



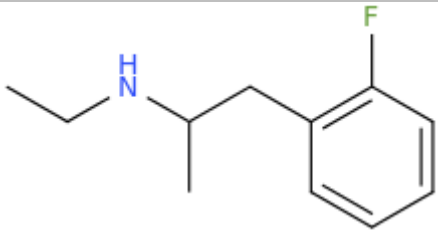
ANALYTICAL REPORT

2-FEA (C₁₁H₁₆FN)

ethyl[1-(2-fluorophenyl)propan-2-yl]amine

Remark – other NPS detected: **none**

Sample ID:	2006-18
Sample description:	powder
Sample type:	test purchase /NFL- purchasing
Date of sample receipt (DD/MM/YYYY):	25/10/2018
Date of entry (DD/MM/YYYY) into NFL database:	22/01/2019
Report updates (if any) will be published here:	http://www.policija.si/apps/nfl_response_web/seznam.php

Substance identified - structure ¹ (base form)	
Systematic name	ethyl[1-(2-fluorophenyl)propan-2-yl]amine
Other names	o-Fluoroethamphetamine; 2-Fluoro-ethyl Amphetamine
Formula (per base form)	C ₁₁ H ₁₆ FN
M _w (g/mol)	181.25
Salt form/anions detected	HCl
StdInChIKey (per base form)	NIWYLNFSJJLYHU-UHFFFAOYSA-N
Other NPS detected	none
Additional info (purity..)	>95% purity by ¹ H NMR

¹ 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 7.1 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⁻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 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⁻¹.

5. IC (anions) (Thermo Scientific, Dionex ICS 2100), Column: IonPac AS19, 2 x 250mm; Eluent: 10mM KOH 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

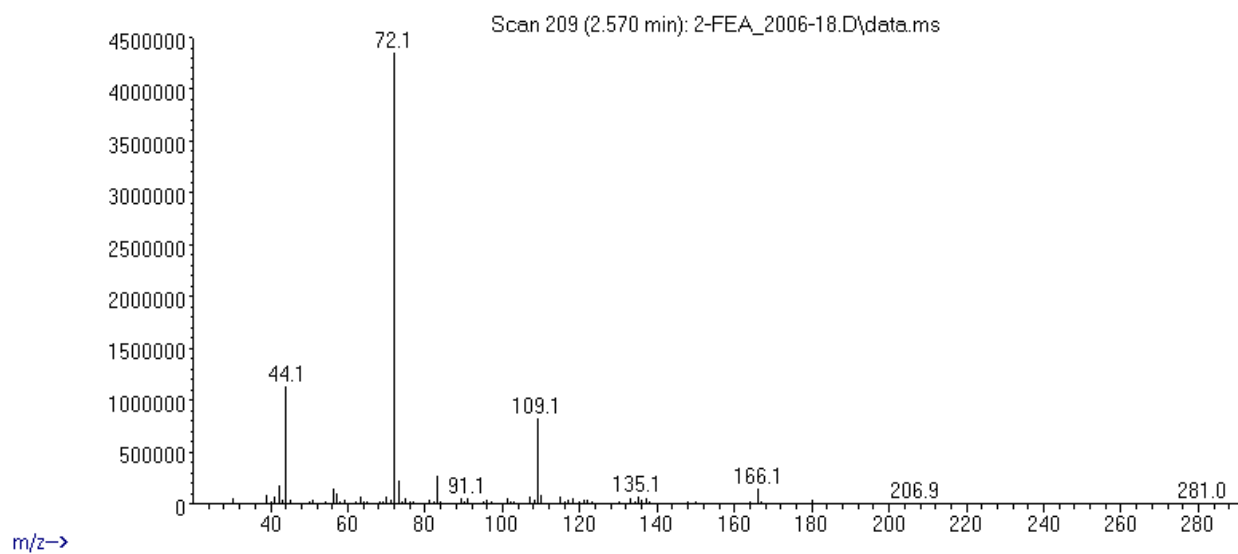
Solubility in	result/remark
CH ₂ Cl ₂	soluble
MeOH	soluble
H ₂ O	soluble

