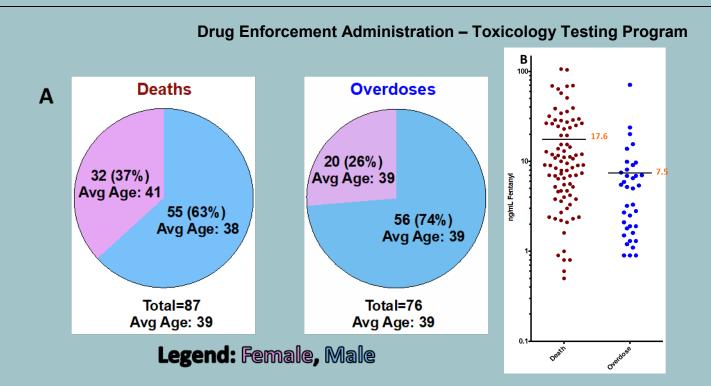


Figure 9. Gender and age (A) and blood fentanyl concentration (B) distribution in death and overdose cases

The concentration ranges for fentanyl found in blood samples from various states in death and overdose cases are presented in Table 10.

State	Death Case	S	Overdose Cases	
	Concentration	Frequency	Concentration	Frequency
	(ng/mL)*		(ng/mL)*	
Alabama	10	1		
Colorado			1.9	1
Florida	5.2-69.5	2		
Georgia	0.8- 104 (Avg = 16.6)	8		
lowa	6.5	1		
Kentucky	11.1	1	0.9-8.1 (Avg = 3.8)	11
Massachusetts			0.9	1
Maryland			20.1	1
Nebraska	0.9- 106 (Avg = 21.5)	13	5	1
Nevada			1.2-1.3	2
New Mexico	26.2	1		
North Carolina			7.5	1
Oregon			1.8-2.8 (Avg = 2.2)	3
Pennsylvania			0.5-71 (Avg = 14.6)	11
South Carolina			2.7	1
Tennessee	0.5- 69 (Avg = 16.7)	59		
Texas	0.8	1	1.1-5.9	2
Utah			1.6- 5.2	2

* Average is given in parenthesis for detection frequency ≥ 3

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Polydrug use is prevalent in cases involving fentanyl (Fig. 10). Of the 174 cases, 163 were biological samples; 91% of these cases (149 of 163) have two or more drugs confirmed.

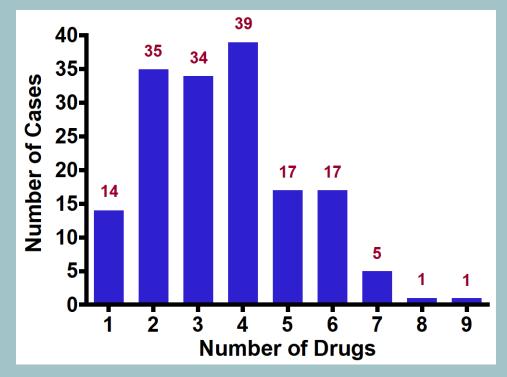


Figure 10. Distribution of the number of NPS and TID in fentanyl cases

In 163 cases, biological samples were analyzed. Fentanyl was detected along with other NPS and TID in these cases. Drugs that co-occurred with fentanyl in more than 10 cases are summarized in Table 11.

Drug	Co-occurrence Frequency		
	Overall	Deaths	Overdoses
Methamphetamine	76	39	37
Cocaine/ Benzoylecgonine	61	26	35
para-Fluorofentanyl	54	47	7
Metonitazene	12	11	1
Morphine	26	24	2
Oxycodone	11	9	2
Bromazolam	19	17	2
Clonazolam	16	14	2
Delta-9 THC	13	5	8
Xylazine	22	15	7
Quinine	11	5	6

Table 11. Co-occurrence freq	quency of other drug	us and adulterants in fenta	nvl cases

The distribution between overdose and death cases in some of these co-occurrences and the average concentration of co-occurring drug and fentanyl in death cases are presented in Fig. 11.

