



NACIONALNI FORENZIČNI LABORATORIJ  
NATIONAL FORENSIC LABORATORY

Vodovodna 95  
1000 Ljubljana  
SLOVENIJA

T: +386 (0)1 428 44 93  
E: [nfl@policija.si](mailto:nfl@policija.si)  
[www.policija.si](http://www.policija.si)

## ANALYTICAL REPORT<sup>1</sup>

### Methoxypiperamide (C13H18N2O2) 1-(4-methoxybenzoyl)-4-methylpiperazine

Remark – other NPS detected: **none**

Sample ID:	1439-16
Sample description:	powder - white
Sample type:	collected /Institute of Forensic medicine, University Freiburg, Germany
Date of sample receipt (M/D/Y):	1/14/2016
Date of entry (M/D/Y) into NFL database:	8/16/2016
Report updates (if any) will be published here:	<a href="http://www.policija.si/apps/nfl_response_web/seznam.php">http://www.policija.si/apps/nfl_response_web/seznam.php</a>

Substance identified - structure <sup>2</sup> (base form)	
Systematic name	1-(4-methoxybenzoyl)-4-methylpiperazine
Other names	(4-Methoxyphenyl)(4-methylpiperazin-1-yl)methanone; Methoxypiperamide; MEXP, MeOP
Formula (per base form)	C13H18N2O2
M <sub>w</sub> (g/mol)	234,3
Salt form/anions detected	chloride
StdInChIKey	DWPVVZZGGGCRMM-UHFFFAOYSA-N
Compound Class	Piperazine derivates
Other NPS detected	none
Add.info (purity..)	pure by HPLC-TOF, GC-MS,

<sup>1</sup> This report has been produced with the financial support of the Prevention of and fight against crime Programme of the European Union (grant agreement number JUST/2013/ISEC/DRUGS/AG/6413). 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.

<sup>2</sup> 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 (9.258 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 µm. Carrier gas He: flow-rate 1.2 ml/min. GC oven program: 170 °C for 1 min, followed by heating up to 190 °C at rate 8 °C/min, then 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 quadropole temperatures 280 °C and 180 °C, respectively. Scan range m/z scan range: from 50 (30 until 6 min.) to 550 (300 until 6 min) 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<sup>-1</sup>; resolution 4cm<sup>-1</sup>

**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 quadropole temperatures 280 °C and 180 °C, respectively. Scan range m/z scan range: from 50 (30 until 6 min.) to 550 (300) amu.

IR (condensed (solid) phase): IR scan range 4000 to 650, resolution 4 cm<sup>-1</sup>.

**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

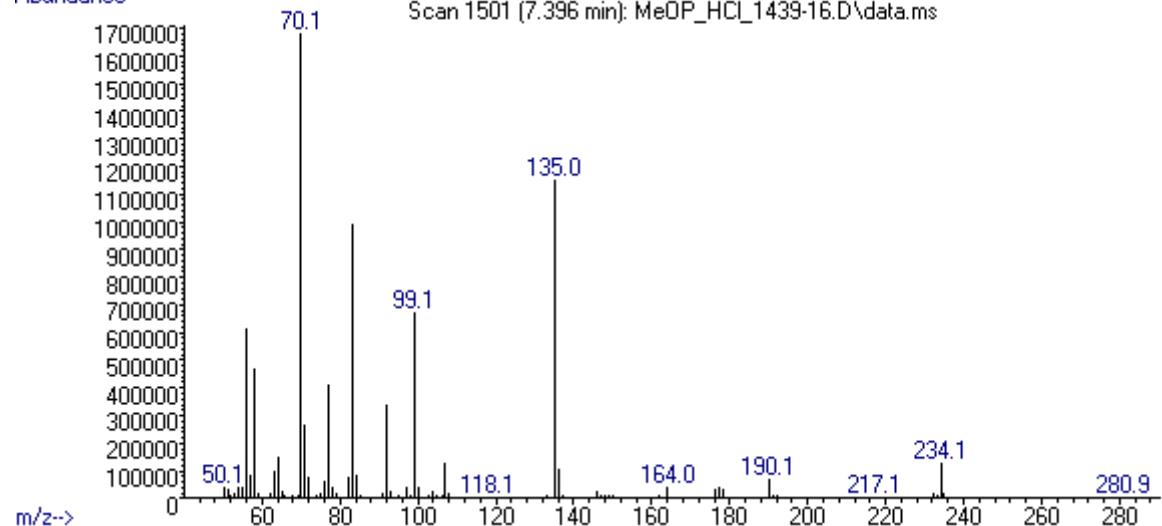
Solubility in	result/remark
CH <sub>2</sub> Cl <sub>2</sub>	soluble
MeOH	soluble
H <sub>2</sub> O	soluble

