



UNODC

United Nations Office on Drugs and Crime

The UNODC Early Warning Advisory On NPS

2nd Annual Meeting Copolad II

Martin Raithelhuber

Global SMART Programme

Laboratory and Scientific Section

November 2017



Laboratory and Scientific Section

- Established in 1954 by **GA Resolution 834 (IX)** as the United Nations Narcotics Laboratory
- Several GA, ECOSOC, CND and CCPCJ Resolutions have mandated areas of work such as:
 - Strengthening **national drug testing laboratories**
 - Developing reliable **field and laboratory testing methods**
 - Establishment a central source of **reference standards** of major drugs
 - Developing **internationally acceptable guidelines** for forensic science practice



Support to national laboratories

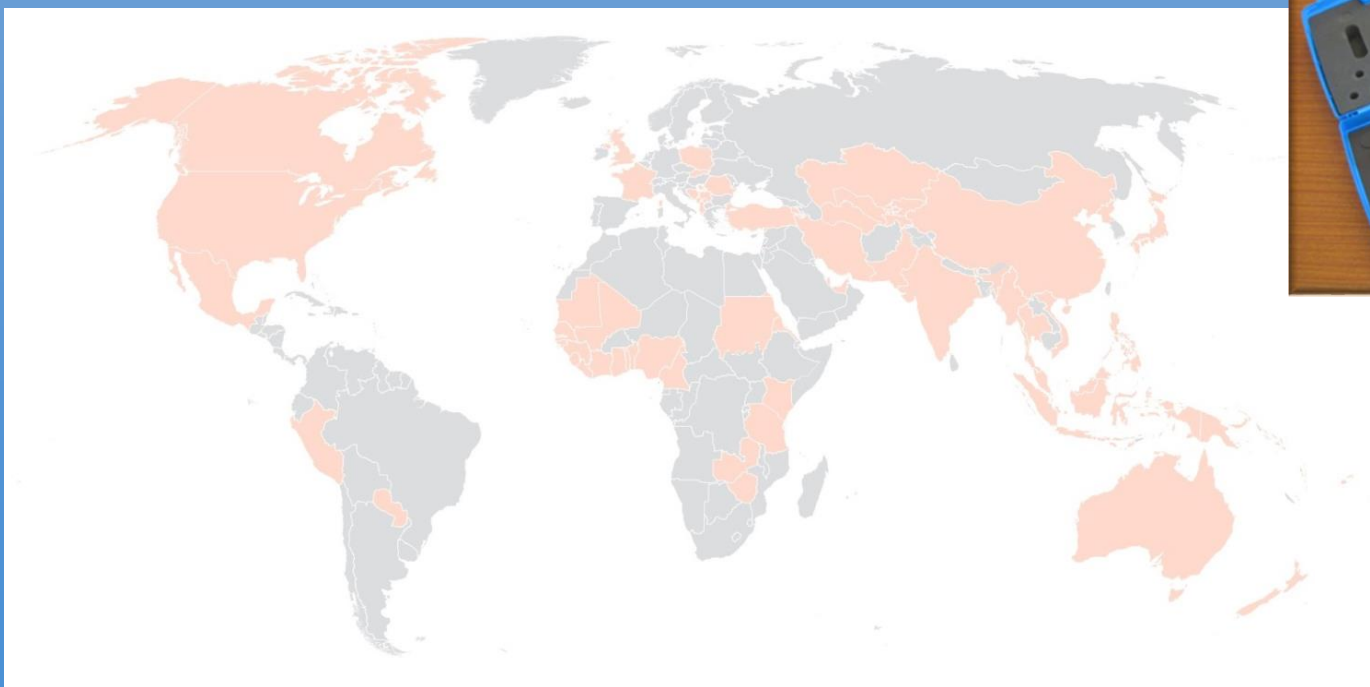
- Onsite training of national drug testing experts in Vienna as part of UNODC regional programmes





Strengthening law enforcement capacity

- Provision of over 3800 drugs and precursors field testing kits to 77 Member States





Strengthening law enforcement capacity

- Introducing modern hand-held technologies in field drug and precursor testing trainings
- Field-based trainings and CBTs on field drug testing