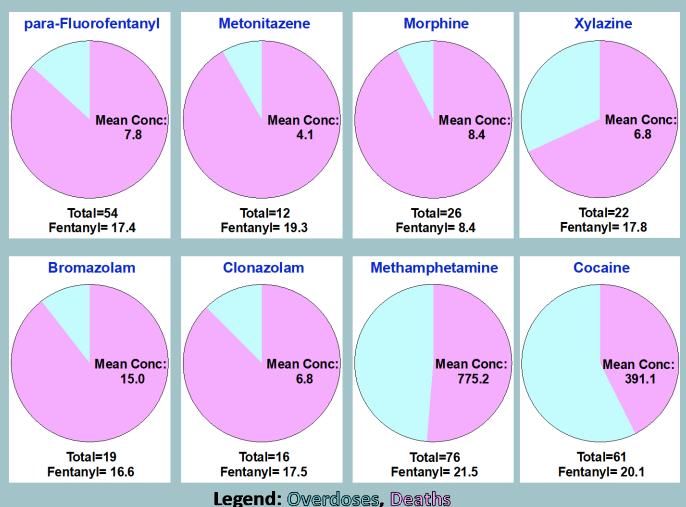


Figure 11. Distribution between deaths and overdoses in co-occurrence of fentanyl with selected drugs along with the average concentration in ng/mL of the co-occurring drug (in pie chart) and fentanyl in death cases

Drug Enforcement Administration – Toxicology Testing Program para-Fluorofentanyl

DEA TOX confirmed *para*-fluorofentanyl in 74 samples from 74 cases (20% of total cases) submitted by 8 states (Fig. 12). The samples consisted of 58 whole blood, 4 serum, 7 urine and 5 drug products.

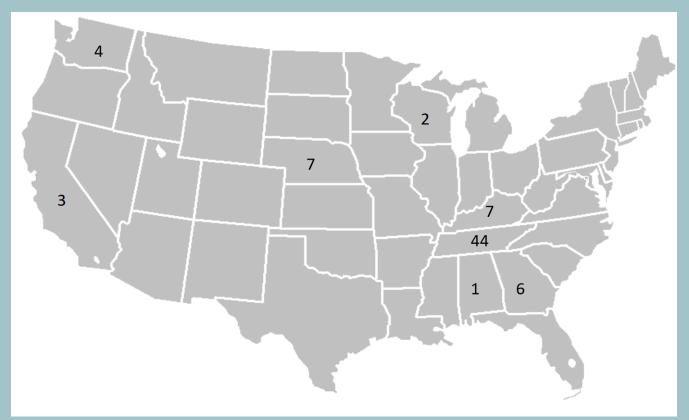


Figure 12. Geographic distribution of cases with confirmed para-fluorofentanyl

There were 60 deaths and 10 overdoses. The gender distribution and average age in these cases are presented in Figure 13A. 60 samples were submitted in death cases, while two samples were submitted in overdose cases. The distribution in blood *para*-fluorofentanyl concentrations observed in death cases is shown in Fig. 13B. The mean blood concentration in the death cases is 10.4 ng/mL.

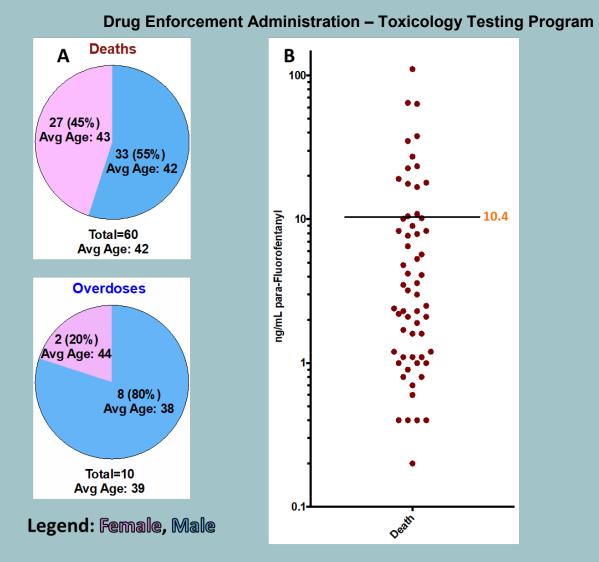


Figure 13. Gender and age (A) and blood para-fluorofentanyl concentration (B) distribution in deaths

The concentration ranges for *para*-fluorofentanyl found in blood samples from various states in death and overdose cases are presented in Table 12.