Analytical technique:	applied	remarks
GC-MS (EI ionization)	+	NFL GC-RT (min): 7,4 BP(1): 70; BP(2): 135,BP(3) :83,
HPLC-TOF	+	Exact mass (theoretical): 234,1368; measured value Δppm:-2,36; formula:C13H18N2O2
FTIR-ATR	+	direct measurement (sample as received)
FTIR (condensed phase) always as base form	+	
IC (anions)	+	
NMR (in FKKT)	-	
validation		
other		MS spectrum consistent by spectrum in SWGDRUG library (QM =98)

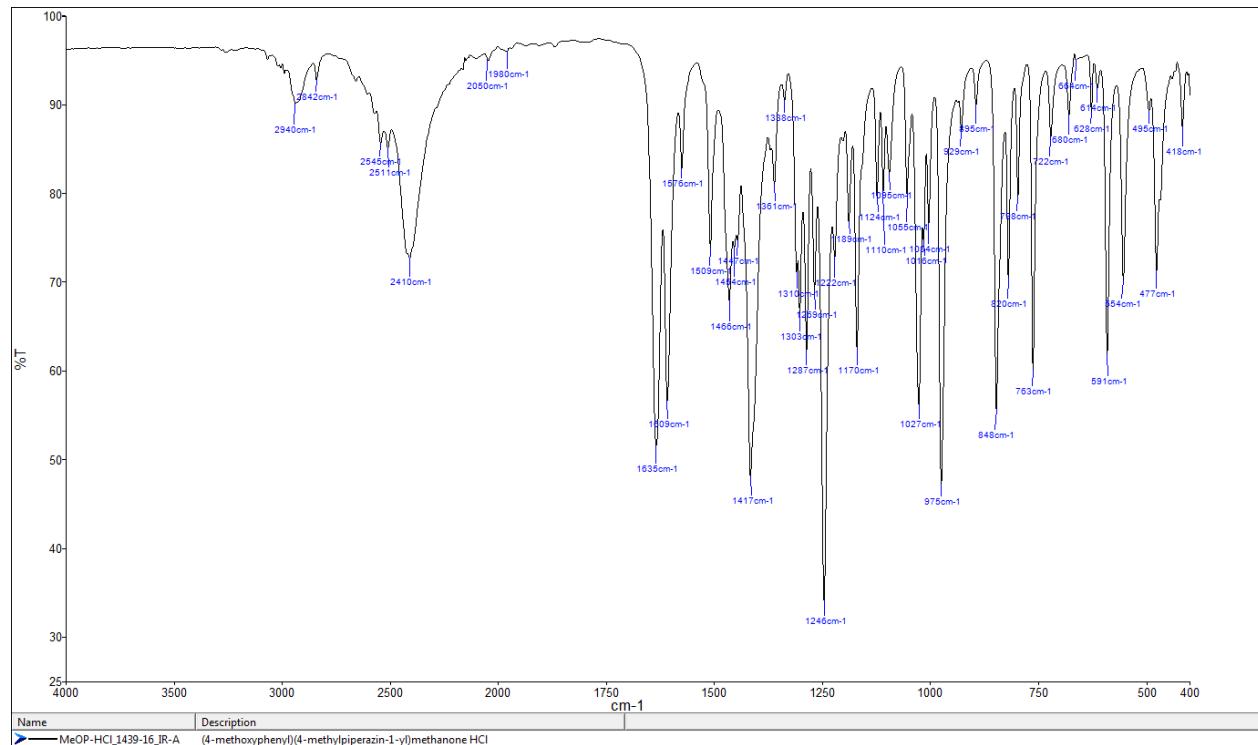
## ANALYTICAL RESULTS

MS (EI)