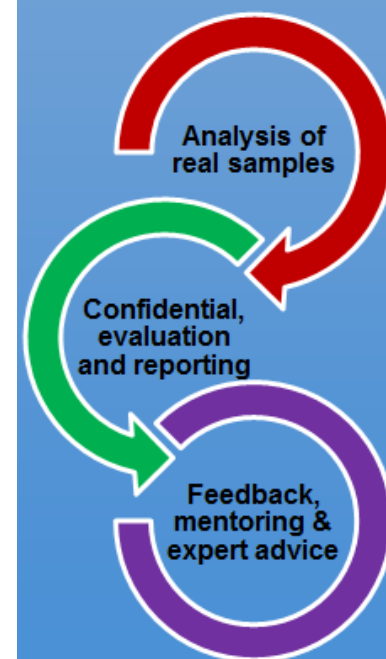
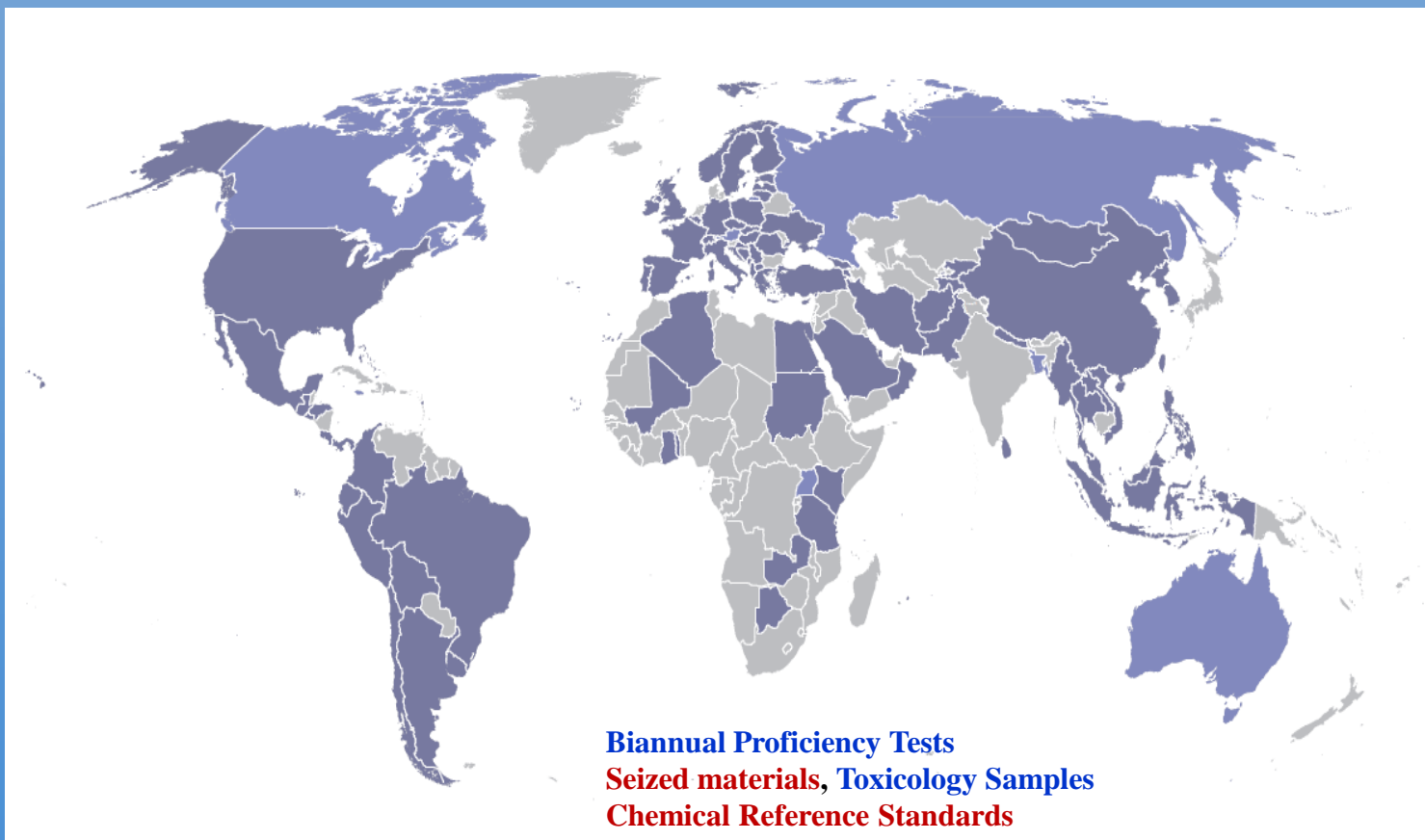
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Global SMART
Programme

Enhancing National Forensic Laboratory Capacity: The UNODC International Collaborative Exercises





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Global SMART
Programme

The Global Synthetics Monitoring: Analysis, Reporting, Trends (SMART) Programme



- As a response to the synthetic drug problem, the UNODC launched the Global SMART Programme in 2008
- The Programme seeks to enhance the capacity of Member States and authorities in priority regions:
 - to generate, manage, analyse and report synthetic drug information
 - and to apply this scientific evidence-based knowledge to design the policies and programmes



“Early warning” at the global level: Understanding a new phenomenon

- 2012, NPS came to the attention of the Commission on Narcotic Drugs as an emerging problem
- In March 2013, the Commission tasked UNODC with the development of an early warning mechanism on NPS
- In June 2013, the UNODC Early Warning Advisory on NPS (EWA) was launched, starting with information from the network of national forensic laboratories
- Task: provide information on NPS trends (which substances where and when) for the Commission to take action
- Product: password protected web platform with information on substance, country, year



“Early Warning” at the global level: Review of NPS for international control

- Challenge: huge data gaps in some regions, sometimes no methodologies available, chemical reference standards not available or too costly, ...
- Enhancement of EWA to include analytical information, molecular structures, development of technical guidelines for analysis of synthetic cannabinoids, piperazines, synthetic cathinones, ...
- Support prioritization of NPS for international review (since 2015): most prevalent, persistent and harmful NPS
- Development of a toxicology module for the EWA to capture information on fatalities and hospitalizations associated with the use of NPS



UNODC Early Warning Advisory on New Psychoactive Substances

- + What are NPS?
- + NPS Substance Groups
 - Aminolindanes
 - Ketamine & Phencyclidine-type substances
 - Other substances
 - Phenethylamines
 - Piperazines
 - Plant-based substances
 - Synthetic cannabinoids
 - Synthetic cathinones
 - Tryptamines
- + Legal Responses
- + Resources
- + Global SMART Programme
- + EWA Partners
- + Latest News on NPS
- + ICE-Portal

UNODC Early Warning Advisory (EWA) on New Psychoactive Substances (NPS)

The EWA provides access to basic information on new psychoactive substances. Specific information on NPS including trend data, chemical details on individual substances, supporting documentation on laboratory analysis and legislative responses can be accessed by [registered users](#) only.



What are NPS?

