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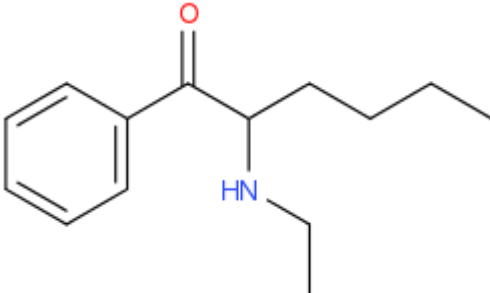
ANALYTICAL REPORT¹

N-ethylhexedrone (C₁₄H₂₁NO)

2-(ethylamino)-1-phenylhexan-1-one

Remark – other NPS detected: **none**

Sample ID:	1503-16 ; case reference 233-108/2016; sample label:32
Sample description:	powder - white-pale
Sample type:	collected /NGO-collected
Date of sample receipt (M/D/Y):	1/6/2016
Date of entry (M/D/Y) into NFL database:	2/10/2016
Report updates (if any) will be published here:	http://www.policija.si/apps/nfl_response_web/seznam.php

Substance identified - structure ² (base form)	
Systematic name	2-(ethylamino)-1-phenylhexan-1-one
Other names	Ethyl-hexedrone, HEXEN, HEX EN, NEH
Formula (per base form)	C ₁₄ H ₂₁ NO
M _w (g/mol)	219,33
Salt form/anions detected	HCl, traces of bromides
StdInChIKey	CWNKMHJETKEBCA-UHFFFAOYSA-N
Compound Class	Cathinones
Other NPS detected	none
Add.info (purity..)	pure by GC, HPLC-TOF, NMR

¹ Sample was collected and GC-MS screened in the frame of EU co-funded project I-SEE (JUST/2013/ISEC/DRUGS/AG/6426) and the compound structure was further verified in the frame of the RESPONSE project (JUST/2013/ISEC/DRUGS/AG/6413). Both projects run with the financial support of the Prevention of and fight against crime Programme of the European Union. The contents of this report are the sole responsibility of the National Forensic Laboratory and can in no way be taken to reflect the views of the European Commission.

² Created by OPSIN free tool: <http://opsin.ch.cam.ac.uk/> DOI: 10.1021/ci100384d

Report updates

date	comments (explanation)

Instrumental methods (if applied) in NFL

1. GC-MS (Agilent): GC-method is RT locked to tetracosane (RT=9.53 min). Injection volume 1 ml and split mode (1:50). Injector temperature: 280 °C. Chromatographic separation: on column HP1-MS (100% dimethylpolysiloxane), length 30 m, internal diameter 0.25 mm, film thickness 0.25 mm. Carrier gas He: flow-rate 1.2 ml/min. GC oven program: 170 °C for 1 min, followed by heating up to 293 °C at a rate of 18 °C/min, hold for 6.1 min, then heating at 50 °C/min up to 325 °C and finally 6.1 min isothermal. MSD source EI = 70 eV. GC-MS transfer line T= 235 °C, source and quadrupole temperatures 280 °C and 180 °C, respectively. Scan range m/z scan range: from 50 (40) to 550 amu.

2. HPLC-TOF (Agilent): 6230B TOF with Agilent 1260 Infinity HPLC with binary pump, column: Zorbax Eclipse XDB-C18, 50 x 4.6 mm, 1.8 micron. Mobile phases (A) 0.1% formic acid and 1mM ammonium formate in water; (B) 0.1% formic acid in methanol (B). Gradient: starting at 5% B, changing to 40% B over 4 min, then to 70% over 2 min and in 5 min to 100%, hold 1 min and back to 5%, equilibration for 1.7 min. The flow rate: 1.0 ml/min; Injection volume 1 µl. MS parameters: 2GHz, Extended Dynamic range mode to a maximum of 1700 amu, acquisition rate 1.30 spectra/sec. Sample ionisation: by Agilent Jet Stream technology (Dual AJS ESI). Ion source: positive ion scan mode with mass scanning from 82 to 1000 amu. Other TOF parameters: drying gas (N2) and sheath temperature 325 °C; drying gas flow rate 6 l/min; sheath gas flow rate 8 l/min; nebulizer 25 psig; Vcap. 4000 V; nozzle 2000 V; skimmer 65 V; fragmentor 175 V and Octopole RF 750 V.