Analytical technique:	applied	remarks
GC-MS (EI ionization)	+	NFL GC-RT (min): 2.57 BP(1): 72; BP(2): 44,BP(3) :109,
HPLC-TOF	+	Exact mass (theoretical): 181.1267; measured value Δppm:-3.36; formula:C11H16FN
FTIR-ATR	+	direct measurement (sample as received)
FTIR (solid phase) always as base form	+	
IC (anions)	+	
NMR (in FKKT)	+	
validation		
other		

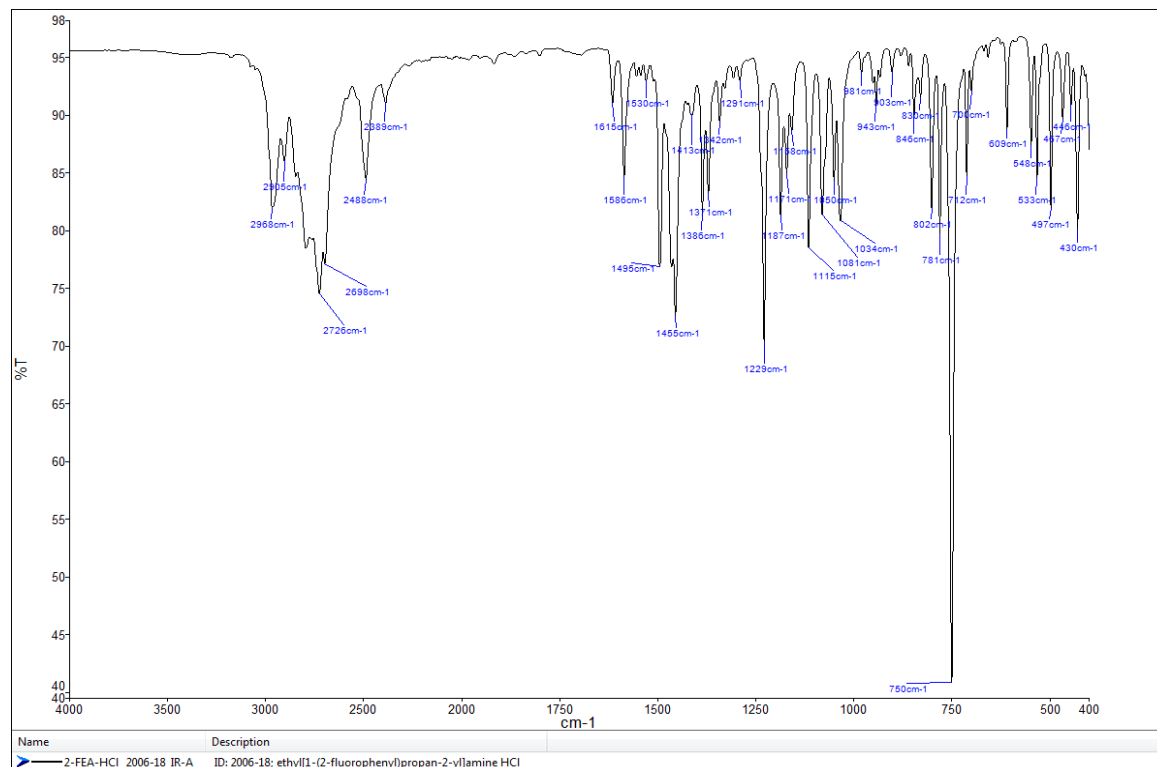
ANALYTICAL RESULTS

MS (EI)