New psychoactive substances have been known in the market by terms such as 'legal highs', 'herbal highs', 'bath salts', 'research chemicals'

NPS Substance Groups

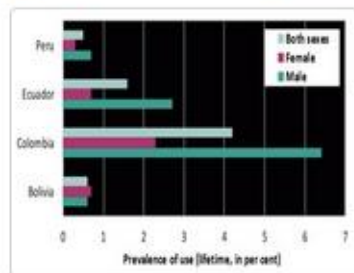
NPS differ greatly in terms of their adverse effects, the ways in which they are abused and their historical background



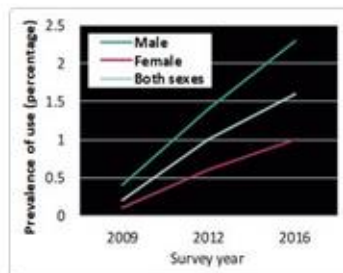
Legal Responses

Countries have adopted various legislative measures to tackle the NPS problem

Latest News on NPS ([view all](#))



November 2017 – Peru: University



October 2017 – Peru: Rise in LSD



October 2017 – UNODC-SMART:



UNODC

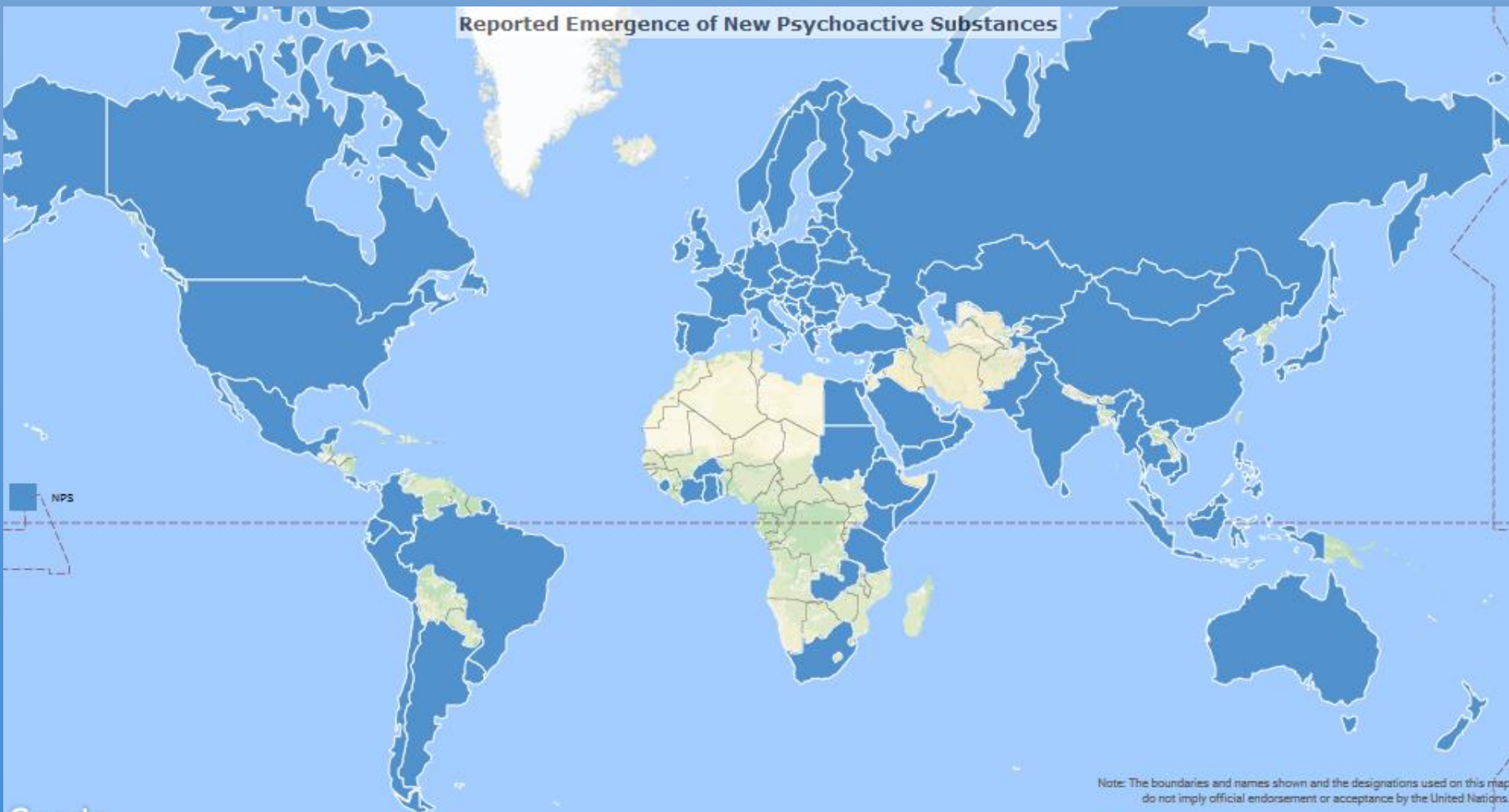
United Nations Office on Drugs and Crime