3. FTIR-ATR (Perkin Elmer): scan range 4000-400 cm⁻¹; resolution 4cm⁻¹

4. GC- (MS)-IR condensed phase (GC-MS (Agilent) & IR (Spectra analyses-Danny)

GC-method: Injection volume 1 ml and split mode (1:5). Injector temperature 280 °C. Chromatographic separation as above **(1)**. Split MS : IR = 1:9.

MSD source EI = 70 eV. GC-MS transfer line T= 235 °C, source and quadrupole temperatures 280 °C and 180 °C, respectively. Scan range m/z scan range: from 50 (40) to 550 amu.

IR (condensed phase): IR scan range 4000 to 650, resolution 4 cm⁻¹.

5. IC (anions) (Thermo Scientific, Dionex ICS 2100), Column: IonPac AS19, 2 x 250mm; Eluent: 10mM from 0 to 10 min, 10-58 mM from 10 to 40min; Flow rate: 0.25 ml/min; Temperature: 30°C; Suppressor: AERS 500 2mm, suppressor current 13mA; Inj. Volume: 25 µl

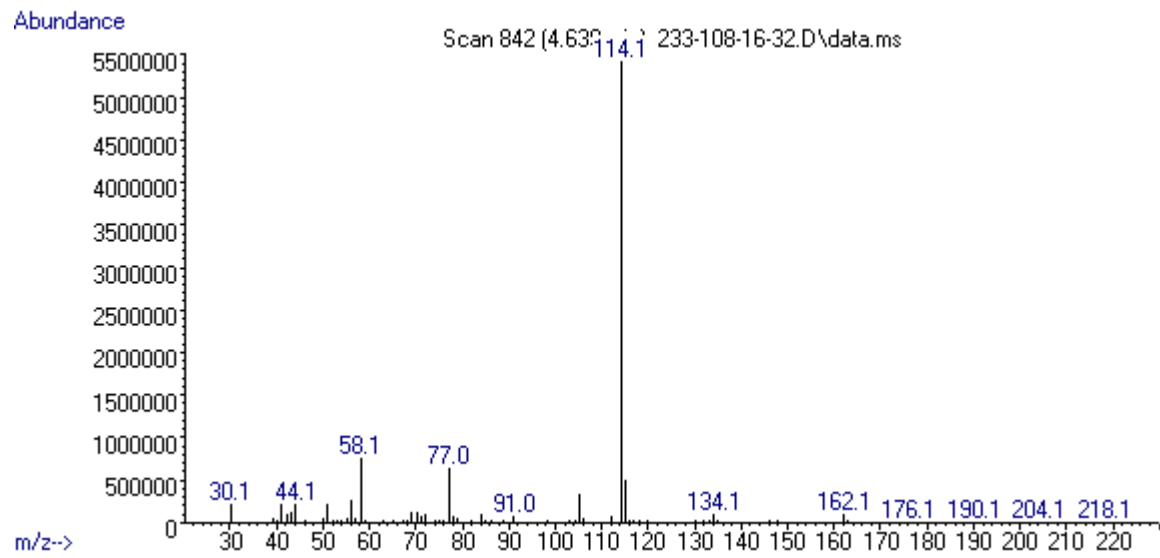
Supporting information

Solubility in	result/remark