Table 12. Blood concentration ranges of <i>para</i> -fluorofentanyl observed in cases from
different states

State	Death Cases Overdose Cases		e Cases	
	Concentration (ng/mL)	Frequency	Concentration (ng/mL)	Frequency
Alabama	6.5	1		
California	1.6-8.3 (Avg = 6.8)	3		
Georgia	0.7-22.7 (Avg = 7.6)	6		
Kentucky	3.2	1		
Nebraska	0.4-64.6 (Avg = 21.4)	5	14-14.2	2
Tennessee	0.2-111 (Avg = 10)	44		

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Polydrug use is also prevalent in *para*-fluorofentanyl cases (Fig. 14). Of the 69 cases with biological samples two or more drugs were confirmed in 67 cases (97%).

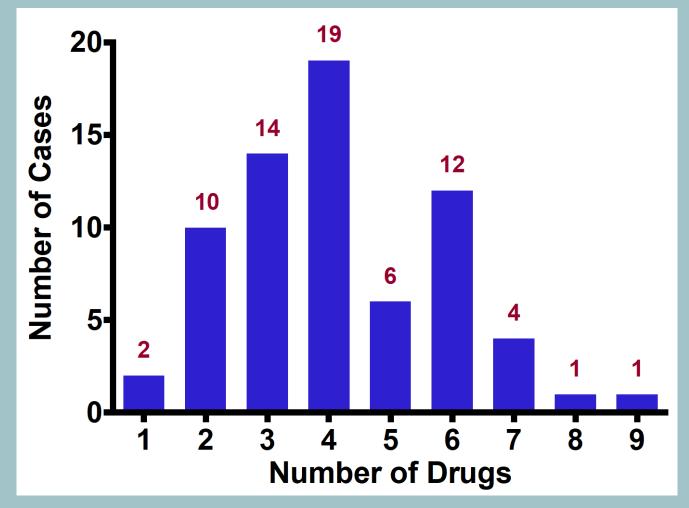


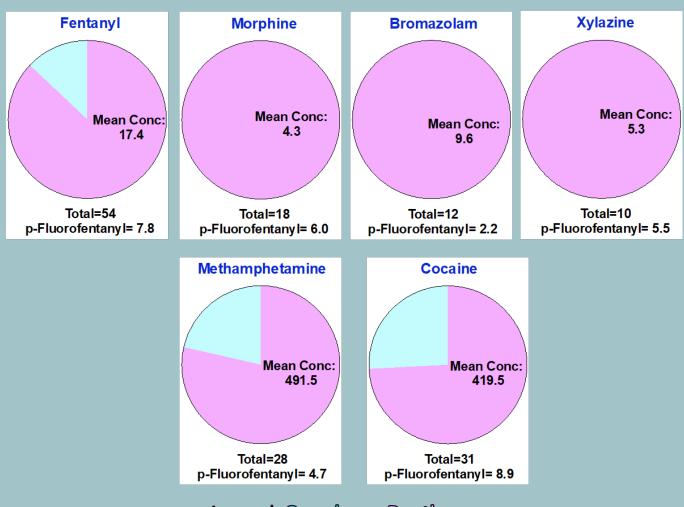
Figure 14. Distribution of the number of NPS and TID in para-fluorofentanyl cases

Like fentanyl, other drugs are confirmed with *para*-fluorofentanyl. Drugs with 7 or more cooccurrences are summarized in Table 13.

Table 13. Co-occurrence frequency of para-fluorofentanyl with other drugs and
adulterants

Drug	Co-occurrence Frequency		
	Overall	Deaths	Overdoses
Methamphetamine	28	22	6
Cocaine/Benzoylecgonine	31	23	8
Fentanyl	54	47	7
Morphine	18	18	0
Bromazolam	12	12	0
Xylazine	10	10	0
Quinine	7	5	2

The distribution between overdose and death cases in these co-occurrences and the average concentration of co-occurring drug and *para*-fluorofentanyl in death cases are presented in Fig. 15.



