

# „PARTY DRUGS“ TESTING ON THE DANCE FLOOR: EQUIP-

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## Introduction

Since 1998 the mobile lab unit of the Office of the Cantonal Pharmacist (Health & Social Welfare Department, State of Berne, Switzerland) is testing so called „Party Drugs“ on the dance floor. At more than 220 events the team has tested over 4700 samples in cooperation with „Streetwork Zurich“, „Contact Berne“, „Suchthilfe Region Basel“ and „Prémiere ligne“ Geneva.

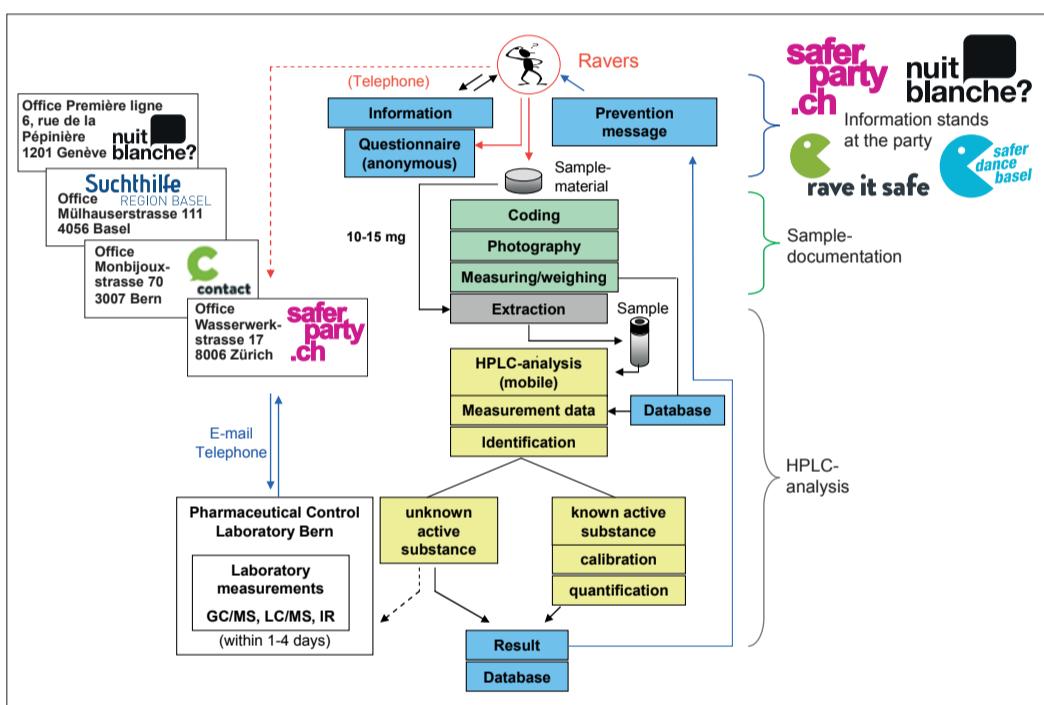
The mobile lab consists of four custom made subunits mounted in steel framed racks on wheels, one for weighing and documentation, one for sample preparation and two with the equipment for chemical analysis (HPLC-DAD).

The lab is operated by two experienced technicians. Before analysis the interested customer is asked by the lab crew to fill out a questionnaire concerning information about the sample; thereafter every sample is digitally documented and characterized by physical appearance (form, weight, dimensions etc.).



