

FORENSICS MARKET

A FORENSIC TOXICOLOGY METHOD FOR THE DETERMINATION OF DESOMORPHINE, HEROIN, METHADONE, BUPRENORPHINE AND METABOLITES IN URINE USING LC/MS QQQ

Desomorphine, also known by its street name Krokodil, is an emerging synthetic opioid, used as a heroin substitute and produced by back street illicit laboratories.

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ABSTRACT

The analytical measurement of opioid panels commonly include a number of compounds such as heroin, methadone and buprenorphine. Research methods with sufficient flexibility are of value to respond to the need to measure new compounds resulting from clandestine production.

INTRODUCTION

Desomorphine, also known by its street name Krokodil, is an emerging synthetic opioid, used as a heroin substitute and produced by back street illicit laboratories. Little or no literature exists presently describing an analytical approach to determining desomorphine in samples.

This application brief describes the development of an LC/MS QQQ research method that aims to rapidly screen for to and measure desomorphine together with heroin, buprenorphine, methadone and their metabolites 6-acetylmorphine, norbuprenorphine and EDDP.

A three minute analytical research method has been developed with results obtained for five batches of spiked urine samples over the linear concentration range of 1-5000ng/ml. Precision data obtained, LLOQ, calibration linearity and accuracies will be reported for each analyte.

SAMPLE PREPARATION

Spiked Urine Sample Preparation (Dilute & Shoot):

1. Internal Standard Diluent Preparation (per individual batch):

a) Take 36ml of deionized water;

b) Add 10uL of each internal standard (100ug/ ml) or 1uL (1mg/ml) to the water;

c) Vortex the ~ 25ng/ml ISTD solution.

2. Negative Urine Pre-treatment:

a) Split 5ml of neat negative urine between 10x nanosep 3K centrifuge filters;

- b) Centrifuge @ 12000 for 20 min;
- c) Pool filtered urine into single tube or vessel;

d) Draw 4ml of the filtered urine and add to the 36ml ISTD solution (1);

e) This solution was used as calibrator diluent containing 25ng/ml of each ISTD.

6-Monoacetylmorphine-D6	Heroin-D9
Buprenorphine-D4	Methadone-D9
Desomorphine-D3	Norbuprenorphine-D3
EDDP-D3	

Table 1 – Internal Standards Utilized

HPLC Method Parameters (Agilent 1290 system):			
Column:	Agilent Poroshell 120 EC C18, 2.1 x 50mm (2.7µm)		
Column temperature:	55°C		
Injection volume:	1 uL		
Autosampler temp:	4°C		
Needle wash:	flushport (100% metha- nol), 5 sec		
Mobile phase:	A = 5mM NH4for- mate/0.01% formic acid in water		
	B = 0.01% formic acid in methanol		
Gradient flow rate:	0.5 mL/min		

Table 2 - HPLC Method Parameters

Time (min)	% A	%B
0.00	90	10
1.10	75	25
1.60	5	95
2.00	5	95
2.01	90	10
3.00	90	10

Table 3 – HPLC Gradient Profile

Mass Spectrometer Conditions & Configuration:			
Configuration:	Agilent 6460 QqQ MS		
Ion Source Conditions			
Ion Mode:	ESI/ Positive		
Capillary Voltage:	2500 V		
Nozzle Voltage:	0 V		
Drying gas (nitrogen):	9 L/min		
Drying gas temperature:	250 °C		
Nebulizer gas (nitrogen):	35 psi		
Sheath Gas temperature:	380 °C		
Sheath Gas flow:	11 L/min		

Table 4 - Mass Spectrometer Source Conditions

Cpd Name	Prec Ion	Prod Ion	Dwell	Frag (V)	CE (V)	Cell Acc (V)
6-monoacetyl morphine	328.2	165.1	20	158	41	3
6-monoacetyl morphine	328.2	43.1	20	158	60	3
6-monoacetyl morphine-D6	334.2	211	20	168	25	3
Buprenorphine	468.3	83.7	35	210	53	3
Buprenorphine	468.3	55.1	35	210	65	3
Buprenorphine-D4	472.3	88.1	35	220	49	3
Buprenorphine-D4	472.3	59.1	35	220	61	3
Desomorphine	272.2	165.2	20	161	65	3
Desomorphine	272.2	152.2	20	161	65	3
Desomorphine-D3	275.2	165.2	20	156	65	3
Desomorphine-D3	275.2	152.2	20	156	65	3
EDDP	278.2	234.1	20	158	33	7
EDDP	278.2	219.1	20	158	45	7
EDDP-D3	281.2	234.1	20	168	29	7
Heroin	370.2	268.1	20	149	37	7
Heroin	370.2	165	20	149	61	7
Heroin-D9	379.2	61.2	20	168	29	7
Methadone	310.2	265.1	20	112	9	7
Methadone	310.2	105	20	112	29	7
Methadone-D9	319.3	268.1	20	89	13	7
Norbuprenorphine	414.3	83.1	35	178	57	3
Norbuprenorphine	414.3	55.2	35	178	65	3
Norbuprenorphine-D3	417.3	101.1	35	198	45	3
Norbuprenorphine-D3	417.3	83.1	35	198	57	3