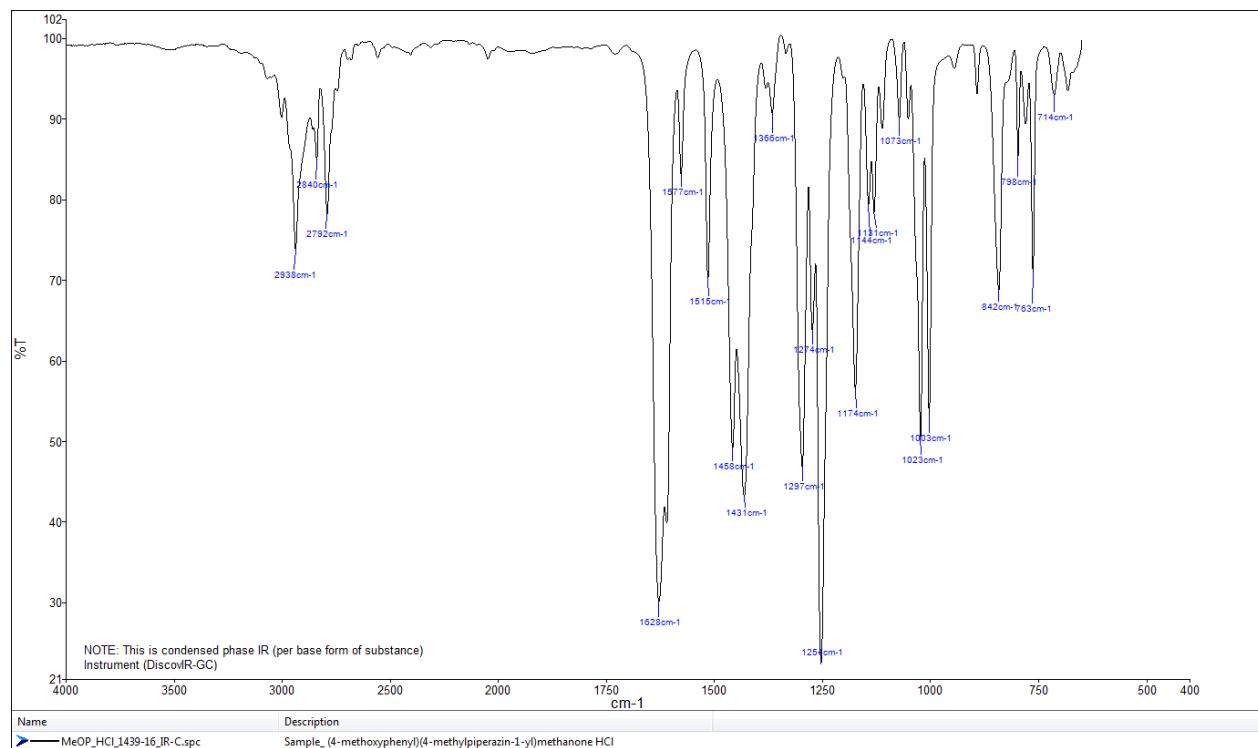
Abundance



### FTIR-ATR - direct measurement (sample as received)



### IR (condensed phase – after chromatographic separation)



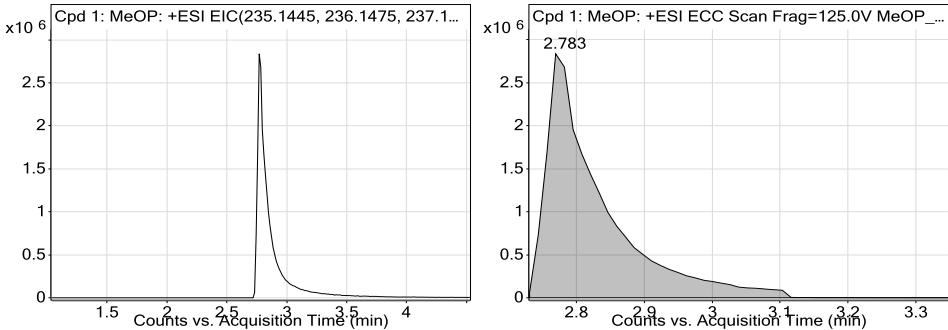
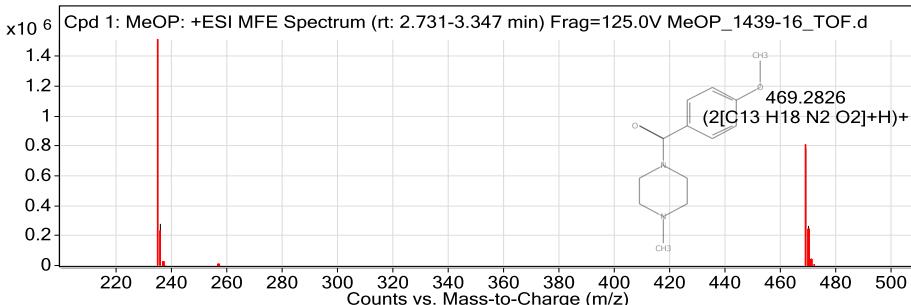
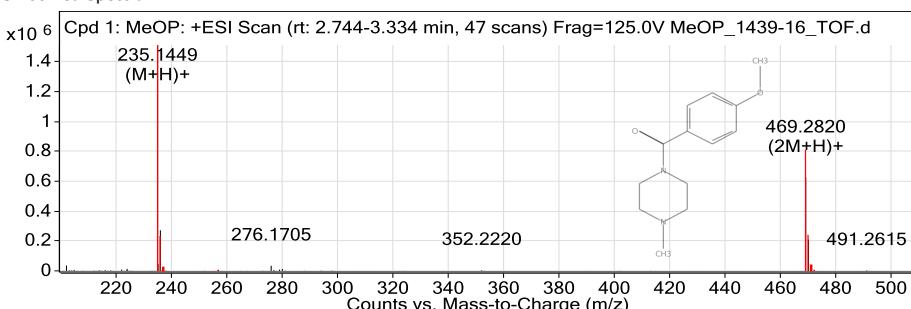
# TOF REPORT

<b>Data File</b>	MeOP_1439-16_TOF.d	<b>Sample Name</b>	ID_1439-16
<b>Sample Type</b>	Sample	<b>Position</b>	P1-E6
<b>Instrument Name</b>	6230B TOF LC-MS	<b>User Name</b>	TG
<b>Acq Method</b>	general-1512015-XDB-C18-ESI-pozi.pod.m	<b>Acquired Time</b>	2/23/2016 9:33:22 AM
<b>IRM Calibration Status</b>	Success	<b>DA Method</b>	Drugs_NFL.m
<b>Comment</b>	extract in MeOH		

**Compound Table**

Label	Compound Name	MFG Formula	Obs. RT	Obs. Mass
Cpd 1: MeOP	MeOP	C13 H18 N2 O2	2.783	234.1374

Name	Obs. m/z	Obs. RT	Obs. Mass	DB RT	DB Formula	DB Mass	DB Mass Error (ppm)
MeOP	235.1446	2.783	234.1374	2.78	C13 H18 N2 O2	234.1368	-2.36