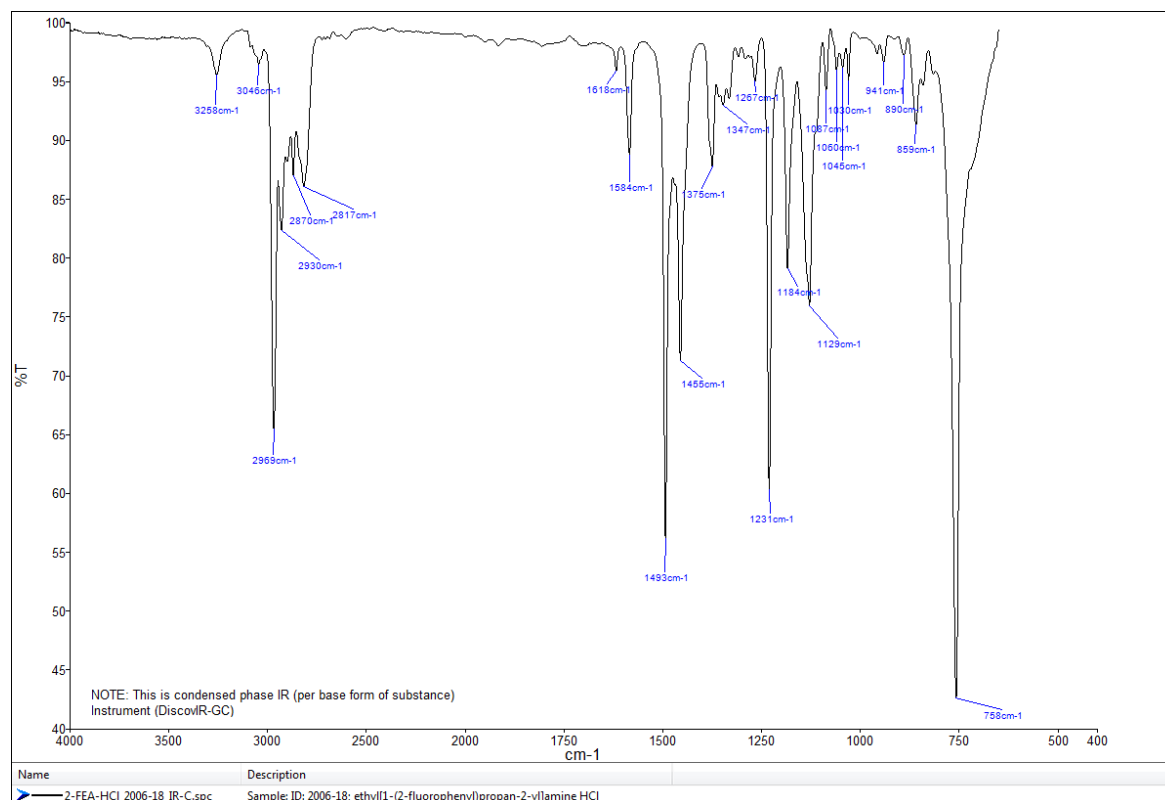
Abundance



FTIR-ATR - direct measurement (sample as received)



IR (solid phase – after chromatographic separation)



TOF REPORT

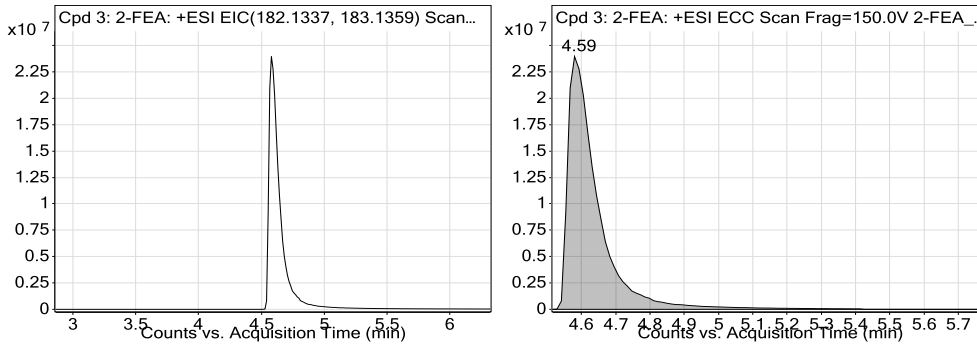
Data File	2-FEA_2006_18.d	Sample Name	ID 2006-18
Sample Type	Sample	Position	P2-C5
Instrument Name	6230B TOF LC-MS	User Name	TG
Acq Method	general-19_10_2018-XDB-C18-ESI+.m	Acquired Time	11/1/2018 5:27:30 PM
IRM Calibration Status	Success	DA Method	a-Drugs_NFL.m
Comment	MeOH		