Global SMART
Programme

Prioritization of NPS: monitoring emergence, prevalence and persistence

Reported Emergence of New Psychoactive Substances

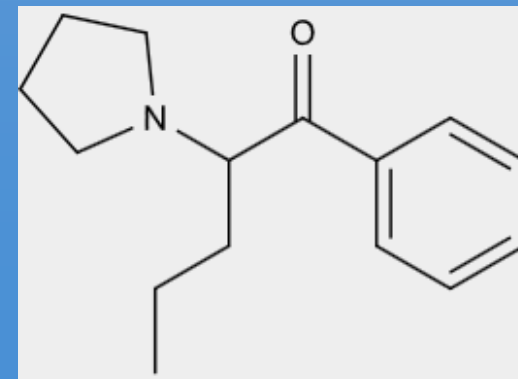
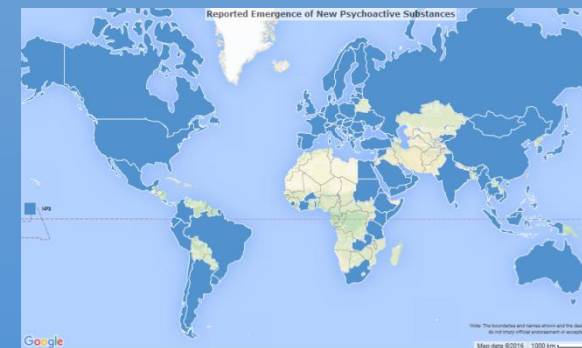


Note: The boundaries and names shown and the designations used on this map do not imply official endorsement or acceptance by the United Nations.



Early Warning Advisory: Clients and needs

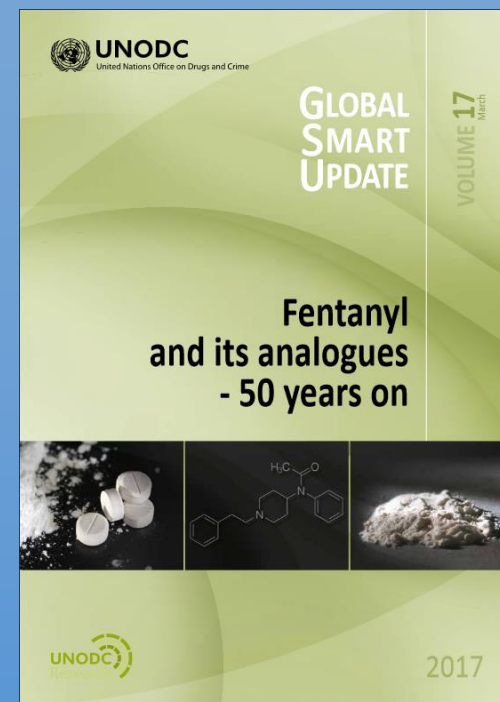
- International bodies and organisations (CND, WHO Expert Committee on Drug Dependence, INCB, ...):
 - Global reference point as a basis for discussion and decisions, NPS trends, maps, terminology, effects, and harms
- National experts, policy and decision-makers:
 - Trend-analysis data and legal approaches
- Laboratories:
 - Spectra for NPS identification, methodologies for analysis, chemical reference standards
- Law enforcement:
 - Tools to detect NPS





Highlighting risks and adverse effects

- For example injecting use of synthetic cathinones (see Global SMART Update volume 16, March 2017)
- EWA newsclips: intoxications, NPS treatment, NPS use trends seizures, national and international legislation



March 2017 - Austria: Fifth International Conference on NPS to be held 23-24 October 2017

Vienna, AUSTRIA – March 2017: We are pleased to announce the Fifth International Conference on NPS to be held from 23 to 24 October 2017 at the United Nations in Vienna, Austria. The conference is jointly organised by the United Nations Office on Drugs and Crime (UNODC), the European Monitoring Centre for Drugs and Drug Addiction (EMCDDA), the World Anti-Doping Agency (WADA), University of Hertfordshire and Sapienza University of Rome.

The first four International Conferences on NPS, held in Budapest (2012); 2015) Swansea (2013) and Rome (2014), were extremely successful events to share knowledge and strengthen collaboration on NPS among multidisciplinary professionals at the international level.

The main objectives of this fifth event are to share evidence-based information on NPS and latest trends of misuse; improve the understanding of clinical treatment and management of NPS use; explore policy and legislative responses to NPS; develop innovative prevention measures for vulnerable individuals; identify public health implications of NPS and explore motivations and socio-cultural factors underlying NPS use. To attend this conference please [register](#) on the conference webpage. [Abstract submissions](#) will open on Wednesday, 1st March 2017 and close on Sunday, 30th April 2017.



June 2016 - Kyrgyz Republic: the synthetic cathinone pentedrone identified and seized based on new NPS legislation in the Kyrgyz Republic

BISHKEK, Kyrgyz Republic – June 2016. On 7 June 2016, during a search operation carried out by employees of State Service on Drug Control (RCKH) and State Customs Service (GTC) in Bishkek, Kyrgyz Republic, two persons were detained on suspicion of involvement in the illicit trafficking of psychotropic substances. During the personal search a plastic bag containing white powder was seized. The State Service on Drug Control experts identified the seized substance as pentedrone, a synthetic cathinone with stimulant effect. The net weight of the seized powder was 29 grams with an estimated value on the illicit market of 100 thousand Kyrgyzstani soms (about USD 1,476).

The Kyrgyz Republic legislation prohibits NPS trafficking and use. On December 4, 2015, through the Decree No 831, the Government of the Kyrgyz Republic amended the list of controlled substances contained in the Decree "On narcotic drugs, psychotropic substances and precursors subject to control in the Kyrgyz Republic", dated November 9, 2007, № 543, to include a number of new psychoactive substances. Over 100 NPS, including pentedrone, were placed under control.





EWA Newsletter – Volume 12 (October 2017)



UNODC Early Warning Advisory on New Psychoactive Substances

You are receiving this newsletter because you are registered to one of the online portals of the UNODC Laboratory and Scientific section and/or are on the mailing list of the Global Synthetics Monitoring: Analyses Reporting and Trends (SMART) Programme.