CH ₂ Cl ₂	partially
MeOH	soluble
H ₂ O	soluble

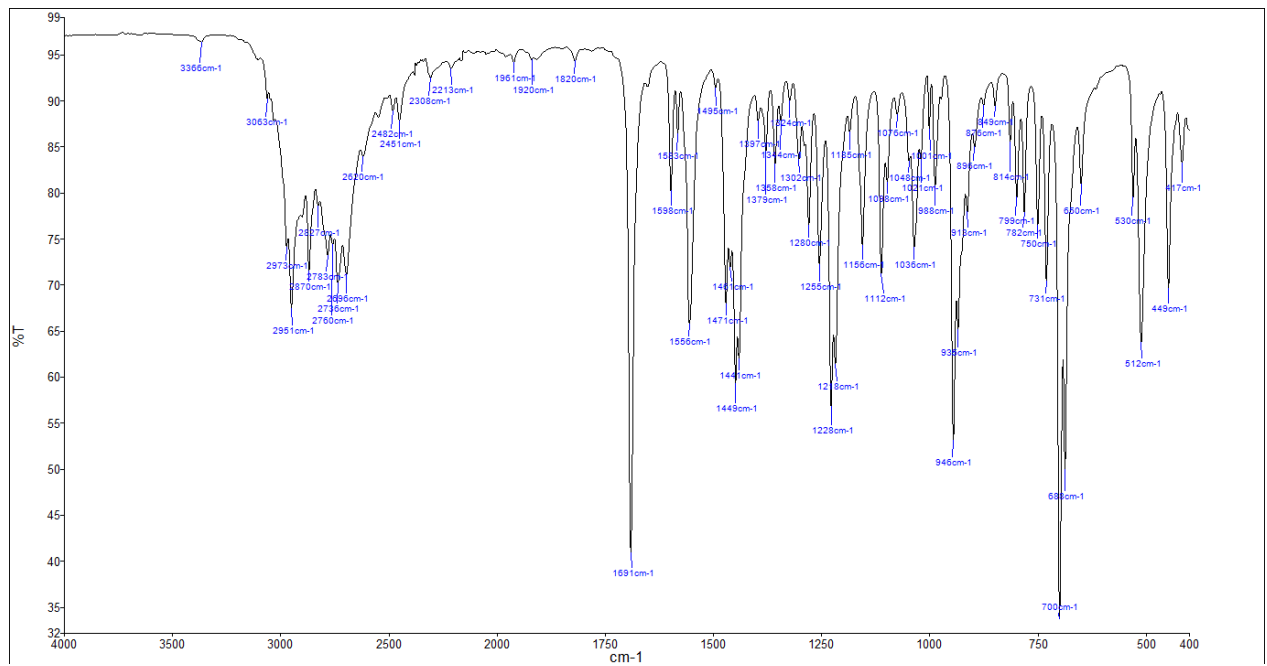
Analytical technique:	applied	remarks
GC-MS (EI ionization)	+	NFL GC-RT (min): 4,65 BP(1): 114; BP(2): 58, BP(3) :77,
HPLC-TOF	+	Exact mass (theoretical): 219,1623; measured value Δppm:-1,4; formula:C ₁₄ H ₂₁ NO
FTIR-ATR	+	direct measurement (sample as received)
FTIR (condensed phase) always as base form	-	
IC (anions)	+	graph not enclosed
NMR (in FKKT)	+	
validation		
other		

ANALYTICAL RESULTS

MS (EI)



FTIR-ATR - direct measurement (sample as received)



TOF REPORT

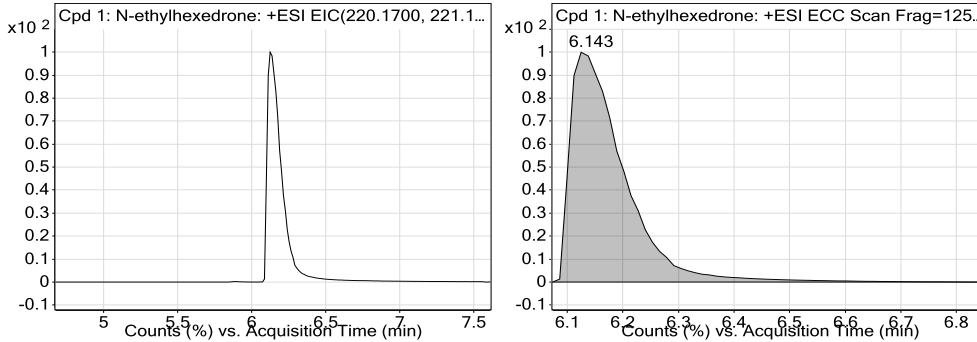
Data File	233-108-16-vzorec 32.d	Sample Name	vzorec 32
Sample Type	Sample	Position	P1-E6
Instrument Name	6230B TOF LC-MS	User Name	
Acq Method	general-1512015-XDB-C18-ESI-poz.m	Acquired Time	1/15/2016 9:11:11 AM
IRM Calibration Status	Success	DA Method	Drugs_NFL.m
Comment	ekstrakcija v MeOH		