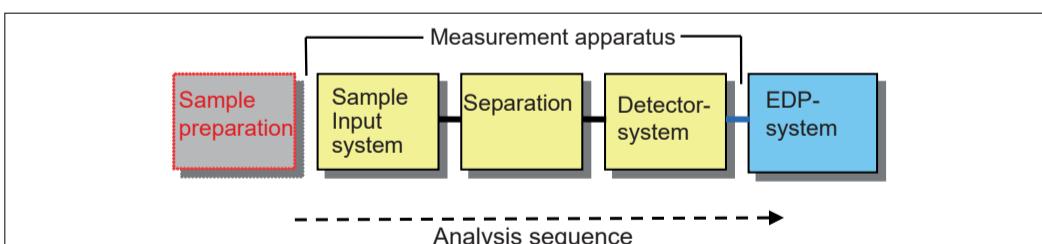
## Flow diagram

Collaboration between the prevention-teams and the laboratory



## Chemical analysis requirements

Chemical analysis of “Party Drugs” gives one of three possible results. The findings may be a single active agent, several active agents, or indeed no active agent. Reliable analysis of the substances in a sample cannot be achieved by means of a simple “quick test” and is only possible by the use of a complex measurement chain, as shown in the following basic diagram:



This arrangement applies to most modern chemical-analysis measurement systems.

### Sample preparation:

Due to the very sensitive analytical methods, only a representative part of the sample is used for further analysis. Sample preparation is quick and effective. The material is pulverised in a mortar and dissolved in methanol with the aid of an ultrasonic extractor.

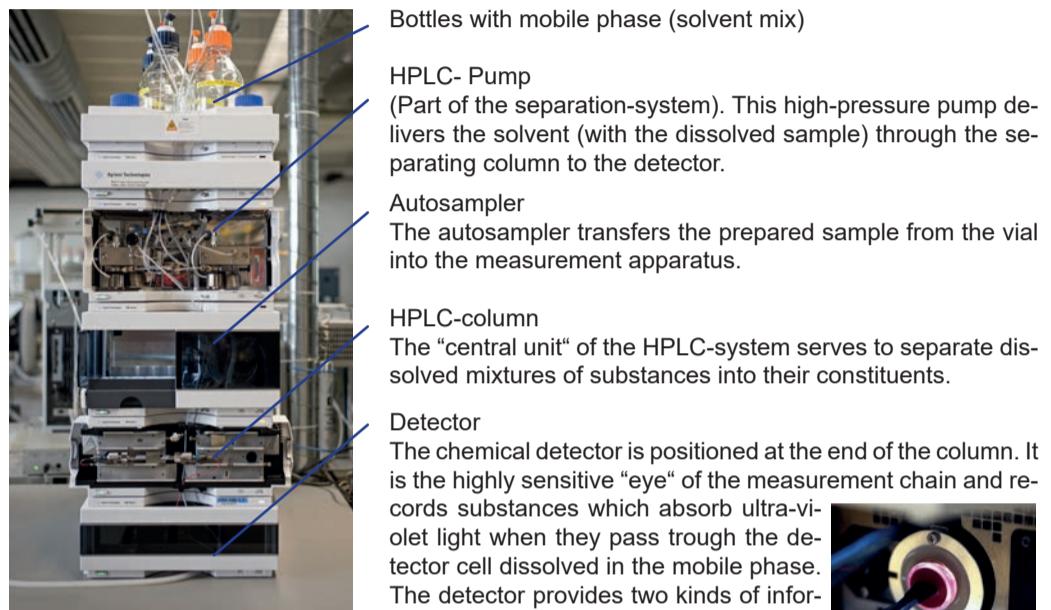


An internal standard is added as control. In most cases this extract still contains insoluble components, which must be filtered off before analysis. The clear sample solution obtained is transferred into a sample vial.



## HPLC (measurement apparatus)

For chemical analysis of party drugs separation and detection of different constituents of mixtures (active ingredients and fillers) is necessary. With the mobile lab we are using High Performance Liquid Chromatography (HPLC) for the separation process. Our computer controlled HPLC-systems are equipped with DAD/UV-Vis Spectrometers (Diode Array Detector).



their identity and their quantity. The whole process is continuously monitored with a computer system. Measurement signals are converted into graphical displays appearing on the computer screen. The two most important displays are the chromatogram and the UV-spectrum. The chromatogram is a representation of the separation process. The UV-spectrum is a characteristic constant for a particular substance.