NEWSLETTER on New Psychoactive Substances
October 2017 Vol. 13

UNODC HIGHLIGHTS

[CND decision on international control of 4-MEC, Ethylone, Pentedrone, Ethylphenidate, MPA, MDMB-CHMICA, 5F-APINACA, and XLR-11 enters into force](#): The decision of the Commission on Narcotic Drugs (CND) during its 60th Session from 13 to 17 March 2017 to add 4-Methylethcathinone (4-MEC), Ethylone, Pentedrone (α -Methylaminovalesterophenone), Ethylphenidate (EPH), Methiopropamine (MPA), MDMB-CHMICA, 5F-APINACA (5F-AKB-48) and XLR-11 to Schedule II of the Convention on Psychotropic Substances of 1971 has entered into force. Previously, on 22 April 2017, the decision adopted by the CND during its 60th Session to add U-47700 and Butyrfentanyl to Schedule I of the Single Convention on Narcotic Drugs of 1961 entered into force.

[2017 Global Synthetic Drugs Assessment is launched](#): The [2017 Global Synthetic Drugs Assessment](#) provides an analysis of the global synthetic drugs market in two parts. The first part consists of regional overviews that highlight context-specific dynamics relating to the demand and supply of Amphetamine-Type Stimulants (ATS) and New Psychoactive Substances (NPS) in Africa, Central and Southwest Asia, East and South-East Asia and Oceania, Europe, the Near and Middle East, North and Central America, and South Central America. The second part of this report presents a global thematic analysis of the key trends and emerging developments of the synthetic drugs market.

[Countries worldwide reported the emergence of 5F-PB-22, UR-144, AB-CHMINACA, AB-PINACA, and ADB-FUBINACA to UNODC](#): The emergence of the five synthetic cannabinoids 5F-PB-22, UR-144, AB-CHMINACA, AB-PINACA, and ADB-FUBINACA, which are among the 16 substances under review by the WHO Expert Committee on Drug Dependence (ECDD), have been reported to the UNODC Early Warning Advisory (EWA) from countries across the world. With the exception of ADB-FUBINACA, each substance has been reported by countries in North and South America, Europe, Central Asia, the Near and Middle East, and East and South-East Asia and Oceania, with the largest number of countries reporting the emergence of 5F-PB-22 (41 countries), followed by UR-144 (32 countries), AB-CHMINACA (31 countries) and AB-PINACA (27 countries). The emergence of ADB-FUBINACA has been reported in 20 countries in North America, Europe, Central Asia, and East and South-East Asia. In September 2017, the EMCDDA and Europol also published the results of a [joint assessment](#) on the availability and spread of AB-CHMINACA within the European Union.

[9th Regional SMART Workshop for East and South-East Asia addresses the rise of methamphetamine and NPS](#): Key trends on New Psychoactive Substances (NPS) and emerging information on synthetic



What does the Early Warning Advisory Offer?

Home About Dashboard Round Management Search NPS Data Maintenance Area

UNODC Early Warning Advisory on New Psychoactive Substances (NPS)

→ Register New Client

→ Clients

→ What are NPS?

→ NPS Substance Groups

- Aminoindanes
- Ketamine & Phencyclidine-type substances
- Other substances
- Phenethylamines
- Piperazines
- Plant-based substances
- Synthetic cannabinoids
- Synthetic cathinones
- Tryptamines

→ NPS Chemical Information

→ NPS Briefs

→ Legal Responses

→ Resources

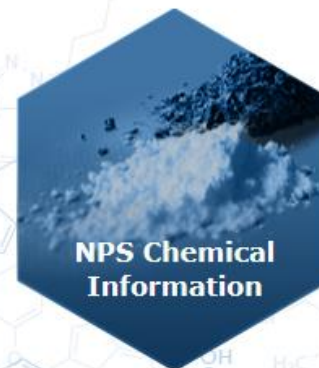
→ EWA Guide

→ Global SMART Programme

→ EWA Partners

→ Latest News on NPS

→  ICE-Portal





UNODC Early Warning Advisory on New Psychoactive Substances



Home Dashboard Search NPS Data

Search NPS Data

Filter:

Year:

<all>
2015
2014
2013

(Ctrl+Click to multi-select)

Substance:

type to select substance

Regions:

-- no region filter --
Africa
Americas
Asia

(Ctrl+Click to multi-select)

Filter

Clear Filter

Show Own/All:

Show All

Substance Group:

Synthetic cathinones

Countries:

-- all --
AFGHANISTAN (Asia)
ÅLAND ISLANDS (Europe)
ALBANIA (Europe)

(Ctrl+Click to multi-select)

Page Size:

30

Selected Countries (ISO-codes):

AF, AM, AZ, BH, BD, BT, BN, KH, CN, CD, KP, GE, HK, IN, ID, IQ, IR, IL, JP, JO, OM, PK, PH, QA, KR, SA, SG, LK, SY, TW, TJ, TH, TL, TM, AE, UZ, VN, YE

Access filtered data:



