



DEA TOX

DRUG ENFORCEMENT ADMINISTRATION
TOXICOLOGY TESTING PROGRAM

QUARTERLY REPORT

1st Quarter – 2021



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

Contents

Introduction.....	3
Summary.....	4
NPS Discovered via DEA TOX	5
New Psychoactive Substances	7
Traditional Illicit Drugs	9
Prescription and Over the Counter Drugs	10
Contact Information.....	12

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, fentanyl-related substances, 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 THC are exempted). DEA TOX may approve leftover un-used biological samples (or biological samples) for testing from a medical facility or law enforcement partner only.

Once DEA TOX is contacted (DEATOX@USDOJ.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. The CTEB laboratory currently maintains a comprehensive drug library consisting of 910 new psychoactive substances (NPS), 167 traditional illicit drugs (TID), and 92 prescription or over-the-counter (OTC) drugs.

This publication presents the results of cases analyzed and completed by the CTEB laboratory from January 1, 2021 through March 31, 2021.

For more information, please visit:

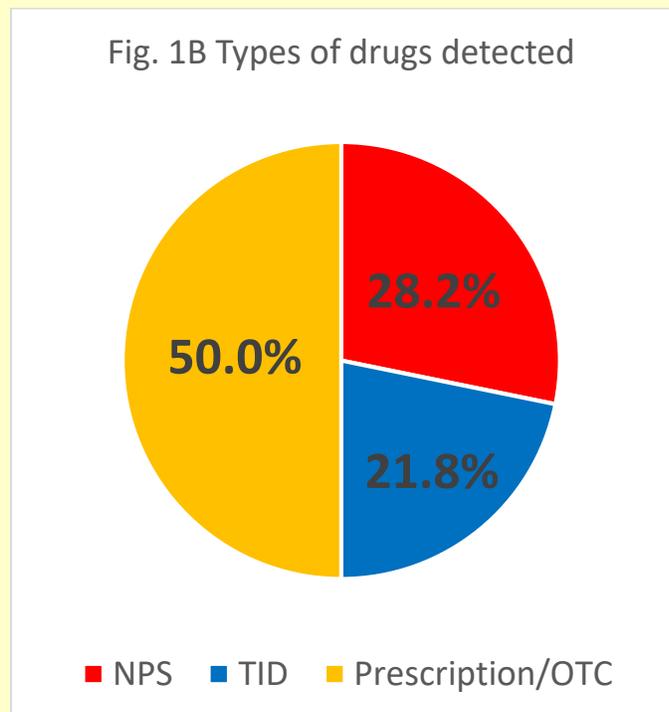
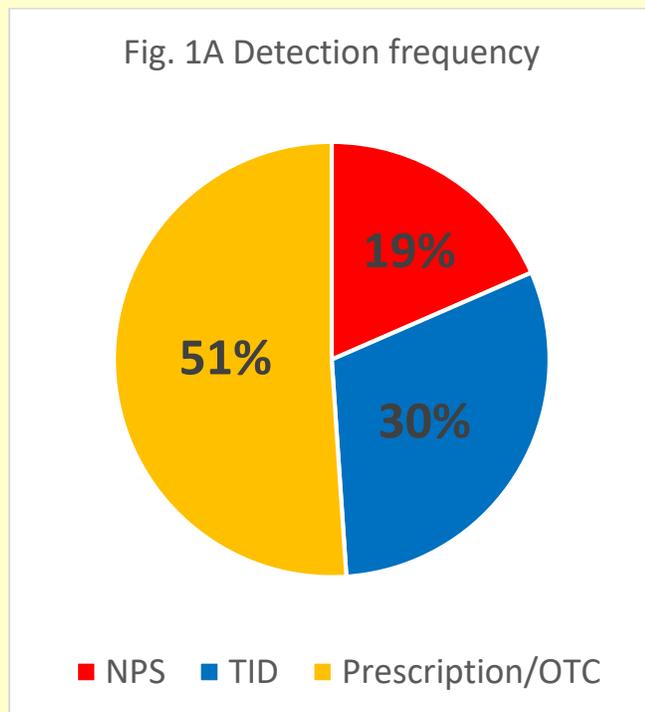
https://www.deadiversion.usdoj.gov/dea_tox/index.html

Summary

Between January 1, 2021 through March 31, 2021, biological samples from 105 cases originating from eight states namely Alabama (38), California (1), Georgia (26), Kansas (27), Kentucky (9), New York (1), North Carolina (1) and Oregon (2) were submitted to DEA TOX. These samples were analyzed for NPS, TID, and prescription or OTC drugs.