Legend: Overdoses, Deaths

Figure 15. Distribution between deaths and overdoses in co-occurrence of *para*fluorofentanyl with selected drugs along with the average concentration in ng/mL of the co-occurring drug (in pie chart) and *para*-fluorofentanyl in death cases

Metonitazene

DEA TOX confirmed metonitazene in 15 whole blood samples from 15 cases in 2022. All are death cases except for one overdose case. All death cases were referred from Tennessee while the overdose case was from Kentucky. The gender distribution and average age and the distribution of blood metonitazene concentrations are shown in Fig. 16A and 16B, respectively.

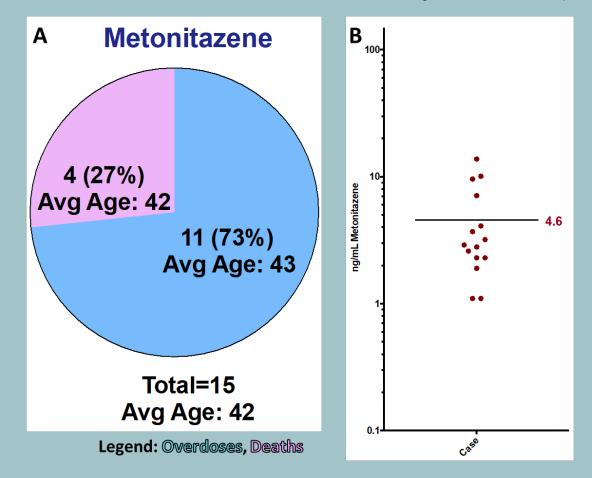


Figure 16. Gender and age (A) and blood concentration (B) distribution in metonitazene cases

Metonitazene was confirmed in these cases along with other drugs. The drugs that often cooccurred with metonitazene are summarized in Table 14. Fentanyl and methamphetamine are detected in 80% of metonitazene cases.

Table 14. Co-occurrence frequency and blood concentration ranges of other drugs with
metonitazene

Drug	Detection	Blood Concentration Range,
	Frequency	ng/mL
Fentanyl	12	1.6- 63.8 (Avg = 18.2)
Methamphetamine	12	1.1-4510 (Avg = 604.1)
Morphine	5	0.6- 4.2 (Avg = 2.2)
Xylazine	5	1.7- 5.9 (Avg = 3.9)

Drug Enforcement Administration – Toxicology Testing Program Bromazolam and Clonazolam

Bromazolam and clonazolam (detected as the metabolite 8-aminoclonazolam) are the most detected designer benzodiazepines in DEA TOX in 2022. The sources of the cases along with the blood concentration ranges observed involving these drugs are shown in Table 15.

Table 15. Detection frequency (DF) and blood concentration ranges of bromazolam and8-amino clonazolam observed in cases from different states

State	Bromazolam		8-Amino Clonazolam	
	Blood Concentration (ng/mL)	DF	Blood Concentration (ng/mL)	DF
Georgia	17.5-46.5	2	0.4-7.4	3
Kansas			4.6	1
Kentucky	5-22.1	2		1
Louisiana			0.4	1
Maryland			11.6	1
Nebraska			0.7	1
New Mexico	2.1	1		
Tennessee	0.9-62.9	15	1.6-47.2	11
Utah	3	1	0.4	1

The average age and gender distribution as well as the distribution between death and overdose cases involving these drugs are shown in Fig. 17.

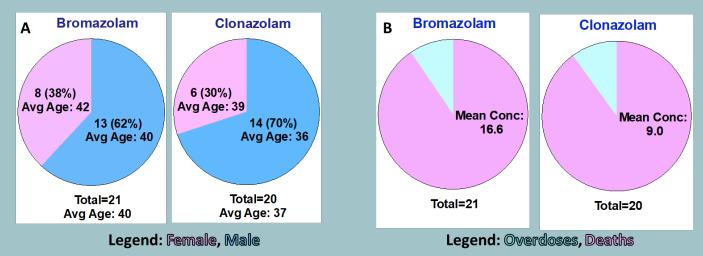


Figure 17. A. Gender distribution and average age in cases with bromazolam and clonazolam. B. Distribution between deaths and overdoses and average blood concentration (ng/mL) in death cases with bromazolam and clonazolam

Other drugs are confirmed in cases with clonazolam and bromazolam. The more frequently cooccurring drugs are summarized in Table 16. Bromazolam co-occurred more frequently with fentanyl (90%) than clonazolam (75%).