### Analytical method (new version, introduction from june 2019)

#### HPLC-System

Autosampler: HP-1100, Typ G1313A  
Pump: HP-1100, Binary Pump, Typ G1312A  
Detector: HP-1100, UV-Vis DAD-Detector, Typ G1315B

Instrument control & integration: ChemStation for LC 3D software

HPLC- Conditions: Spherisorb 80-3 ODS-1 (Waters)

Stationary phase: 125 x 4.0 mm

Column dimension: 1.002 ml

Hold-up volume: Gradient

Elutiontype: 8,50 g ortho-phosphoric acid 85%  
+ 560 µl hexylamine  
+ purified water ad 1000 ml

Eluent A: 4,25 g ortho-phosphoric acid 85%  
+ 280 µl hexylamine  
+ 45,75 g purified water  
+ 351 g acetonitrile

#### Gradient program

Time in min	% A	%B
0.00 – 0.10	95	5
0.10 – 6.00	95 - 44	5 - 56
6.00 – 9.00	44	56
9.00 – 9.33	44 - 95	56- 5
9.33 – 11.33	95	5

#### Measuring parameters:

Flowrate:	1.5 ml/min
Pressure:	190 bar
Injection volume:	2.0 µl
Column temperature:	40 °C
Detection:	UV 198 nm
Signal-Range:	190-400 nm

## Report

Analytical results are available within about 12 minutes. The computer prints the results of the analysis as a report. The report consists of 3 parts: The header, the chromatogram, and the results (including identification and quantification)

### Header:

The header contains details like: File-name, sample-number, date of analysis, name of method etc.

### Chromatogram:

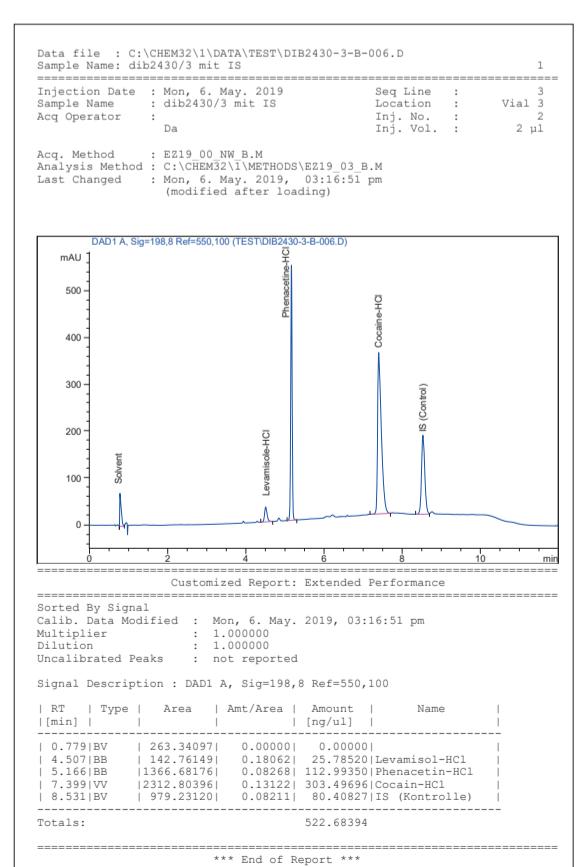
The chromatogram is a graphic visualisation of the separation process. Detected substances appear as “Peaks”.

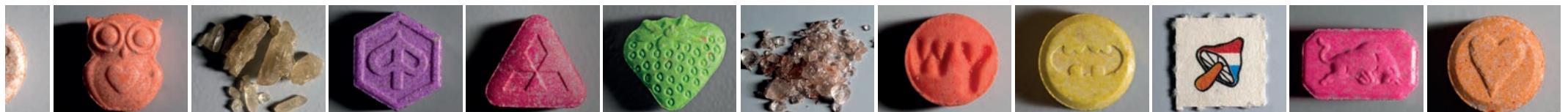
### Results (identification and quantification):

The system compares the area-value of an integrated Peak with the corresponding calibration of the active substance in the specified methods.