Compound Table

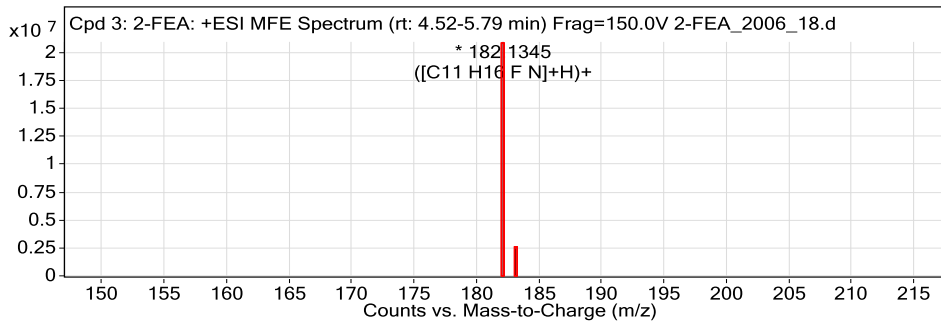
Label	Compound Name	MFG Formula	Obs. RT	Obs. Mass
Cpd 3: 2-FEA	2-FEA	C11 H16 F N	4.59	181.1273

Name	Obs. m/z	Obs. RT	Obs. Mass	DB RT	DB Formula	DB Mass	DB Mass Error (ppm)
2-FEA	182.1345	4.59	181.1273	4.59	C11 H16 F N	181.1267	-3.36

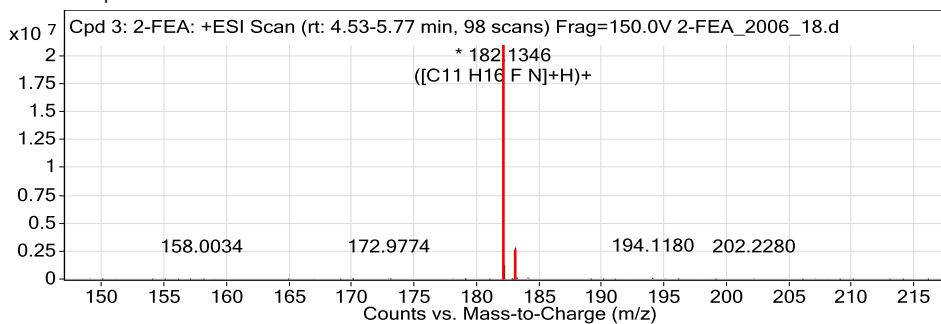
Compound Chromatograms



MFE MS Zoomed Spectrum



MS Zoomed Spectrum



MS Spectrum Peak List

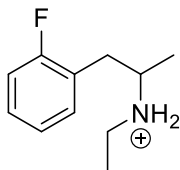
Obs. m/z	Charge	Abund	Formula	Ion/Isotope
182.1345	1	20939726	C11 H16 F N	(M+H)+
183.138	1	2465452.72	C11 H16 F N	(M+H)+