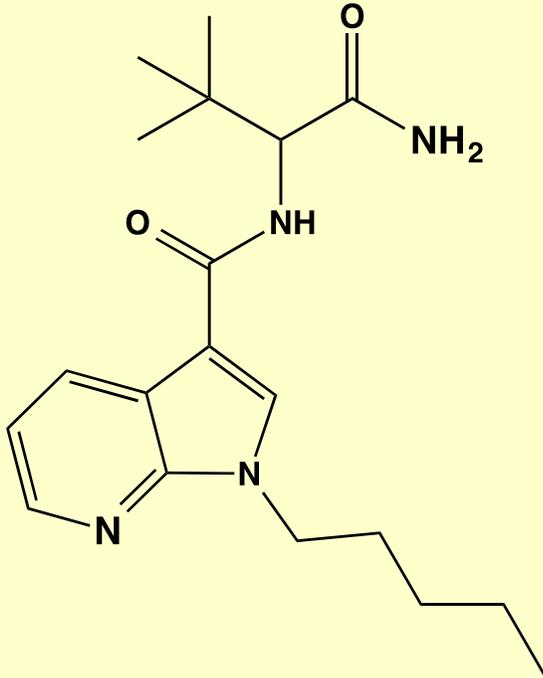
DEA TOX identified and confirmed a total of 325 drugs and metabolites that consisted of 60 NPS detections, 99 TID detections, and 166 prescription or OTC drug detections during this reporting period (Fig. 1A). While some drugs identified could be placed in more than one category, for purposes of this report and for consistency, DEA TOX placed such substances in a single category only. Substances that are not approved by the Food and Drug Administration for medical use within the U.S. are considered NPS.

A breakdown of the 325 total drug and metabolite confirmations demonstrated 78 different drugs, which consisted of 22 NPS, 17 TID, and 39 prescription and OTC drugs (Fig. 1B).



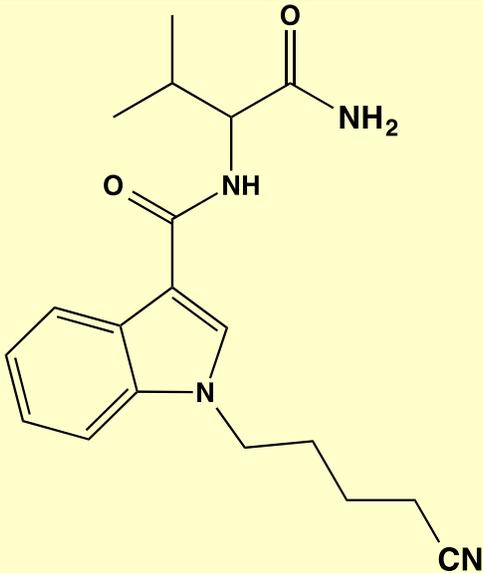
NPS Discovered via DEA TOX

ADB-P7AICA (synthetic cannabinoid)



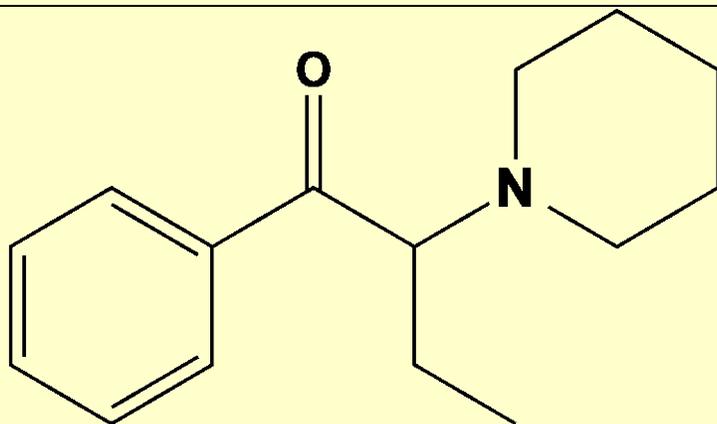
https://www.dea diversion.usdoj.gov/dea_tox/ADB-P7AICA.pdf

4CN-AB-BUTICA (synthetic cannabinoid)



https://www.dea diversion.usdoj.gov/dea_tox/4CN-AB-BUTICA.pdf

alpha-Piperidinobutiophenone (α -PipBP) (synthetic cathinone)



<https://www.dea.gov/divisions/cead/announcements/alpha-PipBP.pdf>

New Psychoactive Substances

DEA TOX confirmed 60 detections comprising of 20 NPS (Table 1) from nine different classes of drugs (Figure 2A) in the 1st quarter of 2021. The total encounters for each NPS class are summarized in Figure 2B.