Compound Table

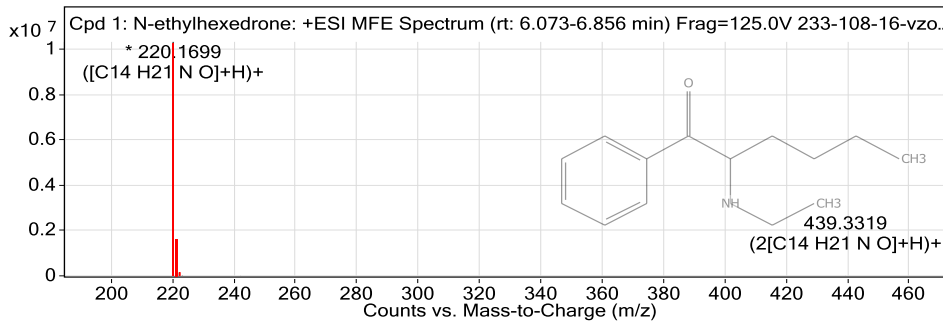
Label	Compound Name	MFG Formula	Obs. RT	Obs. Mass
Cpd 1: N-ethylhexedrone	N-ethylhexedrone	C14 H21 N O	6.143	219.1626

Name	Obs. m/z	Obs. RT	Obs. Mass	DB RT	DB Formula	DB Mass	DB Mass Error (ppm)
N-ethylhexedrone	220.1699	6.143	219.1626	6.14	C14 H21 N O	219.1623	-1.4

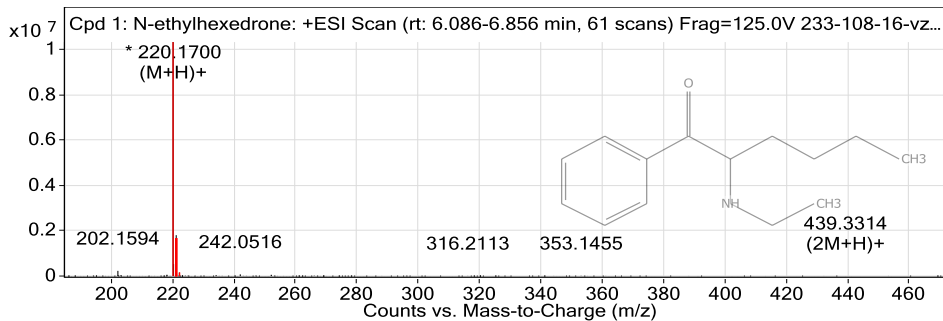
Compound Chromatograms



MFE MS Zoomed Spectrum



MS Zoomed Spectrum



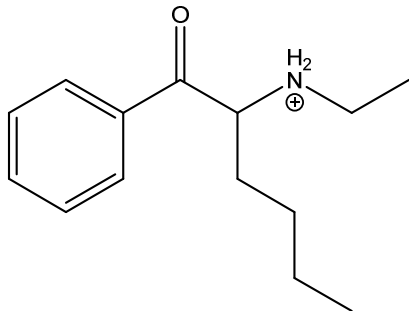
MS Spectrum Peak List

Obs. m/z	Charge	Abund	Formula	Ion/Isotope
220.1699	1	10344070	C14 H21 N O	(M+H)+
221.1731	1	1548135.77	C14 H21 N O	(M+H)+
222.1763	1	147617.28	C14 H21 N O	(M+H)+
223.1786	1	9624.81	C14 H21 N O	(M+H)+
242.1512	1	2481.56	C14 H21 N O	(M+Na)+
439.3319	1	1584.93	C14 H21 N O	(2M+H)+