--- End Of Report ---

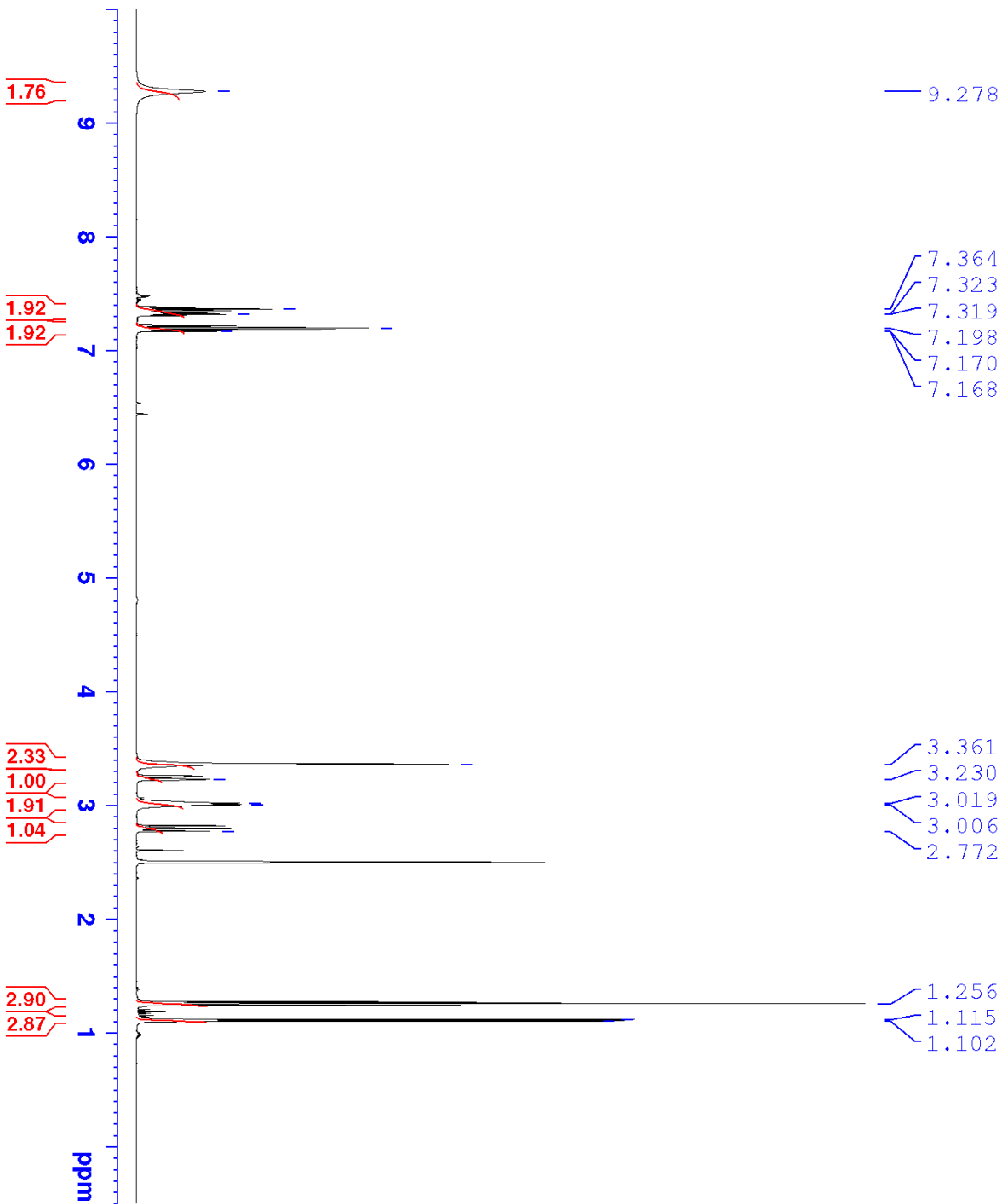
University
of Ljubljana
Faculty of Chemistry
and Chemical Technology



R E P O R T

Contract No.	C1714-17-460078 (Republic of Slovenia, Ministry of the Interior, POLICE)
Sample ID:	2006-18
Received date:	November 29, 2018
Our notebook code:	NFL-2006-18
NMR sample preparation:	20.0 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, ¹⁹ F
Proposed structure with formula, exact mass, molecular weight:	 Chemical Formula: C₁₁H₁₇FN⁺ Exact Mass: 182,1340 Molecular Weight: 182,2619
Chemical name:	<i>N</i> -protonated <i>N</i> -ethyl-1-(2-fluorophenyl)propan-2-amine
Comments:	- Structure elucidation based on 1D and 2D NMR spectra and HRMS. - >95% purity of a sample based on ¹H NMR spectrum
Supporting information:	Copies of ¹ H and ¹³ C NMR spectra, ¹ H and ¹³ C FIDs.
Principal investigator:	Prof. Dr. Janez Košmrlj
Date of report:	December 12, 2018

