Announcement of a Newly Identified Synthetic Cannabinoid
4CN-AB-BUTICA – February 25, 2021

The US Drug Enforcement Administration in collaboration with the University of California San Francisco Clinical Toxicology and Environmental Biomonitoring (CTEB) Laboratory has identified a new synthetic cannabinoid, 4-CYANO-AB-BUTICA (4CN-AB-BUTICA), in urine samples submitted to our New Psychoactive Substances (NPS) surveillance program.¹

**Cohort:** In November 2020, prisoners under a work release program in Alabama were suspected of using tianeptine and a suspected synthetic cannabinoid-laced drug product inmates referred to as “No Show”.

**Drugs Detected:** 4CN-AB-BUTICA was detected, confirmed, and quantified in the cohort. The majority of urine samples from these individuals contained tianeptine and 4-CN-AMB-BUTINACA. Other prominent substances detected in these samples include cathinones (pentedrone, N-butylpentylone and N-ethylbuphedrone (NEB)), amphetamines (ethylamphetamine and PMMA²), and phenibut.

**4CN-AB-BUTICA:** This new synthetic cannabinoid is related to a number of previously reported synthetic cannabinoids such as 4F-MDMB-BUTICA (4F-MDMB-BICA). 4F-MDMB-BICA was first reported in the United States in July 2020. It was reported to have affinity (EC₅₀ = 121 nM) and efficacy (Eₘₐₓ = 253% relative to JWH-018) towards cannabinoid receptor 1 (CB1) binding.³

**Analysis:** 4CN-AB-BUTICA was detected, confirmed and quantified in urine samples using liquid chromatography-quadrupole time-of-flight mass spectrometry. Details of the method used along with the chromatogram and mass spectra associated with this substance are presented below.

**Reference Standard:** There is currently no commercially available reference standard for 4CN-AB-BUTICA. A reference standard of this substance was synthesized by our program collaborator at the University of Sydney, Dr. Samuel Banister. Chemical characterization of the reference standard was performed by Mr. Eric Sparkes and Dr. Adam Ametovski and is detailed below. This reference standard is part of the prophetic cannabinoids standards library proactively synthesized to allow our surveillance to identify and confirm previously unreported NPS in real time.

**Pharmacological Data:** 4CN-AB-BUTICA has been shown to bind to CB1 receptors (Kᵢ = 548 nM) in a binding assay performed by our program collaborators, Ms. Shuli Chen and Dr. Michelle Glass at the University of Otago.

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¹ This report was prepared by Roy Gerona, Ross Ellison, Eric Sparkes, Adam Ametovski, Shuli Chen, Michelle Glass, Samuel Banister and Jordan Trecki.
² PMMA [1-(4-methoxyphenyl)-N-methylpropan-2-amine; para-methoxymethamphetamine]
4CN-AB-BUTICA

I General Information

Synonyms: 4CN-AB-BICA, AB-4CN-BUTICA, AB-4CN-BICA

IUPAC Name: \((\text{S})-N-(1\text{-amino-3-methyl-1-oxobutan-2-yl})-1\text{-(4-cyanobutyl)-1\text{-H-indole-3-carboxamide}}\)

InChI String: InChI=1S/C19H24N4O2/c1-13(2)17(18(21)24)22-19(25)15-12-23(11-7-3-6-10-20)16-9-5-4-8-14(15)16/h4-5,8-9,12-13,17H3,6-7,11H2,1-2H3,(H2,21,24)(H,22,25)/t17-/m0/s1

InChI Key: BEQGVLCHHJZNFY-KRWDZBQOSA-N

SMILES: \(\text{O=C(N}\,[\text{C@@H}](\text{C}(\text{C})(\text{C})(\text{N})=\text{O})\text{C}1=\text{CN}(\text{CCCCC}\#\text{N})\text{C}2=\text{C}1\text{C}=\text{CC}=\text{C2}\)

CAS Number: Not available

CFR: Not scheduled (02/2021)