--- End Of Report ---



REPORT

Sample ID:	233-108-16-32
Our notebook code:	P-233-108-16-32
NMR sample preparation:	15 mg dissolved in 0.7 mL DMSO- <i>d</i> ₆
NMR experiments:	¹ H, ¹³ C, ¹ H- ¹ H <i>gs</i> -COSY, ¹ H- ¹³ C <i>gs</i> -HSQC, ¹ H- ¹³ C <i>gs</i> -HMBC, ¹ H- ¹⁵ N <i>gs</i> -HMBC.
Proposed structure:	
Chemical name:	N-ethyl-1-oxo-1-phenylhexan-2-aminium
Comments:	- Structure elucidation based on 1D and 2D NMR spectra - Sample is pure by NMR.
Supporting information:	Copies of ¹ H and ¹³ C NMR spectra
Author:	Prof. Dr. Janez Košmrlj, Doc. Dr. Krištof Kranjc
Date of report:	January 25, 2016

P-233-108-16-32

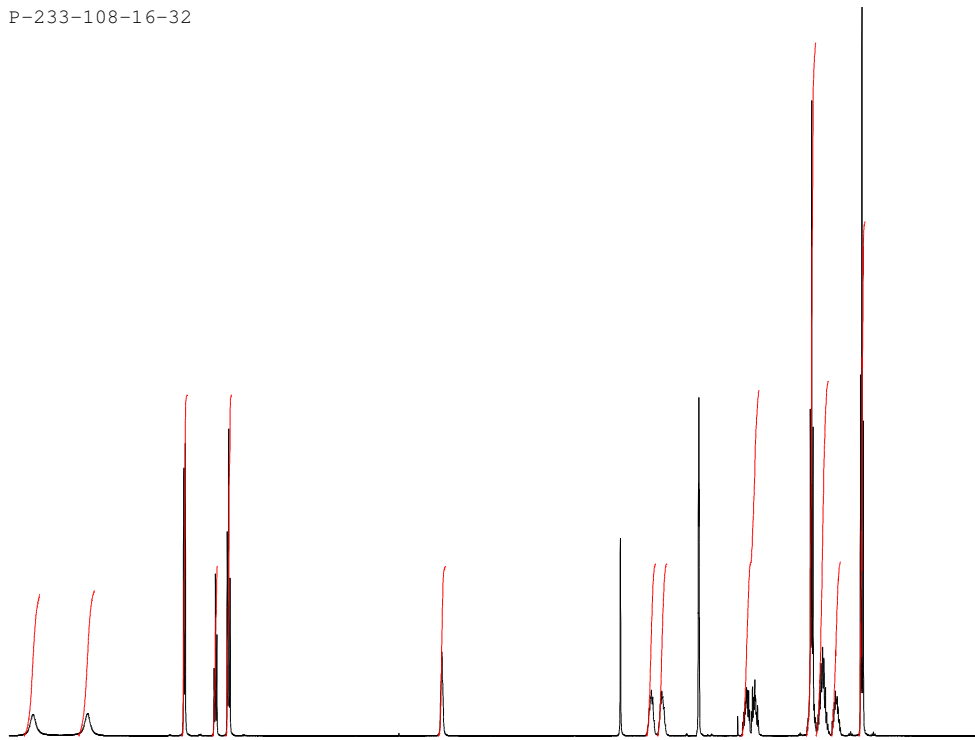


Current Data Parameters
NAME P-233-108-16-32
EXPNO 1
PROCNO 1

F2 - Acquisition Parameters
Date_ 20160124
Time 15.02
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT DMSO
NS 16
DS 2
SWH 10000.000 Hz
FIDRES 0.152588 Hz
AQ 3.2768500 sec
RG 57
DW 50.000 usec
DE 6.50 usec
TE 298.0 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 500.1330885 MHz
NUC1 1H
P1 8.90 usec
PLW1 26.00000000 W