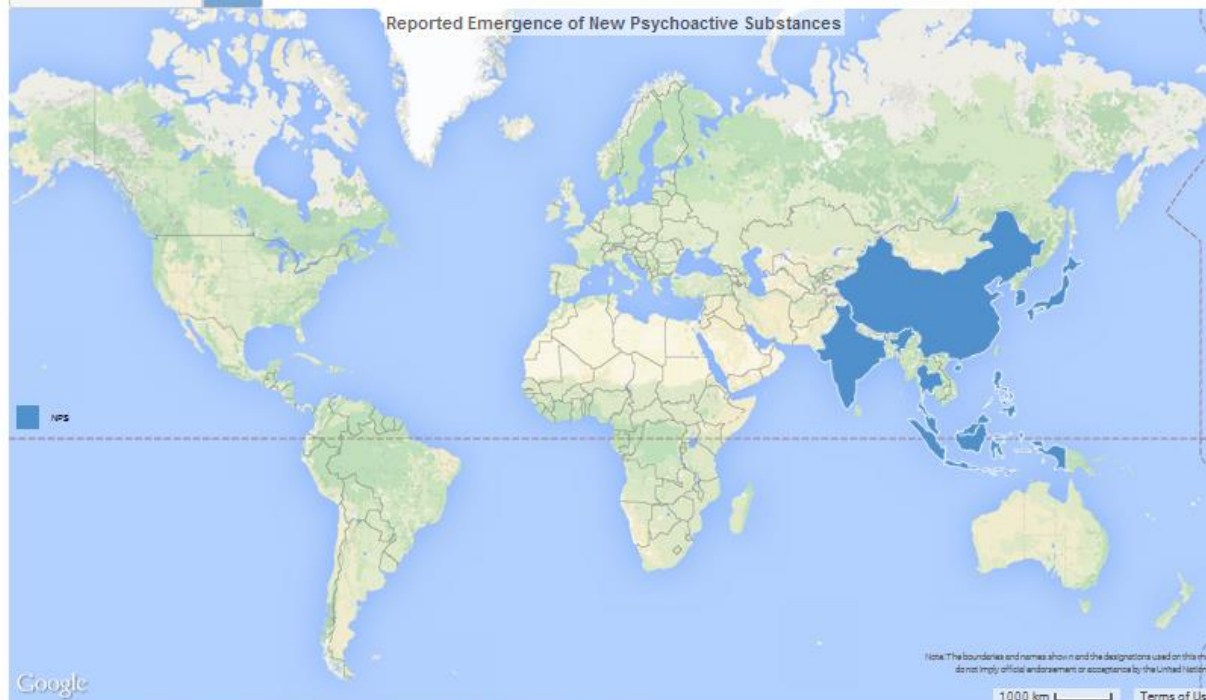
Summary Report (charts/tables)



Detailed Substance Finding List

Substance Finding List

Map



Use filters to check
which NPS emerged
where and when!

PDF summary
report, excel list

Create an
online map of
NPS
emergence
based on
your search
criteria!



UNODC Early Warning Advisory on New Psychoactive Substances



NPS Substance Finding

Substance ([open comprehensive list](#))

Quantity

g or ml

Finding Date

Means Of Identification (Ctrl-click to multiselect)

Analytical technique(s)

Labelling

Online database

Packaging

Description

ATC Identification / Comment

ATC Confirmation / Comment

Comment

Attachment

add attachment

Please ...

select substance

select means of identification

Delete

Cancel

Submit a NPS finding
in your country with
analytical information
using the online form!



UNODC Early Warning Advisory on New Psychoactive Substances

Home Dashboard Search NPS Data

NPS Chemical Information

Find more
information on
individual NPS

NPS Chemical
Information

Name	Substances
Aminoindanes	2,3-Dihydro-1H-Inden-1-amine, 1-Aminoindan
	2-Aminoindane, 2,3-Dihydro-1H-inden-2-amine (2-AI) [2975-41-9]
	5,6-Methylenedioxy-2-aminoindane, 6,7-Dihydro-5H-cyclopenta[f][1,3]benzodioxol-6-amine (MDAI) [132741-81-2]
	5,6-Methylenedioxy-N-methyl-2-aminoindane, N-methyl-6,7-dihydro-5H-cyclopenta[f][1,3]benzodioxol-6-amine (MDMAI)
	5-Iodo-2-aminoindane, 5-Iodo-2,3-dihydro-1H-inden-2-amine (5-IAI) [132367-76-1]
	5-Methoxy-6-methyl-2-aminoindane, 5-Methoxy-6-methyl-2,3-dihydro-1H-inden-2-amine (MMAI)
	N-Ethyl-5-trifluoromethyl-2-aminoindane, N-Ethyl-5-(trifluoromethyl)-2,3-dihydro-1H-inden-2-amine (ETAI)
	N-Methyl-2-Aminoindane, 2,3-dihydro-N-methyl-1H-inden-2-amine (NM-2AI) [10408-85-2]
	((E)-1-(1-(2-Morpholino-1-yl)ethyl)indol-3-yl)-3,4,4-trimethylpent-2-en-1-one (A-796,260 isomer)
	1-(5-fluoropentyl)-3-(2-ethylbenzoyl)indole
	1-(5-fluoropentyl)-3-(2-methylbenzoyl)indole
	1-Butyl-1H-indol-3-yl)(naphthalen-1-yl)methanone (JWH-073) [208987-48-8]
	1-ethyl-1H-indol-3-yl)naphthalen-1-yl-methanone (JWH-071)
	1-Pentyl-1H-indazol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone
	1-pentyl-1H-indazol-3-yl)piperazin-1-yl)methanone
	1-Pentyl-1H-indol-3-yl)(pyridin-3-yl)methanone
	1-pentyl-1H-indol-3-yl)piperazin-1-yl)methanone
	(2,2,3,3-tetramethylcyclopropyl)[1-(4,4,4-trifluorobutyl)-1H-indol-3-yl]-methanone (XLR-12)
	1-(3-chloropentyl)-1H-indol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone (UR-144 N-(3-chloropentyl)analog)
	1-(4-fluorobenzyl)-1H-indol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone (FUB-144, FUB-UR-144)
	1-(5-fluoropentyl)-1H-indol-3-yl)tricyclo[3.3.1.1^{3,7}]dec-1-yl-methanone (5F-AB-001)
	1-(5-fluoropentyl)-N-(quinolin-8-yl)-1H-indazole-3-carboxamide, 5-fluoro-THJ (5F-THJ, 5F-THJ-018)
	1-Naphthalenyl(1-pentyl-1H-indazol-3-yl)-methanone, JWH-018 indazole analogue (THJ-018)
	1-(2-Morpholin-4-ylethyl)-1H-indol-3-yl)-(2,2,3,3-tetramethylcyclopropyl)methanone (A-796,260)
	1-(5-fluoropentyl)-N-(2-phenylpropan-2-yl)-1H-indazole-3-carboxamide, N-cumyl-1-(5-fluoropentyl)-1H-indazole-3-carboxamide (CUMYL-5FPINACA)
	1-(5-fluoropentyl)-N-(2-phenylpropan-2-yl)-1H-indole-3-carboxamide (CUMYL-5FPICA)
	1-(Cyclohexylmethyl)-2-[(4-ethoxyphenyl)methyl]-N,N-diethyl-1H-benzimidazol-5-carboxamide
	1-(Phenylmethyl)-1H-indole-3-carboxylic acid 8-quinolinyl ester, Quinolin-8-yl-1-(phenylmethyl)-1H-indole-3-carboxylate
	1-(Tetrahydropyran-4-ylmethyl)-1H-indol-3-yl)-(2,2,3,3-tetramethylcyclopropyl)methanone (A-834,735)
	1-Benzyl-N-(quinolin-8-yl)-1H-indazole-3-carboxamide
	1-Benzyl-N-(quinolin-8-yl)-1H-indole-3-carboxamide
	1-butyl-N-(2-phenylpropan-2-yl)-1H-indole-3-carboxamide (CUMYL-BICA)