Table 16. Co-occurrence frequency and average blood concentration of other drugswith clonazolam and bromazolam

Drug	Co-occurrence Frequency (Ave Blood Concentration in Death Cases, ng/mL)	
	Bromazolam	Clonazolam*
Bromazolam	21 (Avg = 16.6)	5 (Avg = 16.4)
Clonazolam	5 (Avg = 7.1)	20 (Avg = 9.0)
Fentanyl	19 (Avg = 16.6)	15 (Avg = 18.7)
para-Fluorofentanyl	12 (Avg = 2.3)	4 (Avg = 2.5)
Methamphetamine	12 (Avg = 763.7)	7 (Avg = 686.6)
Cocaine/Benzoylecgonine	5 (Avg = 213.1)	6 (Avg = 368)
Morphine	5 (Avg = 3.6)	5 (Avg = 26.4)
Oxycodone	2	4
Mitragynine	3	4
Delta-9 THC	2	4
Xylazine	3	4

* Measured as 8-Amino Clonazolam

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Xylazine

DEA TOX confirmed xylazine in 22 cases, 20 of which also had confirmed fentanyl; the other two cases had confirmed *para*-fluorofentanyl. The 22 cases were referred from Alabama (1), Kentucky (5), Nebraska (1) and Tennessee (15). Xylazine was confirmed in 16 whole blood (15 deaths, 1 overdose) and 6 urine (overdoses) samples. The gender and age, case type, and blood concentration distribution in cases with confirmed xylazine are shown in Fig. 18.

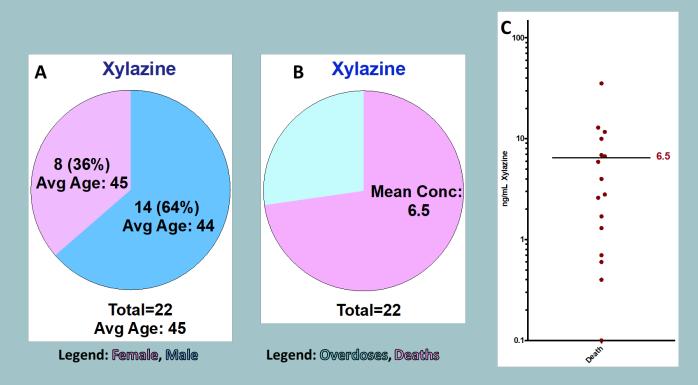


Figure 18. Gender and age (A), case type (B) and blood concentration (C) distribution in xylazine cases. The mean concentration given is in ng/mL.

Although xylazine was often detected as a fentanyl product adulterant, there are other drugs that were frequently confirmed along with it. Drugs that co-occurred five times or more with xylazine along with their blood concentration ranges in death cases are presented in Table 17.

Table 17. Co-occurrence frequency and blood concentration ranges of other drugs with	
xylazine	

Drug	Detection Frequency	Blood Concentration Range in Death Cases, ng/mL
Fentanyl	20	2.1- 57.8 (Avg = 16.7)
para-Fluorofentanyl	11	0.8-35.1 (Avg = 7.0)
Methamphetamine	11	1.1- 4510 (Avg = 705.5)
Cocaine	8	198- 1010 (Avg = 811.2)
Morphine	10	0.6-10.2 (Avg = 3.5)
Metonitazene	5	1.9- 3.7 (Avg = 2.7)
Quinine	5	4.3-25.4 (Avg = 14.8)

Drug Enforcement Administration – Toxicology Testing Program Contact Information

We invite medical and law enforcement facilities to contact our program if you encounter an overdose of a suspected synthetic drug and desire to have any leftover biological samples (blood preferred) analyzed further for such synthetic substances.

Sample Qualification:

 Patients thought to have ingested a synthetic drug, where the traditional drug screen has produced little or no viable options to explain the symptoms exhibited by the patient (alcohol and THC are exempted).

