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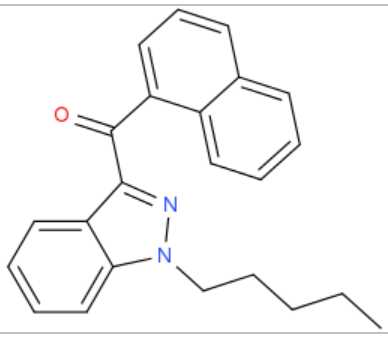
ANALYTICAL REPORT

THJ-018 (C23H22N2O)

3-(naphthalene-1-carbonyl)-1-pentyl-1H-indazole

Remark – other NPS detected:

Sample ID:	1298-15
Sample description:	powder
Sample type:	test purchase /RESPONSE -purchasing
Date of sample receipt (M/D/Y):	10/1/2015
Date of entry (M/D/Y) into NFL database:	11/6/2015
Report[1] 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	3-(naphthalene-1-carbonyl)-1-pentyl-1H-indazole
Other names	naphthalen-1yl-(1-pentyl-1H-indazol-3-yl)-methanone; JWH 018 indazole analog
Formula (per base form)	C23H22N2O
M_w (g/mol)	342,4
Salt form/anions detected	base
StdInChIKey (per base form)	VAKGBPSFDNTMDJ-UHFFFAOYSA-N
Other NPS detected	
Additional info (purity..)	HPLC-TOF possibly contains isomeric form of the same compound (app. 10%), NMR pure

[1] 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.

[2] Created by OPSIN free tool: http://opsin.ch.cam.ac.uk/ **DOI:** 10.1021/ci100384d

Report updates

date	comments (explanation)
January 15, 2018	Typing errors were corrected (molecular formula, MS BP2).

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 0C. Chromatographic separation: on column HP1-MS (100% dimethylpolysiloxane), length 30 m, internal diameter 0.25 mm, film thickens 0.25 µm. Carrier gas He: flow-rate 1.2 ml/min. GC oven program: 170 ^{0}C for 1 min, followed by heating up to 190 ^{0}C at rate 8 ^{0}C/min, then heating up to 293 0C at a rate of 18 ^{0}C/min, hold for 7.1 min, then heating at 50 ^{0}C/min up to 325 ^{0}C and finally 6.1 min isothermal. MSD source EI = 70 eV. GC-MS transfer line T= 235^{0}C, source and quadropole temperatures 280^{0}C and 180^{0}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-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 ^{0}C. Chromatographic separation as above **(1).**
Split MS : IR = 1: 9.
MSD source EI = 70 eV. GC-MS transfer line T= 235^{0}C, source and quadropole temperatures 280^{0}C and 180^{0}C, respectively. Scan range m/z scan range: from 50 (30 until 6 min.) to 550 (300) amu.
IR (condesed (solid) 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