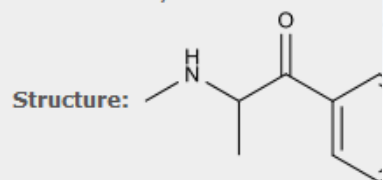
NPS Chemical Information

Methylone | 3,4-Methylenedioxy-N-meth

Details **Analytical Information** **Trend Data**

Names: Methylone | 3,4-Met
2-Methylamino-1-(3,
bk-MDMA | MDMC

IUPAC name: 1-(1,3-benzodioxol-5-yl)-2-methylamino-1-propanone
Substance group: Synthetic cathinones



CAS Number: 196028-79-5

InChI: InChI=1S/C11H13NO3
13)8-3-4-9-10(5-8)
7,12H,6H2,1-2H3

InChI Key: VKEQBMCQRDSRET-U

SMILES: CC(NC)C(=O)c1ccc2

Molecular Formula: C₁₁H₁₃NO₃

Molecular Weight: 207.2258 g/mol

Methylone | 3,4-Methylenedioxy-N-methcathinone

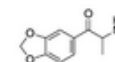
Details **Analytical Information** **Trend Data**

Spectra:

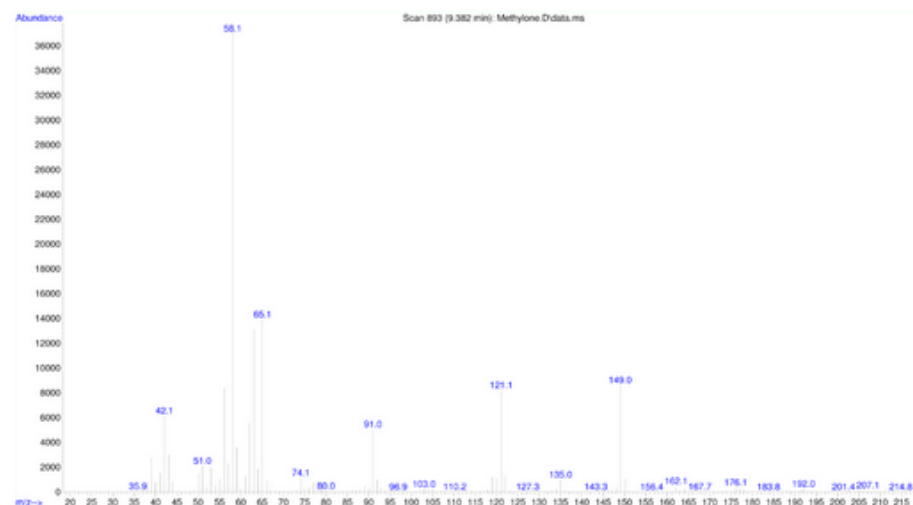
Note: All analytical spectra can only be used for reference purposes.



United Nations Office on Drugs and Crime



Methylone, 1-(1,3-benzodioxol-5-yl)-2-(methylamino)propan-1-one



GC-MS operating conditions

GC oven conditions: Column temp. initially set at 240 °C and held isothermal for 1 min immediately after injection and ramped to 330 °C at a rate of 6 °C/min with a final isotherm of 4 min

Column: TG-5QC, TG-5MS, DB-5MS or equivalent, 30 m x 0.25 mm i.d., 0.25 µm film thickness

Inlet: Mode: splitless (purge flow 30 ml/min at 0.3 min); Temp.: 250 °C; Carrier gas: Helium, 1 ml/min, constant flow; Injection volume: 1 µl

Detector: Ionisation mode: EI mode, 70 eV; Transfer line temp.: 280 °C; Ion source temp.: 225 °C

MS parameters: Solvent delay: 3 min; Scan mode; Scanning mass range: 30 – 600 amu at 2.17 scan/sec

Retention time: 9.4 mins.