This automatic process gives us a precise quantification.

Additionally it is possible to get the UV-spectra of the “Peak” and compare it with our specific UV- Spectra-library.





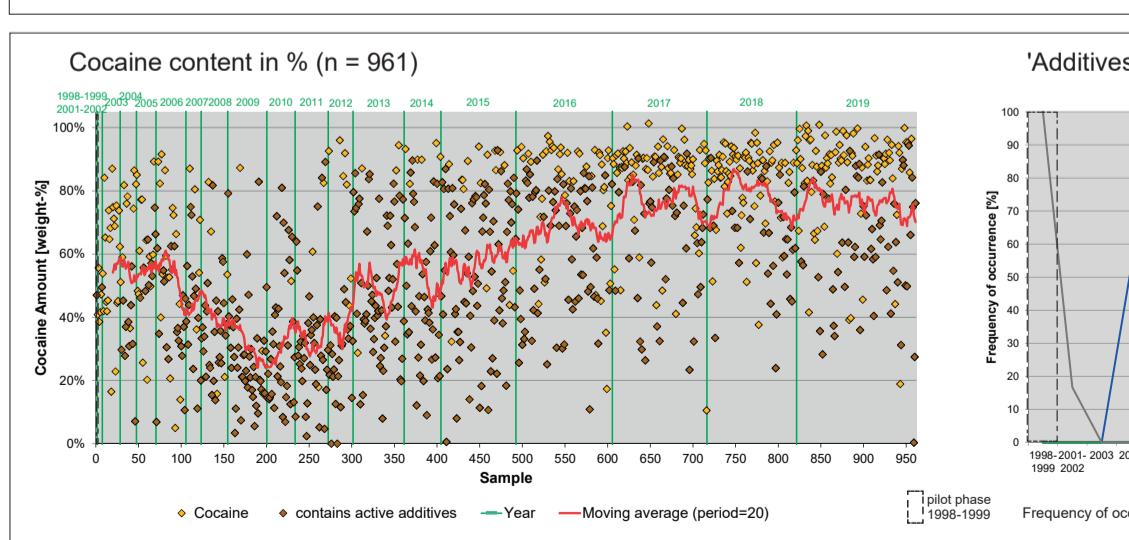