II Physical Properties

Melting Point: 177 – 180 °C

III Chemical Characterization

FTIR Spectrum: \(\nu_{\text{max}} \text{ (cm}^{-1}\text{)}: 3293, 1620, 1514, 1466, 1394, 1269, 1231, 1163, 802, 741, 663, 617, 562.\)
$^1$H NMR Spectrum: (400 MHz, DMSO-$d_6$, 298 K) δ 8.25 (s, 1H), 8.13 – 8.08 (m, 1H), 7.57 (dt, $J = 8.3$, 0.9 Hz, 1H), 7.52 – 7.45 (m, 2H), 7.25 – 7.12 (m, 2H), 7.06 (br s, 1H), 4.35 (dd, $J = 8.9$, 7.0 Hz, 1H), 4.25 (m, 2H), 2.55 (t, $J = 7.1$ Hz, 2H), 2.08 (m, 1H), 1.89 (quin., $J = 7.1$ Hz, 2H), 1.57 (quin., $J = 7.5$ Hz, 2H), 0.93 (dd, $J = 6.8$, 3.0 Hz, 6H)

$^{13}$C NMR Spectrum: (101 MHz, DMSO-$d_6$, 298 K) δ 173.5 ((CO)NH$_2$), 164.0 (CO), 136.1 (quat.), 131.3 (CH), 126.5 (quat), 122.0 (CH), 121.1 (CH), 120.7 (CH), 120.5 (CH), 110.4 (quat.), 109.5 (CH), 57.4 (CH), 45.1 (NCH$_2$), 30.5 (CH), 28.8 (CH$_2$), 22.3 (CH$_2$), 19.5 (CH$_3$), 18.5 (CH$_3$), 15.8 (CH$_2$).
IV LC-QTOF/MS Analysis

**Instrument:** Agilent 1260 Infinity, Agilent 6550 QTOF-MS/MS

**Sample Preparation:** Enzymatic deconjugation with *H. pomatia* glucuronidase followed by dilution

**Chromatography**
- **Column:** Agilent Poroshell 120 EC-C18 (100 mm x 2.1 mm, 2.7 µm)
- **Column Temperature:** 50 °C
- **Injection Volume:** 2.5 µL
- **Mobile Phase:**
  - A: Ammonium formate (5 mM) and Formic Acid (12.6 mM) in H₂O
  - B: Formic Acid (12.6mM) in acetonitrile
- **Flow rate:** 0.5 mL/min
- **Elution Profile:** Gradient- 95A:5B initially; 70A:30B from 0.5 to 1.5 min; 30A:70B from 1.5 to 4.5 min; 0A:100B from 4.5 to 7.5min; 95A:5B from 10.0 to 14.0 min
- **Run Time:** 12 min

**Mass Spectrometry**
- **Ion Source:** Dual Jet Stream Electrospray Ionization
- **Polarity:** Positive
- **TOF MS Scan Range:** 75-1000 Da
- **MS/MS Scan Range:** 50-510 Da
- **Gas Temperature:** 225 °C
- **Drying Gas Flow Rate:** 14L/min
- **Sheath Gas Temperature:** 350 °C
- **Sheath Gas Flow Rate:** 11L/min
- **Nebulizer pressure:** 14psi
- **Capillary Voltage:** 3000 V
- **Nozzle Voltage:** 500 V
- **Skimmer Voltage:** 65 V
- **Octopole RF:** 750 V
- **Fragmentor Voltage:** 380 V
- **Internal Reference Masses:** Purine at m/z 121.0509; HP-921 at m/z 922.0098
- **Data Acquisition:** 2GHz, extended dynamic range
- **Fragmentation:** Auto MS/MS, three maximum precursors (threshold: 500 counts) per cycle with active exclusion after 1 spectrum at a 30s release time
Extracted Ion Chromatogram

*Retention Time*: 5.06 min

TOF-MS Spectrum

*Exact Mass*: 340.1899

*Accurate Mass*: \([M+H]^+ = 341.1967 \text{ (mass error=-0.12ppm)}; [M+Na]^+ = 363.1792 \text{ (mass error= 0.39 ppm)}

MS/MS Spectrum
In response to the ongoing synthetic drug epidemic, the Drug Enforcement Administration (DEA) has initiated a contract with the University of California at San Francisco (UCSF) whereby biological samples generated from overdose victims of synthetic drugs can be further analyzed. In many cases, it can be difficult to ascertain the specific substance responsible for the overdose. In the future, we invite you 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 (i.e. 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.
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