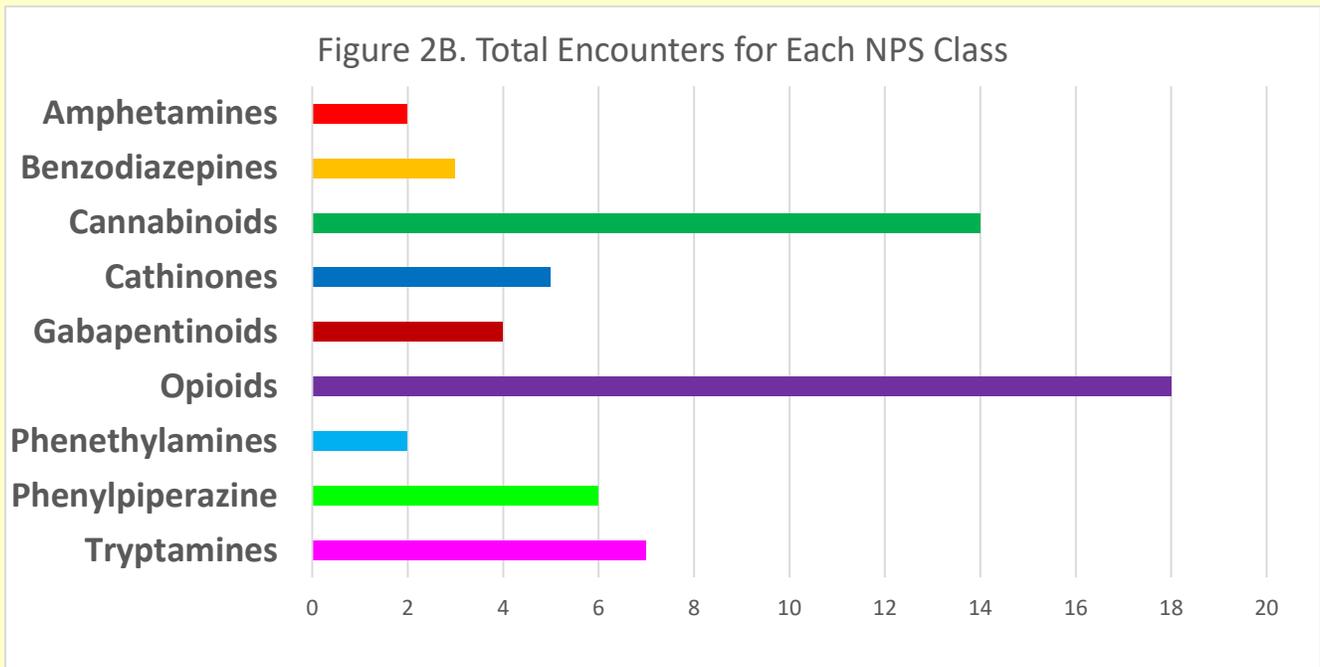
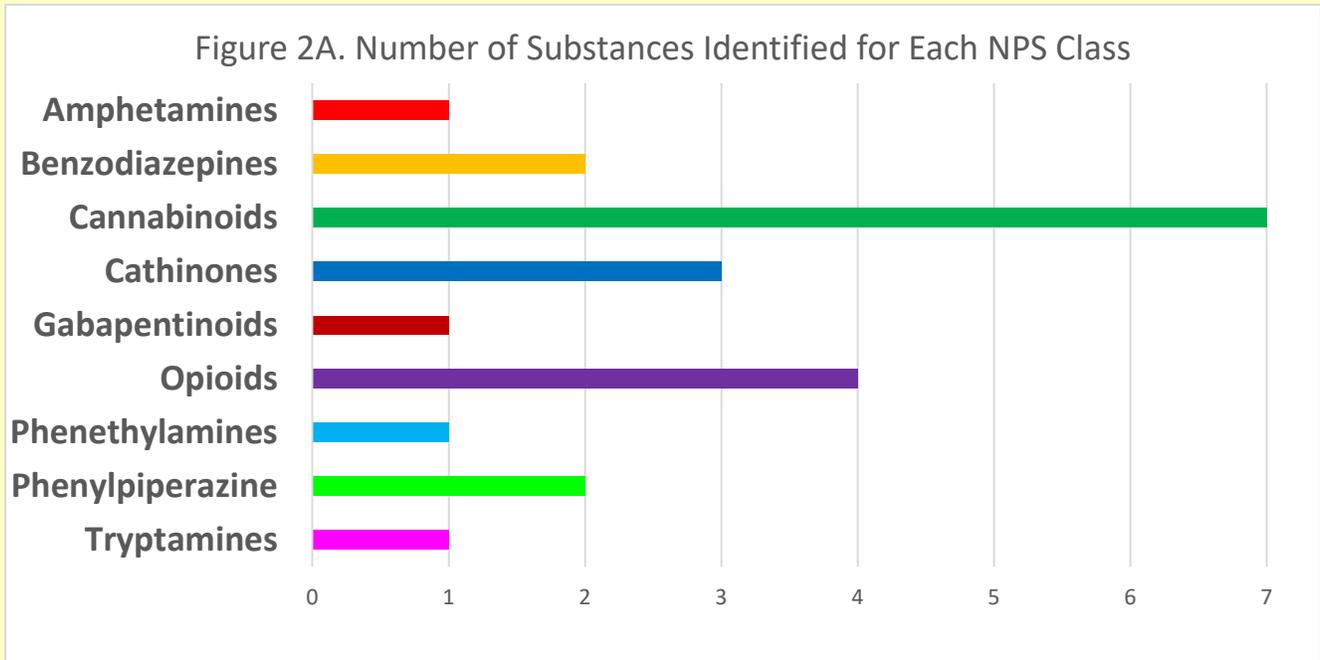