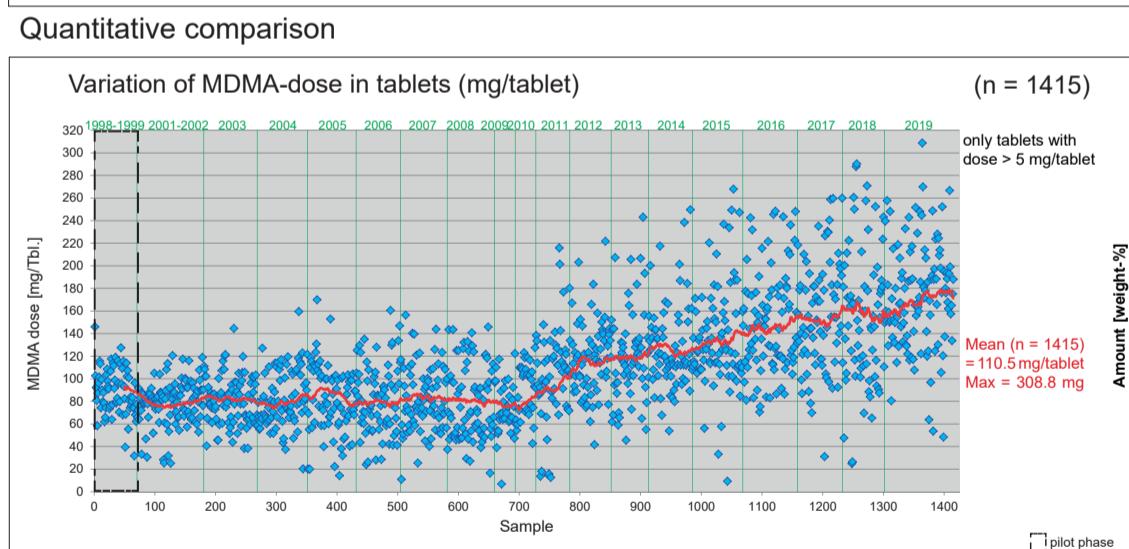
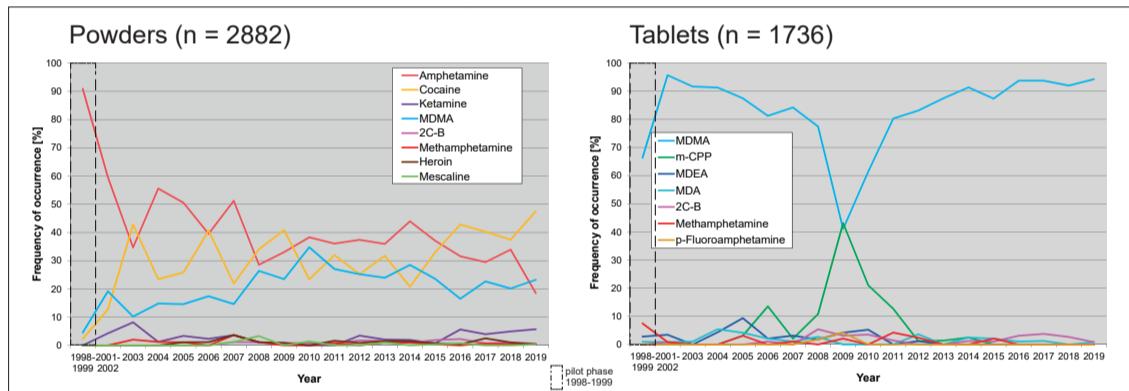
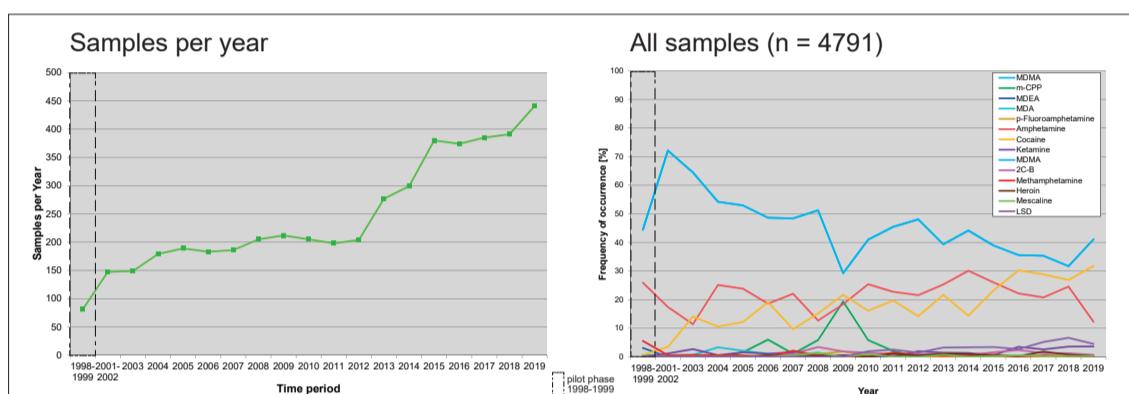
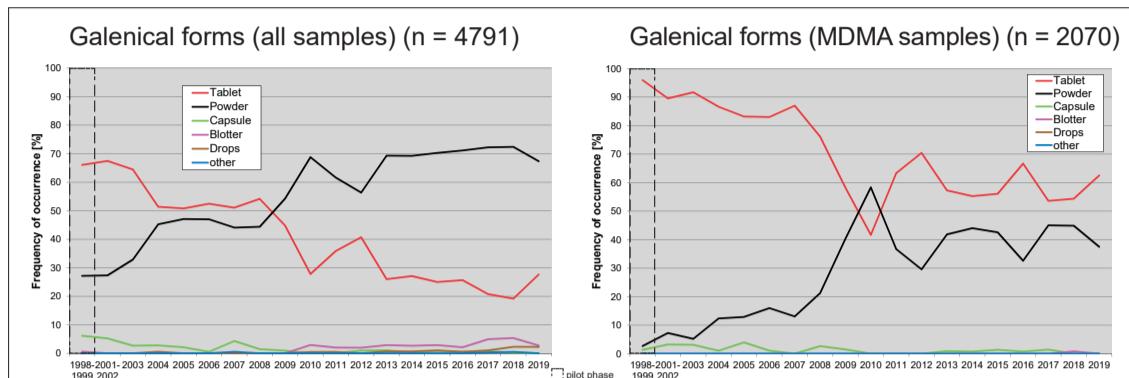
# -MENT, METHODS AND RESULTS