**Compound Chromatograms**

**MFE MS Zoomed Spectrum**

**MS Zoomed Spectrum**

**MS Spectrum Peak List**

Obs. m/z	Charge	Abund	Formula	Ion/Isotope
235.1446	1	1511222.63	C13 H18 N2 O2	(M+H)+
236.148	1	278438.01	C13 H18 N2 O2	(M+H)+
237.1499	1	28690.33	C13 H18 N2 O2	(M+H)+
238.1525	1	1837.43	C13 H18 N2 O2	(M+H)+
257.1259	1	4291.71	C13 H18 N2 O2	(M+Na)+
258.1293	1	692.85	C13 H18 N2 O2	(M+Na)+
469.2826	1	783689.56	C13 H18 N2 O2	(2M+H)+
470.2848	1	264377.49	C13 H18 N2 O2	(2M+H)+
471.2863	1	46716.68	C13 H18 N2 O2	(2M+H)+
472.2889	1	5818.58	C13 H18 N2 O2	(2M+H)+

--- End Of Report ---

### Peak Integration Report

Sample Name:	MeOP_1439-16_IC	Inj. Vol.:	25,00
Injection Type:	Unknown	Dilution Factor:	1,0000
Program:	ANIONI	Operator:	kemija
Inj. Date / Time:	24-jun-2016 / 12:44	Run Time:	42,00

No.	Time min	Peak Name	Peak Type	Area $\mu\text{S}^*\text{min}$	Height $\mu\text{S}$	Amount mg/L
1,00	8,10	Chloride	BMB	26,84	126,35	n.a.
		TOTAL:		26,84	126,35	0,00