Table 1. NPS detected – 1st Quarter 2021*

Amphetamines	Ethylamphetamine (1) (AL)
Benzodiazepines	Etizolam (2) (KY) Flubromazolam (1) (KS)
Cannabinoids	4CN-AB-BUTICA (1) (AL) 4F-MDMB-BUTINACA (1) (KS) 4F-MDMB-BUTINACA acid (4) (KS, GA) 5F-MDMB-PICA acid (1) (KS) 11-nor-9-carboxy-delta-8-THC (2) (KY) ADB-P7AICA (1) (AL) MAB-CHMINACA M11 metabolite (1) (KS) MDMB-4en-PINACA (2) (KS) delta-8 THC (1) (KY)
Cathinones	alpha-PipBP (a-piperidinobutiophenone) (1) (AL) N-ethylhexylone (2) (AL) NRG-3 (2) (AL)
Gabapentinoid	Phenibut (4) (AL)
Opioids	7-hydroxymitragynine (3) (GA, NC) beta-Hydroxyfentanyl (2) (KS, KY) Mitragynine (5) (GA, KY, NC) Norsufentanil (1) (AL) Tianeptine [§] (7) (AL, KY)
Phenethylamine	2C-P (2) (KS)
Phenylpiperazine	meta-Chlorophenylpiperazine (mCPP) (4) (GA, KY) para-Chlorophenylpiperazine (pCPP) (2) (AL)
Tryptamine	alpha-Methyltryptamine (7) (AL, GA)

* Numbers in parentheses refer to the total number of cases positive for the respective substance followed by the state abbreviations where the sample was collected.

§ Tianeptine, a tricyclic antidepressant, is not approved by the Food and Drug Administration for medical use within the U.S. Tianeptine also has been shown to be a mu-opioid receptor agonist and it is currently being abused in high dosages for its opioidergic activity.

AL – Alabama; GA – Georgia; KS – Kansas; KY – Kentucky; NC – North Carolina

Traditional Illicit Drugs

DEA TOX confirmed 99 detections comprising of 17 TIDs (Table 2) in the 1st quarter of 2021.