NFL-2006-18
¹H



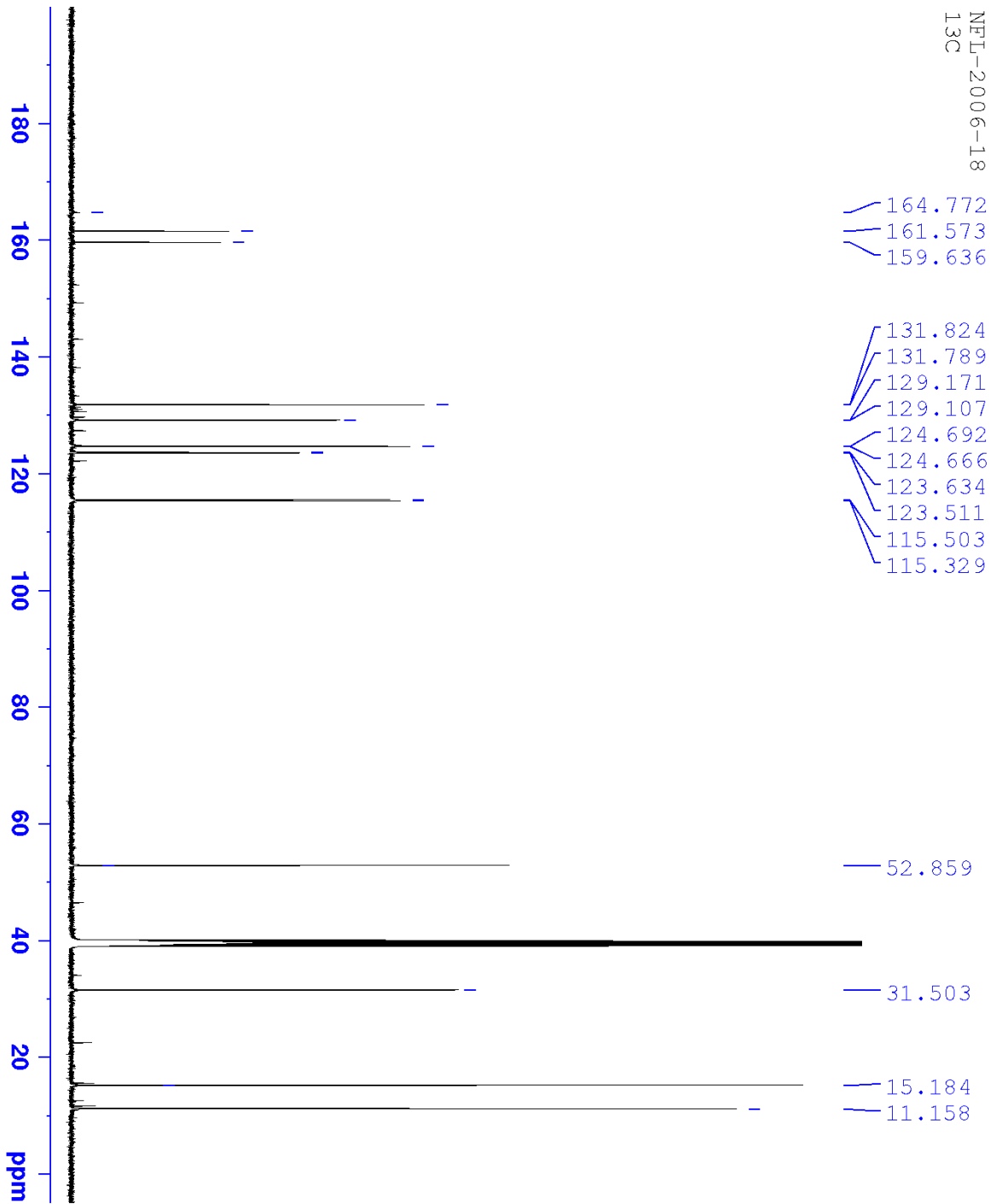
Current Data Parameters
 NAME NFL-2006-18
 EXPNO 1
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20181129
 Time 16.14
 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 3.2767999 sec
 RG 71.8
 DW 50.000 usec
 DE 6.50 usec
 TE 296.0 K
 D1 1.00000000 sec
 TDO 1

===== CHANNEL f1 =====
 SFO1 500.1330885 MHz
 NUC1 ¹H
 P1 8.70 usec
 PLW1 26.00000000 W

F2 - Processing parameters
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 SF 500.1300043 MHz
 WDW EM
 SSB 0
 LB 0.30 Hz
 GB 0
 PC 1.00

NFL-2006-18
13C



Current Data Parameters
NAME NFL-2006-18
EXPNO 3
PROCNO 1

F2 - Acquisition Parameters
Date_ 20181129
Time 20.02
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PROBHD 5 mm PABBO BB-
PULPROG zgpg30
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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 296.0 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

CHANNEL F1
SF01 125.7703637 MHz
NUC1 13C
P1 8.70 usec
PL1 122.0000000 W

CHANNEL F2
SF02 500.1320005 MHz
NUC2 1H
CPDPRG12 waltz16
PCPDZ 80.00 usec
PLW2 26.0000000 W
PLM12 0.36046001 W
PLM13 0.15113001 W

F2 - Processing parameters
SI 32768
SF 125.7578462 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40