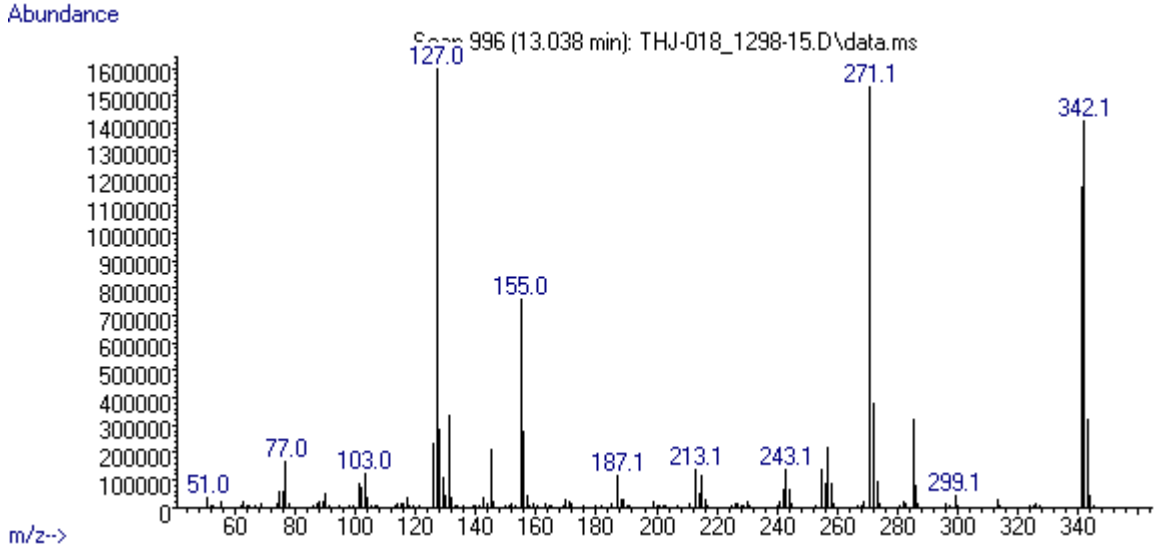
Supporting information

Solubility in	result/remark
CH_2Cl_2	soluble
MeOH	soluble
H_2O	

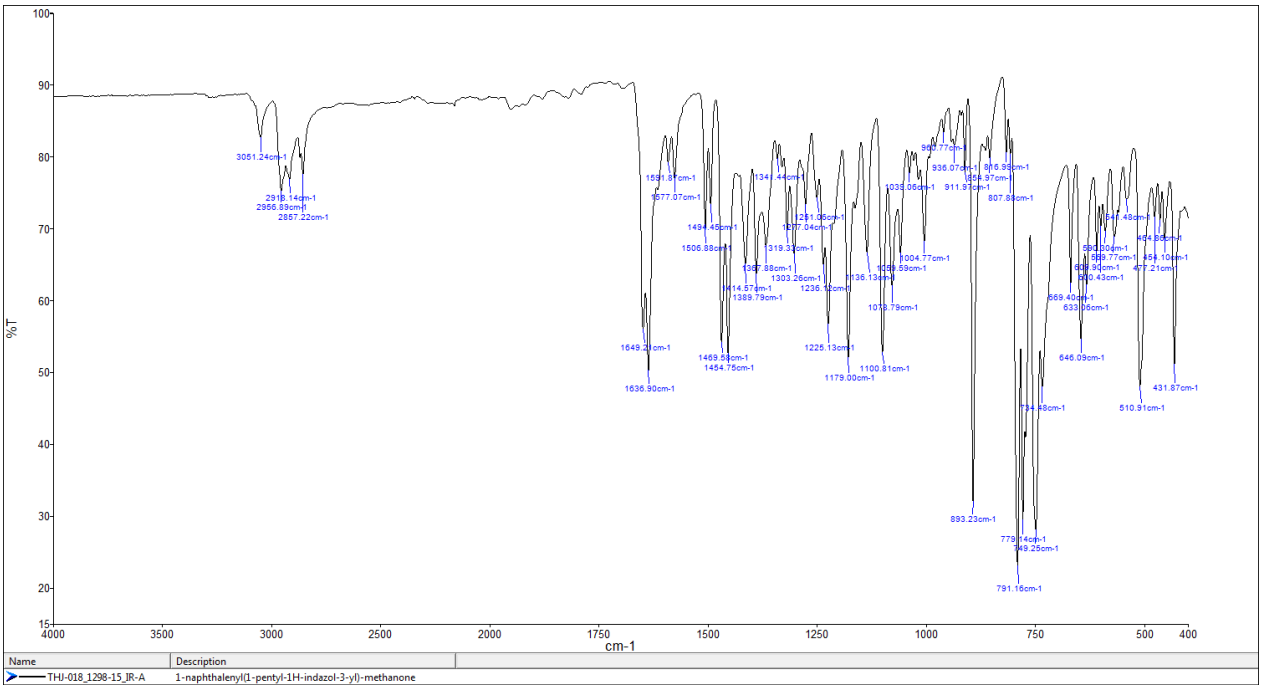
Analytical technique:	applied	remarks
GC-MS (EI ionization)	+	NFL GC-RT (min): 13,04 BP(1): 127; BP(2): 271,BP(3) :342,
HPLC-TOF	+	Exact mass (theoretical): 342,1732; measured value Δppm:-0,7; formula:C23H22N2O
FTIR-ATR	+	direct measurement (sample as received)
FTIR (condensed phase) always as base form	+	
IC (anions)	+	
NMR (in FKKT)	+	
validation		
other		