F2 - Processing parameters
SI 65536
SF 500.1300000 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



9 8 7 6 5 4 3 2 1 ppm

0.83 0.85 2.00 1.00 2.00 0.99 1.01 1.01 2.03 4.06 2.08 1.02 3.01

P-233-108-16-32



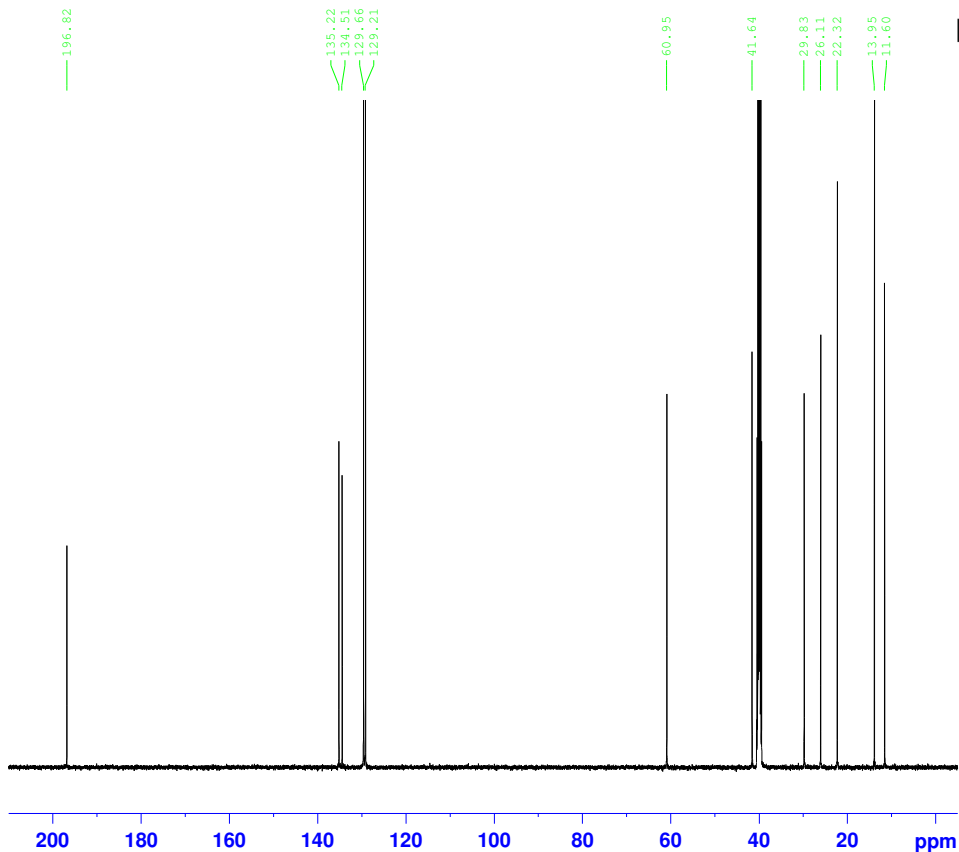
Current Data Parameters
NAME P-233-108-16-32
EXPNO 3
PROCNO 1

F2 - Acquisition Parameters
Date_ 20160124
Time 17.37
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT DMSO
NS 4096
DS 4
SWH 29761.904 Hz
FIDRES 0.454131 Hz
AQ 1.1010048 sec
RG 2050
DW 16.800 usec
DE 6.50 usec
TE 298.0 K
D1 1.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 125.7703637 MHz
NUC1 13C
P1 9.00 usec
PLW1 122.00000000 W

===== CHANNEL f2 =====
SFO2 500.1320005 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 80.00 usec
PLW2 26.00000000 W
PLW12 0.32179001 W
PLW13 0.16186000 W

F2 - Processing parameters
SI 32768
SF 125.7577885 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



200 180 160 140 120 100 80 60 40 20 ppm