Note: The above method represents a general screening method and may not be applicable to separate multiple components in a sample



Search Legal Responses

Legal Responses

To find more information on the various legal responses in place around the world, click [here](#).

Known year of first control:

-- all --
1978
1985
1986

(Ctrl+Click to multi-select)

Type of legislation:

-- all --
Analogue Control
Drug Laws/Individual Listing
Generic Legislation

(Ctrl+Click to multi-select)

Regions:

-- no region filter --
Africa
Americas
Asia

(Ctrl+Click to multi-select)

Countries:

ÅLAND ISLANDS
ALBANIA
ALGERIA
AMERICAN SAMOA

(Ctrl+Click to multi-select)

Selected countries:

AS, AI, AG, AR, AW, BS, BB, BZ, BM, BO, BR, CA, KY, CL, CO, CR, CU, DM, DO, EC, SV, FK, GF, GD, GP, GT, GY, HT, HN, JM, MQ, MX, MS, AN, NI, PA, PY, PE, PR, KN, LC, VC, SR, TT, TC, US, UY, VE, VG, VI

Page Size:

30

Filter

Clear Filter

Drug Laws/Individual Listing for ARGENTINA

Type of legislation: Drug Laws/Individual Listing

Legislation title: Decree 299 of 2010, Update of the list of narcotics and other chemical substances to be included under the Law No. 23.737

Known year of first control: 2010

Description:

In Argentina, the possession and traffic of narcotic and psychotropic substances is regulated by Law No. 23.737 [<http://www.mseq.qba.gov.ar/Investigaciones/DrogasIllicitas/ley%2023737.htm>] and the Decree No. 722 of 18 April 1991 [<http://infoleg.mecon.gov.ar/infolegInternet/resaltaranexos/5000-9999/6027/norma.htm>] where the list of controlled substances is contained. This Decree has been amended several times to update the list of controlled substances. In 2010, a number of new psychoactive substances were added to the list of controlled substances by Decree 299 of 2010 [<http://infoleg.mecon.gov.ar/infolegInternet/anexos/160000-164999/164826/norma.htm>]. Among the substances that were controlled are the phenethylamines 2C-I; 2C-T-2; 2C-T-7, PMMA, TMA-2, ketamine, the synthetic cathinone MPPP.

Links:

[Decree 299 of 2010](#)

If you have any further information or any amendments to the information provided on this page, please send an email to Global SMART (globalsmart@unodc.org).

[Return to list of legislation in ARGENTINA](#)

In most cases
the original law
is hyperlinked



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TIAFT
The International Association
of Forensic Toxicologists

Preview of the Tox-Portal

File Edit View Favorites Tools Help

UNODC Early Warning Advisory
Tox-Portal

Search.. EWA Testaccount Logout

Dashboard

Search Toxicology Report

New Toxicology Report

Help

Preferences

Administration

Dashboard

402
Total published reports

312
Reports last 12 months

190
Total reports with NPS

Gas Chromatography/Mass Spectrometry (GC/MS) (145)
 High Performance Liquid Chromatography (HPLC) (36)
 Gas Chromatography (GC) (36)
Top Screening Techniques

Inhalation (33)
Oral consumption (26)
Injection (26)
Top Methods of Administration

My Draft Incidents Submitted Assigned to me




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TIAFT
The International Association
of Forensic Toxicologists

Preview of the Tox-Portal



UNODC Early Warning Advisory - Tox-Portal

[Dashboard](#) [Search Toxicology Report](#) [+ New Toxicology Report](#) [About EWA](#) [Administration](#)

Unknown Unknown [Logout](#)

Substance

Dimethylcathinone (7)

N-Ethyl-ketamine (2)

AB-PICA (2)

4-Methoxy-alpha-pyrrolidinovalerophenone

5-EAPB (2)

4-Methoxybutyrfentanyl (1)

5,6-Methylenedioxy-2-aminoindane (1)

Deletion state

Not deleted (18)

Deleted (1)

Status

Not published (19)

[+ New Toxicology Report](#) [Export as Excel](#) [Print](#)

1 of 19 United Kingdom - occurred on 07/07/2017

id: 19

Patient: N/A (97)

Results

#	Substance or Metabolite	Matrix	Sampling Location	Concentration
1.	5-EAPB	Blood	Aorta	5%

2 of 19 United Kingdom - occurred on 15/06/2017

id: 18


Patient: Male (97)

Results

#	Substance or Metabolite	Matrix	Sampling Location	Concentration
1.	CP-47,497-C8	Blood/bile	Bladder/peripheral vein	10%

Challenge of capturing all relevant substances

“Found dead having smoked heroin the previous evening. Had access to prescription drugs and "Meow Meow".

 1 of 5 United Kingdom - occurred on 26/07/2016

Patient: Female (22)

Results

#	Substance or Metabolite	Matrix	Sampling Location
1.	Amfetamine	Blood	Femoral
2.	4-Methylethcathinone	Urine	N/A
3.	4-Methylmethcathinone	Urine	N/A
4.	Morphine	Blood	Femoral

Challenge of identifying the contributory substance

2 Substance 4-Methylethcathinone

Substance Details

Substance or metabolite(s) identified*

Matrix*

Sampling location (*if post-mortem*)*

Concentration (*e.g. mg/ml*)

Urine

N/A

N/A

Analytical Method(s)

Primary Screening

Identification*

Confirmation*

High Performance Liquid Chromatography (HPLC) ×

Select.. -

Liquid chromatography/tandem mass spectrometry

Verification Details

Means of Verification*

Route of Administration (*where available*)

Relative/probable contribution of the drug to the outcome of the event

Reference Standard

N/A

Present but non-contributory (low)

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