ANALYTICAL RESULTS

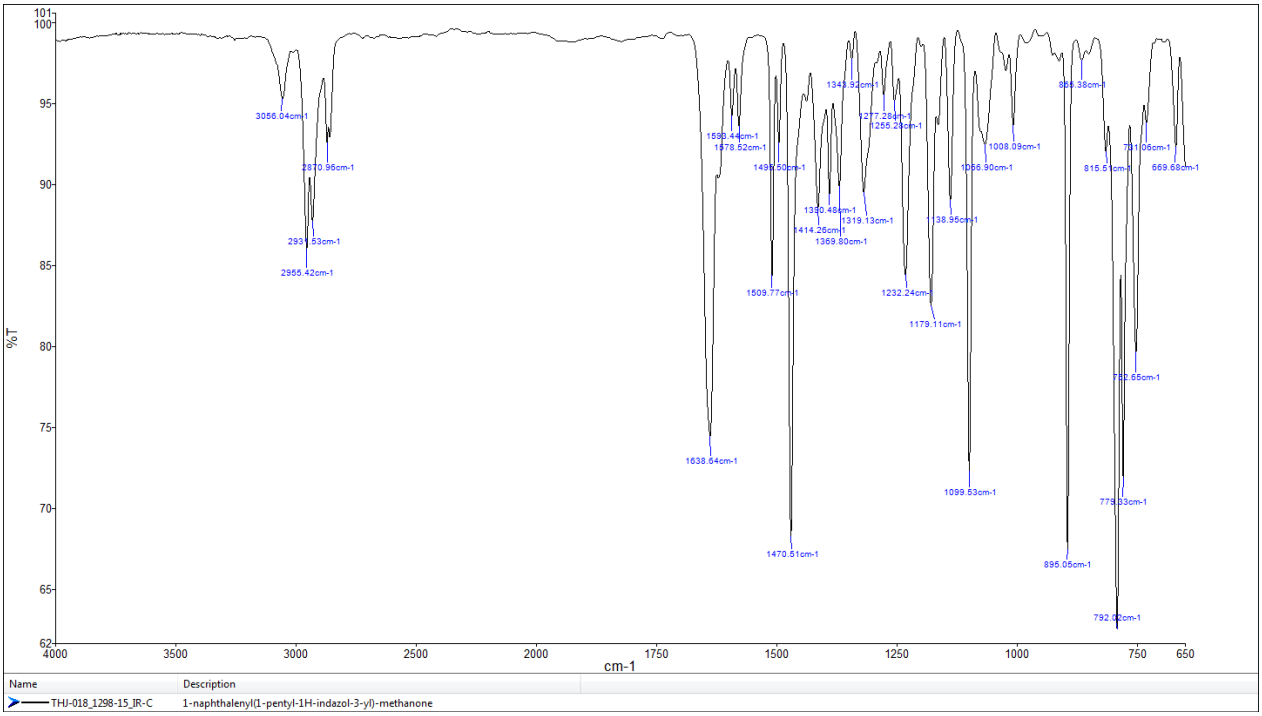
MS (EI)



FTIR-ATR - direct measurement (sample as received)



IR (condensed phase – after chromatographic separation)