Hans-Jörg Helmlin, Samuel Steiner

Update: 31.12.2019

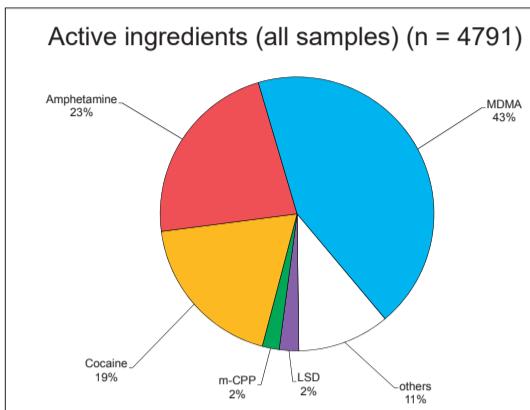
## Results

### Frequency of occurrence

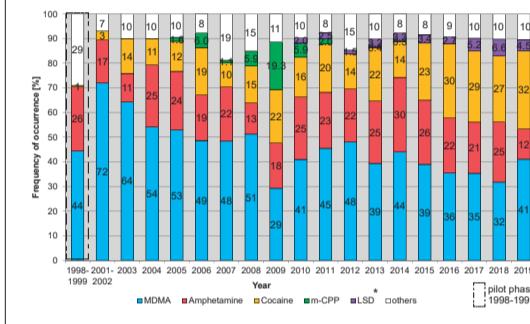


Chemical substances detected	Count	Percentage
MDMA	2108	44.00
Caffeine	1113	23.23
Amphetamine	1088	22.22
Cocaine	967	20.18
Levamisole	363	7.58
Phenacetin	290	6.05
LSD	120	2.59
m-CPP	96	2.00
Ketamine	88	1.84
no active ingredient	86	1.80
Lidocaine	72	1.50
Phencyclidone	64	1.34
Amphetamine synthesis by-product	60	1.25
2C-B	56	1.17
Paracetamol	48	1.00
Unknown	46	0.98
MDA	42	0.88
Domperidone	41	0.86
Methamphetamine	41	0.86
MDEA	37	0.77
Iso-LSD	35	0.73
Heroin	30	0.63
Noradrenaline	29	0.61
Creatine	23	0.48
Acetylsalicylic acid	22	0.46
Metaraminol	22	0.46
MDMA synthesis by-product related substance	19	0.40
3,4-Methylenedioxypheophenyl-2-propanol	17	0.35
Mescaline	16	0.33
2C-H	15	0.31
Hydroxyzine	14	0.29
4-Chloroamphetamine	13	0.27
Ephedrine	13	0.27
Procaine	13	0.27
Baclofene	12	0.25
Tetracaine	12	0.25
4-Hydroxyamphetamine	11	0.23
Buflomedil	10	0.21
N,N-Dimethylamphetamine	9	0.19
Salicylic acid	9	0.19
Mephedrone	8	0.17
4-Chloroamphetamine	8	0.17
Atropine (Hyoscyamine)	6	0.13
Methylenedipropylamine	6	0.13
Dimethyltryptamine	5	0.10
Phenylethylamine	5	0.10
TFMP	5	0.10
Vitamin B6	5	0.10
3-Methylnantholin (3-MMC)	4	0.08
Benzylpiperazine (A2)	4	0.08
DMT	4	0.08
DOA	4	0.08
MDA	4	0.08
Mescaline (bk-MDMA)	3	0.06
MDP4	3	0.06
N-Methyl-3,4-Methylenedioxylbenzylamir	3	0.06
Xylophane	3	0.06
2C-E	2	0.04
4-Chloroamphetamine	2	0.04
4-Chloroamphetamine (4-MEC)	2	0.04
Butylone (bk-MBDB)	2	0.04
Ciprolpold	2	0.04
Clorazepate	2	0.04
Ethchlorophylline	2	0.04
Fumarsure	2	0.04
MDMB	2	0.04
