

SLOVENIJA



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#### ANALYTICAL REPORT

## Furanyl-norfentanyl (C16H18N2O2)

## N-phenyl-N-(piperidin-4-yl)furan-2-carboxamide

Remark – other active cpd. detected: none

Sample ID:	1781-17		
Sample description:	liquid - clear		
Sample type:	RM-reference material		
Comments <sup>1</sup> :	CAY Lot#0491081; RESPONSE -purchasing		
Date of entry:	3/24/2017		

Substance identified- structure <sup>2</sup> (base form)	O NH		
Systematic name:	N-phenyl-N-(piperidin-4-yl)furan-2-carboxamide		
Other names:	N-phenyl-N-4-piperidinyl-2-furancarboxamide		
Formula (per base form)	C16H18N2O2		
M <sub>w</sub> (g/mol)	270,33		
Salt form:	HCI		
StdInChIKey (for base form)	DDSUGTXSJFAMIM-UHFFFAOYSA-N		
Other active cpd. detected	none		
Add.info (purity)	1 mg/ml in MeOH		

Stran 1 od 4

<sup>&</sup>lt;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:  $\underline{\text{http://opsin.ch.cam.ac.uk/}}\,$  DOI: 10.1021/ci100384d

## Report updates

date	comments (explanation)

## Supporting information

Analytical technique:	applied	remarks
GC-MS (El ionization)	+	NFL GC-RT (min): 8,57 BP(1): 95; BP(2): 175,BP(3):82,
FTIR-ATR	+	direct measurement after solvent evaporation
GC-IR (condensed phase)	+	always as base form

- **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  $\mu$ m. 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 0C 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. FTIR-ATR (Perkin Elmer): scan range 4000-400 cm<sup>-1</sup>; resolution 4cm<sup>-1</sup>
- 3. 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  $^{\circ}$ C. Chromatographic separation as above (1). Split MS: IR = 1:9.

MSD source EI = 70 eV. GC-MS transfer line T=  $235^{\circ}$ C, source and quadropole temperatures  $280^{\circ}$ C and  $180^{\circ}$ C, respectively. Scan range m/z scan range: from 50 (30 until 6 min.) to 550 (300) amu.

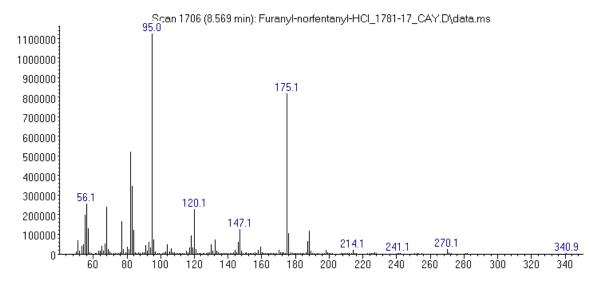
IR (condesed (solid) phase): IR scan range 4000 to 650, resolution 4 cm<sup>-1</sup>.

4. HPLC-TOF for exact monoisotopic mass and empirical formula control - results are not shown in the report.

#### **ANALYTICAL RESULTS**

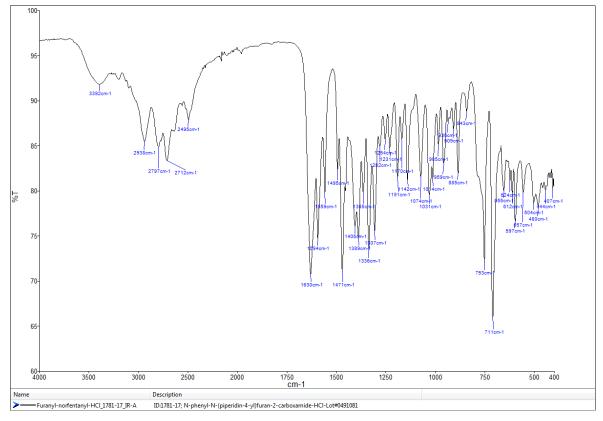
# MS (EI)

#### Abundance



m/z->

# FTIR-ATR - solid sample (MeOH solvent was evaporated)



# IR (condensed phase – after chromatographic separation)