TOF REPORT

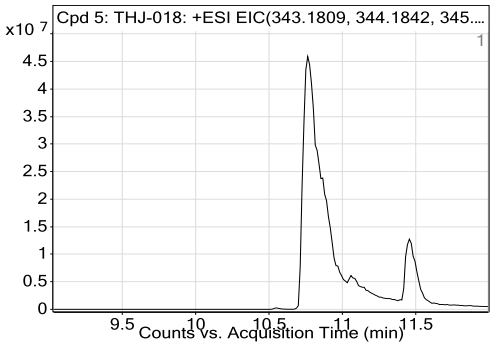
Data File	THJ-018_1298-15_TOF.d	**Sample Name**	THJ-018
Sample Type	Sample	**Position**	P1-B3
Instrument Name	6230B TOF LC-MS	**User Name**	TG
Acq Method	general-28052015-XDB-C18-ESI-poz.m	**Acquired Time**	10/23/2015 8:28:50 AM
IRM Calibration Status	Success	**DA Method**	Drugs_NFL.m
Comment	extract in MeOH		

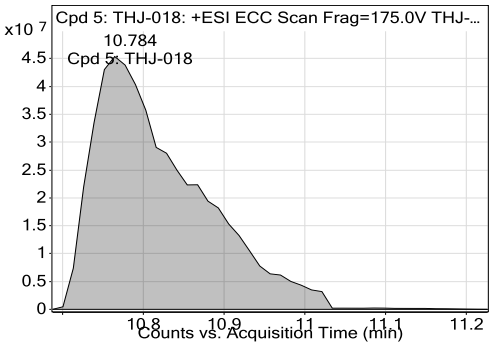
Compound Table

Label	Compound Name	MFG Formula	Obs. RT	Obs. Mass
Cpd 5: THJ-018	THJ-018	C23 H22 N2 O	10.784	342.1734

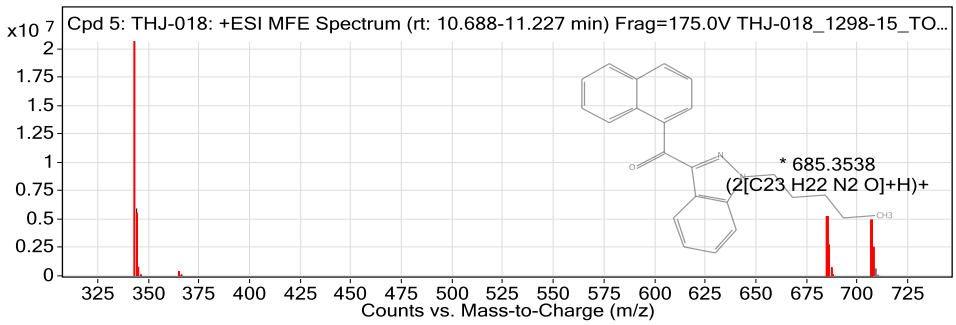
Name	*Obs. m/z*	Obs. RT	Obs. Mass	DB RT	DB Formula	DB Mass	DB Mass Error (ppm)
THJ-018	343.1808	10.784	342.1734	10.8	C23 H22 N2 O	342.1732	-0.7