MDMB-CHMICA	2	0.04
Pentedrone	2	0.04
Phenethylamine	2	0.04
pHET	2	0.04
Pseudoeuphoria	2	0.04
Succrose octacetate	2	0.04
THC	2	0.04
Triglyceride	2	0.04
Tryptophan	2	0.04
1-Benzyl-4-Methylpiperizine (MBZP)	1	0.02
1H-Pyrazole	1	0.02
25B-NBOMe	1	0.02
25C-NBOMe	1	0.02
2C-D	1	0.02
2C-3	1	0.02
2C-1	1	0.02
2C-2	1	0.02
2-Fluor-Ketamine	1	0.02
2-Fluoromethamphetamine (2-FMA)	1	0.02
3-Fluoromethamphetamine (3-FPM)	1	0.02
3-Me-PCP	1	0.02
4-Chloroamphetamine	1	0.02
4-Fluoromethamphetamine	1	0.02
4-Hydroxyamphetamine	1	0.02
4-HO-MEP	1	0.02
4-HO-MPT	1	0.02
4-Methylmethamphetamine	1	0.02
4-Methylmethoxyamine	1	0.02
4-Methylmethoxypyrene	1	0.02
4-Methylmethothrone	1	0.02
5-Me-O-MPT	1	0.02
ALD-52	1	0.02
Alprazolam	1	0.02
Axonotillin	1	0.02
Benzoxic acid	1	0.02
Bisabolol	1	0.02
Carbamazepine	1	0.02
Clozapine	1	0.02
Desmethylketamine	1	0.02
Desmethylmorphinan (DXM)	1	0.02
Dimethyl terephthalate	1	0.02
Diphenidol	1	0.02
Doxylamine	1	0.02
Erdosteine	1	0.02
Entonox	1	0.02
Ethacridine	1	0.02
Ethambutol	1	0.02
Ethylamphetamine	1	0.02
Felphedrine	1	0.02
Gamma-hydroxybutyrate	1	0.02
GHB	1	0.02
Isoniazid	1	0.02
Leucylamphetamin	1	0.02
LSA	1	0.02
MDOPH	1	0.02
Melanin	1	0.02
Medicament	1	0.02
Melatonin	1	0.02
Mescaline	1	0.02
Melatansione	1	0.02
Methadoline	1	0.02
Methaqualone	1	0.02
Methylthiomethamphetamine (MTA)	1	0.02
Mix	1	0.02
N-Methyltyramine	1	0.02
Oxazepam	1	0.02
Phenobital	1	0.02
Piperazine	1	0.02
Pineatamine	1	0.02
Propantheline	1	0.02
Proprianol	1	0.02
Scopolamine	1	0.02
Sedentary derivative	1	0.02
Sildenafil	1	0.02
Tadalafil	1	0.02
Theophylline	1	0.02

### Qualitative results



### Active ingredients (evolution time based)



### Capsules (n = 61)