Table 2. TID detected – 1st Quarter 2021*

Amphetamines	4-hydroxymethamphetamine (3) (AL, KS) Amphetamine (2) (AL) MDA (1) (AL) MDEA (1) (NC) Methamphetamine (10) (AL, GA, KS, NC)
Arylcyclohexylamine	Ketamine (2) (KS, KY) PCP (1) (KS)
Cannabinoid	11-Nor-9-carboxy-THC (13) (KS, KY, NC) THC (2) (OR)
Cocaine	Benzoylcegonine (10) (AL, GA, KS, KY, NC, NY) Cocaine (5) (AL, NY, NC) Ecgonine Methyl Ester (8) (AL, KS, KY, NC, NY)
Ergolines	LSD (1) (KY)
Opioids	6-Acetylmorphine (1) (AL) Acetylcodeine (1) (AL) Codeine (2) (AL) Fentanyl (17) (AL, CA, KS, KY, NC, NY) Hydrocodone (1) (GA) Morphine (2) (AL) Norfentanyl (13) (AL, CA, KS, KY, NC, NY) Oxycodone (2) (AL, KS) Oxymorphone (1) (AL)

* Numbers in parentheses refer to the total number of cases positive for the respective substance followed by the state abbreviations where the sample was collected.

AL – Alabama; CA – California; GA – Georgia; KS – Kansas; KY – Kentucky; NC – North Carolina; NY – New York; OR – Oregon

Prescription and Over the Counter Drugs

DEA TOX confirmed 166 detections comprising of 39 prescription or OTC drugs (Table 3) in the 1st quarter of 2021.

Table 3. Prescription or OTC drugs detected – 1st Quarter 2021*

Anesthetic	Lidocaine (1)
Antibiotic	Levofloxacin (1) Sulfamethoxazole (1)
Anticonvulsant	Gabapentin (4) Lamotrigine (1) Oxcarbazepine (3)
Antidepressant	Citalopram (1) Fluoxetine (3) Mirtazapine (3) Norfluoxetine (1) Paroxetine (2) Sertraline (3) Trazadone (6) Venlafaxine (3)
Antihistamine	Chlorpheniramine (1) Cimetidine (2) Diphenhydramine (8) Hydroxyzine (4)
Antipsychotic	Haloperidol (4) Olanzapine (2) Ziprasidone (3)
Antiviral	Emtricitabine (2)
Benzodiazepine	7-Aminoclonazepam (1) Alpha-hydroxyalprazolam (2) Alprazolam (4) Diazepam (1) Midazolam (7) Nordiazepam (2) Oxazepam (2) Temazepam (2)
Cardiovascular	Amiodarone (1) Atropine (2) Carvedilol (3) Lisinopril (3) Metoprolol (1)

Drug Enforcement Administration – Toxicology Testing Program

	Mexilitine (1) Propranolol (3) Verapamil (1)
Decongestant	Phenylephrine (1) Phenylpropanolamine (1) Pseudoephedrine (1)
Muscle Relaxant	Baclofen (3)
Opioids	Buprenorphine (9) Naloxone (15) Norbuprenorphine (9) Tramadol (1)
Pain Reliever	Acetaminophen (26) Naproxen (3)
Respiratory	Albuterol (2)

* Numbers in parentheses refer to the total number of cases positive for the respective substance.

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 Qualifications:**

- 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 qualifications are satisfied:
 - Email DEATOX@USDOJ.GOV 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 weeks of receipt of the sample at UCSF except in rare occurrences when a novel substance is identified.

PUBLIC DOMAIN NOTICE

All material appearing in this publication is in the public domain and may be reproduced or copied without permission from the DEA. However, this publication may *not* be reproduced or distributed for a fee without the specific, written authorization of the U.S. Drug Enforcement Administration, U.S. Department of Justice. Citation of the source is appreciated.

Suggested citation:

U.S. Drug Enforcement Administration, Diversion Control Division. (2021). *DEA TOX: Quarterly Report – 1st Quarter 2021*. Springfield, VA: U.S. Drug Enforcement Administration.

OBTAINING COPIES OF THIS PUBLICATION

Electronic copies of this publication can be downloaded from the DEA TOX website at:

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

This report was produced in conjunction with the CTEB laboratory at UCSF.



**Clinical Toxicology
and Environmental Biomonitoring Laboratory**

DEA PRB 04-19-21-23