Compound Chromatograms

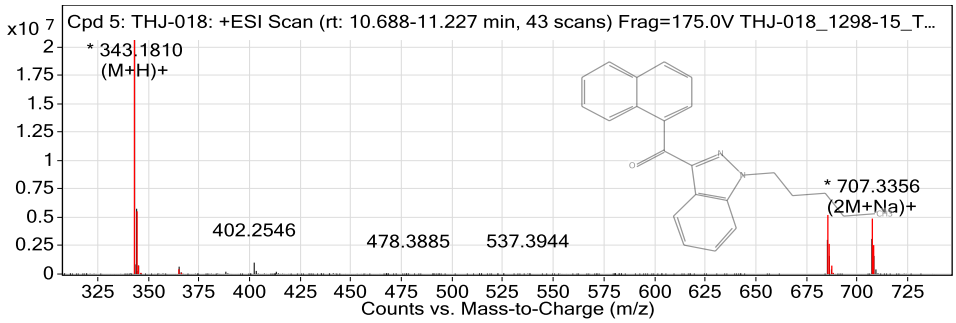




MFE MS Zoomed Spectrum



MS Zoomed Spectrum



MS Spectrum Peak List

Obs. m/z	Charge	Abund	Formula	Ion/Isotope
343.1808	1	20597552	C23 H22 N2 O	(M+H)+
344.1839	1	5931260.23	C23 H22 N2 O	(M+H)+
345.1877	1	798064.08	C23 H22 N2 O	(M+H)+
365.163	1	435592.59	C23 H22 N2 O	(M+Na)+
685.3538	1	5169631.5	C23 H22 N2 O	(2M+H)+
686.3571	1	2753829.5	C23 H22 N2 O	(2M+H)+
687.361	1	706741.33	C23 H22 N2 O	(2M+H)+
707.3358	1	4914007.5	C23 H22 N2 O	(2M+Na)+
708.3391	1	2555431.61	C23 H22 N2 O	(2M+Na)+
709.3429	1	658367.75	C23 H22 N2 O	(2M+Na)+

--- End Of Report ---

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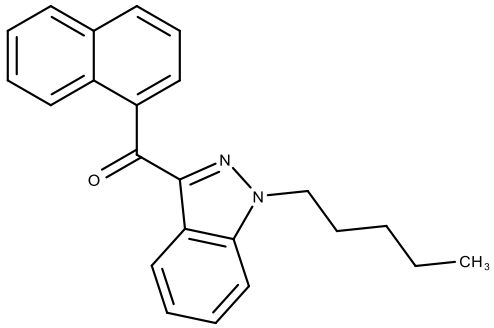
Co-funded by the Prevention of and Fight against Crime Programme of the European Union

REPORT

Sample ID:	**1298-15**
Our notebook code:	P-1298-15
NMR sample preparation:	15 mg dissolved in 0.7 mL DMSO-d_6
NMR experiments:	^{1}H, ^{13}C.
Proposed structure:	
Chemical name:	naphthalen-1-yl(1-pentyl-1H-indazol-3-yl)methanone
Comments:	- Structure elucidation based on 1D NMR spectra - Compound is pure by NMR, containing only a minor amount of impurities.
Supporting information:	Copies of ^{1}H and ^{13}C NMR spectra
Author:	Prof. Dr. Janez Košmrlj, Doc. Dr. Krištof Kranjc
Date of report:	December 8, 2015

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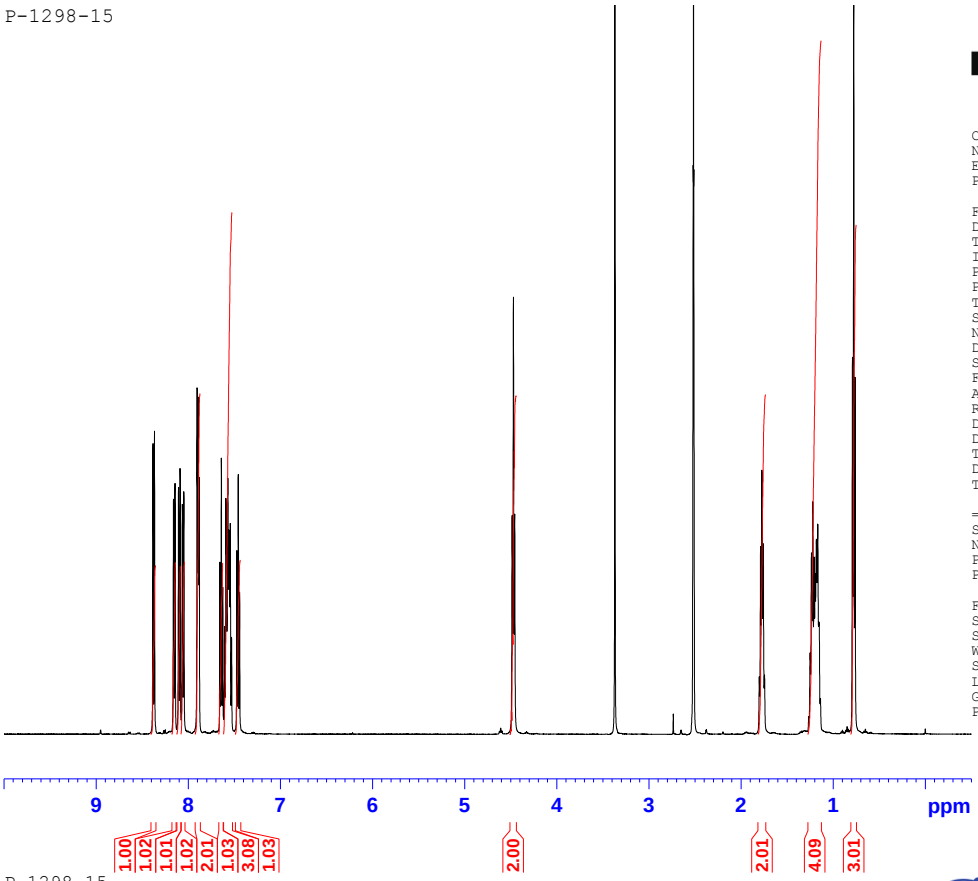
P-1298-15