How to Contact Us and Send Your Samples:

- Once the above qualification is satisfied:
 - Email <u>DEATOX@DEA.GOV</u> with a brief description of the case (including initial toxicology screen and history) and a request for testing.
 - DEA will respond to each inquiry, and if approved, will send the instructions for packing and shipping of sample(s) to UCSF.
 - The main reason for disapproval of a case would be the identification of substances including methamphetamine, heroin, fentanyl, cocaine, LSD, PCP etc. in a routine toxicology screening at your facility.
 - This program's goal is to connect symptom causation to abuse of newly emerging synthetic drugs (e.g. synthetic cannabinoids, synthetic cathinones, fentanyl-related substances, other hallucinogens etc.).
- Ensure that you de-identify and label the sample with a numerical value, sex, date of birth or age, and the date and time the sample was collected in accordance with the labeling instructions (sent with shipping instructions).
- Keep a master list of the patients and the numerical values you allocated to each sample at your institution.

• Cost of Sample Analysis:

- DEA will cover the full cost of testing the patient samples.
 - The sender will only be responsible for paying for packing and shipping samples to UCSF.

• Turn-around Time:

• Results are expected within three to four weeks of receipt of the sample at UCSF except in rare occurrences when a novel substance is identified.

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https://www.deadiversion.usdoj.gov/dea_tox/index.html.

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DEA PRB 05-15-2023-11

Appendix Laboratory Method

Sample Analysis

Each sample is analyzed by targeted and suspect screening through LC-QTOF/MS using non-targeted data acquisition. Confirmed drugs are quantified through targeted LC-QTOF/MS using the isotope dilution method.

Sample extracts or diluted samples are analyzed by LC-QTOF/MS (Agilent LC1260- QTOF/MS 6550, Sta. Cruz, CA) using our published comprehensive drug screening method. Chromatographic separation through an Agilent Poroshell 120 column (2.1X 100mm, 2.7 µm) is achieved by gradient elution. Eluates from the chromatographic column are ionized in the QTOF/MS using an electrospray ionization source in positive polarity on one run and negative polarity on a subsequent run. Using non-targeted data acquisition, TOF-MS (parent ion) and MS/MS (fragment ions) spectra are collected in automated MS/MS mode (information-dependent acquisition). Quantification of confirmed drugs is performed by isotope dilution method using a contemporary six to eight-point calibration curve and deuterated or carbon-13-labelled drug isotopologues as internal standards. In cases where a suspected drug is exceptionally potent or rapidly metabolized, the sample is also analyzed using a higher-sensitivity targeted method.

Data Analysis

To confirm the presence of specific drugs in each sample, the total ion chromatogram (TIC) obtained from the LC-QTOF/MS run is analyzed using Agilent MassHunter Qualitative Analysis software. Both targeted and suspect screening are performed in analyzing each sample using the "Find by Formula" algorithm. For targeted screening, a database of 1193 drugs, including 912 NPS, is used as reference for compound matching using the following criteria: mass error ≤ 10 ppm; retention time ≤ 0.15 min; target score ≥ 70 (indication of isotopic pattern match) for peaks that did not exhibit detector saturation; and, the presence of at least one major fragment ion peak in its MS/MS spectrum. For suspect screening, suspect NPS databases are used with the following criteria for a suspect compound match: mass error ≤ 10 ppm; target score ≥ 70 for peaks that did not exhibit detector saturation; and, retention time plausibility. Suspect compounds are confirmed by verifying that the retention time and mass spectral properties of the suspect compound match those of the relevant reference

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standard, including fragmentation patterns observed in MS/MS data collected from the analyzed samples and standards.

Quantitative analysis of confirmed compounds is performed using the Agilent MassHunter Quantitative Analysis software. A linear regression fit between the peak area ratios of spiked reference standard and relevant internal standard, and the known concentrations of the spiked reference standards in a matrix blank is used to quantify the concentration of the confirmed compound. Because it is too expensive and impractical to buy an internal standard for each of the 1193 drugs in our comprehensive drug library, we use a mixture of 15 internal standards that cover the entire range of retention times of our target drugs. The internal standard with a retention time closest to a target drug is used for quantitation. This is a common approach used for comprehensive drug panels.