Table 5 - MRM acquisition Parameters

DESOMORPHINE RESULTS

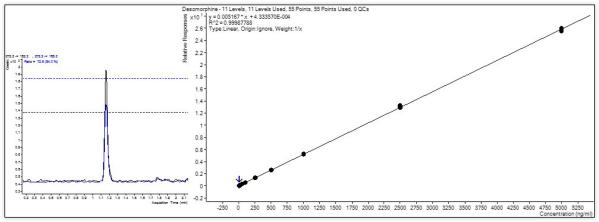


Figure 1 - Chromatography (1ng/ml) and Linear Dynamic Range (1-5000ng/ml.)

Concentration (ng/ml)	% RSD	Concentration (ng/ml)	%RSD
1	6.75	50	0.68
5	2.14	100	0.91
10	1.64	250	0.60
25	1.46	500	0.46

Table 6 – Precision Data for Desomorphine calibrator levels over N=5 Batches.

RESULTS OF OTHER OPIATE ANALYTES

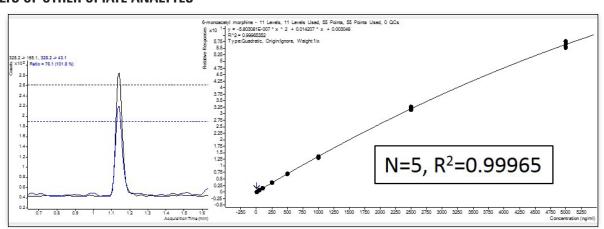


Figure 2 - 6-monoacetylmorphine (1ng/ml) and Dynamic Range (1-5000ng/ml.)

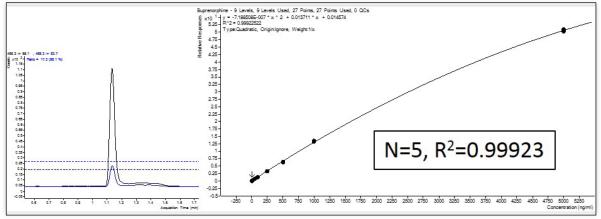


Figure 3 - Buprenorphine (1ng/ml) and Dynamic Range (1-5000ng/ml.)

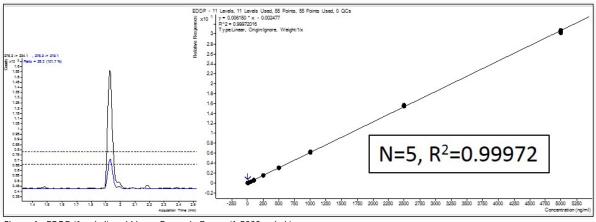


Figure 4 - EDDP (1ng/ml) and Linear Dynamic Range (1-5000ng/ml.)

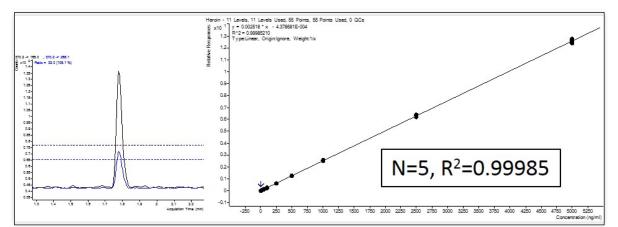


Figure 5 - Heroin (1ng/ml) and Linear Dynamic Range (1-5000ng/ml.)

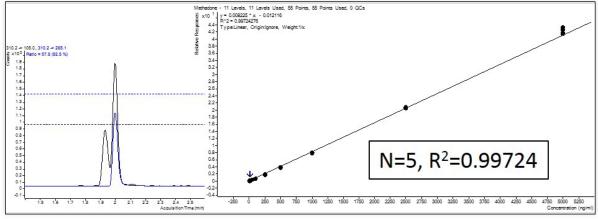


Figure 6 - Methadone (1ng/ml) and Linear Dynamic Range (1-5000ng/ml.)

SUMMARY OF OPIATE ANALYTICAL METHOD RESULTS:

i. LOQ values for all opiates in this study were typically <1ng/ml (except Buprenorphine which was 5ng/ml.)

ii. Over five separately spiked batches of samples, %RSD values for desomorphine were <6.75% across the Linear range.

iii. This rapid three minute method illustrates the feasibility of distinguishing between the illicit street opiates inclusive of desomorphine.

CONCLUSIONS

The analytical method developed herein has demonstrated the feasibility of:

• Analyzing illicit opiate/opioids & metabolites directly and individually in a rapid three minute chromatographic method;

• Demonstrated the ability to detect desomorphine from a LOQ < 1ng/ml across a wide linear dynamic through at least 5000ng/ml;

• Illustrated that low limits of quantitation can be achieved for opiates and their metabolites with minimal sample preparation;

• Wide linear dynamic ranges of up to 4 orders can be achieved for the analytes in this study.



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