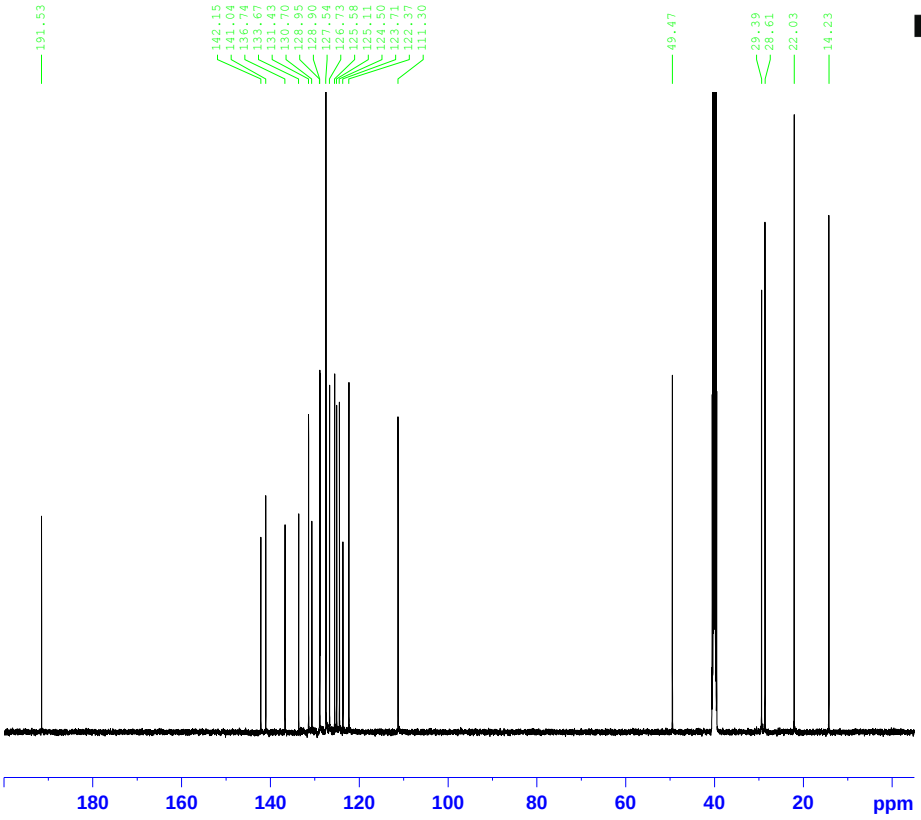
Current Data Parameters
NAME p-1298-15
EXPNO 1
PROCNO 1

F2 - Acquisition Parameters
Date_ 20151207
Time 22.36
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 64
DW 50.000 usec
DE 6.50 usec
TE 296.0 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 500.1299959 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

P-1298-15



Current Data Parameters
NAME P-1298-15
EXPNO 2
PROCNO 1

F2 - Acquisition Parameters
Date_ 20151208
Time 0.28
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT DMSO
NS 3072
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 296.0 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