

Automated Sample Preparation for a Comprehensive Drug Research Panel from Oral Fluid Using Intercept i2® Device

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Overview

- Automation of clean extraction, high recovery oral fluid sample preparation method
- Sample prep with no human interaction or supervision
- · High precision and accuracy for low and high QCs

Introduction

There is a growing interest in oral fluid testing over other test matrices such as urine and blood because it is non-intrusive, convenient, and observable - making adulteration or substitution difficult. In a recent technical note (www.phenomenex.com/ TN100) we developed a unique sample preparation procedure that encompasses a wide range of illicit and pain management drugs that result in a very clean extract with high recovery. 1,2 Here, we extend our scope in automating the procedure by utilizing a liquid handler. The speed of data analysis has improved significantly in the past decade due to increased detection capabilities of mass spectrometry coupled with liquid chromatography. Despite these advances, the biggest time constraint in sample processing is the manual process around sample preparation. One survey attributes 61 % of an analytical chemist's time is spent on sample processing3. Automated liquid handling can increase lab productivity and circumvents human error.

Materials and Methods

Reagents and Chemicals

Analytical reference standards and human saliva were purchased from Cerilliant® Corporation (Round Rock, TX) and BioreclamationIVT® (Chestertown, MD). The Intercept i2® oral fluid collection devices was obtained from OraSure Technologies, Inc. (Bethlehem, PA). All other chemicals were obtained from the Sigma-Aldrich Company (St. Louis, MO). D.I. water via Sartorius® arium® Comfort II, courtesy of Sartorius Corporation (Bohemia, NY). Liquid Handling via Tecan Freedom EVO® 100. (San Jose, CA)

Methods

Sample collection

1.0 mL of saliva was pipetted onto the application tip of the oral fluid collection device. The saturated pad was then placed into the transport tube containing the buffer solution.

Sample pretreatment

The Intercept i2 applicator tip that absorbed about 1 mL of oral fluid was transferred to the transport tube containing the preservative buffer. After breaking the plastic nipple at the end of the transport tube, it was placed in a centrifuge tube and centrifuge at 600 g for 15 min to collect the supernatant. The transport tube was placed directly on the automation platform. Liquid handler pipetted 0.5 mL from the top of the sample, to avoid transfer of debris onto the SPE cartridge.



Sean Orlowicz Manager, PhenoLogix When not in the lab, Sean enjoys just about anything involving the outdoors: hiking, climbing, surfing, etc. He is especially at home in the mountains, being an avid skier and motorcyclist.







Sample Prep Automation

The method, found on page 2, can be amenable to many different automation platforms. We selected single carrier configuration with inserts that process up to eight samples in parallel with industry standard 3mL cartridges. This selection accommodates loading of 40 SPE cartridges simultaneously. Because all wash solvents used in the extraction steps for both acidic and basic compounds were the same, automation allowed for simultaneous extraction. For the elution solvent, the liquid handler loaded two elution solvent separately for the cationic and anionic sorbent media respectively. The duration and amount of positive pressure application to the extracted cartridges are all software controlled. Calibrators for the 7-point linearity curve were prepared by serial dilution of the spiked oral fluid. The curve spanned a total of seven concentration levels. The QC samples for extraction were prepared at two concentration levels. Addition of acidic (methanolic) solution to the eluted samples were necessary to prevent loss of free bases in the dry down step.





SPE Method

Step	Basic analyte extraction	Acidic analyte extraction
Product Name:	Strata®-X-C, 30 mg in 3 mL cartridge	Strata-X-A, 30 mg in 3 mL cartridge
Part No.:	8B-S029-TBJ	8B-S123-TBJ
Condition:	1 mL 100 % Methanol	1 mL 100 % Methanol
Equilibrate:	1 mL DI Water	1 mL DI Water
Load:	Combine 0.5 mL of pretreated sample with 1 mL 1 % formic acid, mix/vortex 5-10 sec and load on Strata-X-C.	Combine 0.5 mL of pretreated sample with 1 mL 1 % ammonium hydroxide, mix/vortex 5-10 sec and load on Strata-X-A.
Wash 1:	1 mL DI Water	1 mL DI Water
Wash 2:	1 mL Acetone/Water (50:50)	1 mL Acetone/Water (50:50)
Dry Cartridge:	2-3 minutes under positive pressure	2-3 minutes under positive pressure
Elute:	2 x 500 µL Methanol/Acetonitrile/ 28-30 % Ammonium Hydroxide (5:5:2)	2 x 500 µL Methanol/Acetonitrile/ Conc Formic acid (50:50:5)
Optional Addition*:	30 µL of 50 mM HCI/Methanol	-
Dry down:	Evaporate to dryness under gentle stream N ₂ at 45-50 °C	Evaporate to dryness under gentle stream of N ₂ at 45-50 °C
Reconstitute:	With 125 µL initial mobile phase	With 125 µL initial mobile phase

Combine both fractions into a single autosampler vial

*to help prevent the loss of free bases during evaporation

LC/MS Conditions

The LC/MS/MS method utilized a Kinetex® Biphenyl 2.6 µm, 50 x 3.0 mm column (Part No.:00B-4622-Y0) with a simple mobile phase consisting of 0.1% formic acid in water and methanol. A fast LC gradient resulted in total run time of 5 min. The detection was carried out on a SCIEX API 5000™ equipped with ESI source. For basic compounds the MS was operated under positive polarity and in a separate injection, all acidic compounds (except lorazepam) were analyzed in negative polarity.

Positive ESI Panel

Column: Kinetex® 2.6 µm Biphenyl

Dimensions: 50 x 3.0 mm Part No.: 00B-4622-Y0 SecurityGuard™ Ultra Cartridge: AJ0-9208 SecurityGuard Ultra Holder: AJO-9000

Mobile Phase: A: 0.1 % Formic acid in Water

B: 0.1 % Formic acid in Methanol

Gradient: Time (min) % B 10 Λ 4 95 95 5 5.01 10 7.5 10

Flow Rate: 500 µL/min Temperature: Ambient Injection Volume: 10 µL Detection: ESI+ Detection System: SCIEX API 5000

Negative ESI Panel

Column: Kinetex 2.6 µm Biphenyl

Dimensions: 50 x 3.0 mm Part No.: 00B-4622-Y0 Guard Cartridge: AJ0-9208 Guard Holder: AJ0-9000

Mobile Phase: A: 0.1 mM Ammonium formate in Water

B: 100 % Methanol

Gradient: Time (min) % B 0 10 95 5 95 5.01 10 10

Flow Rate: 500 µL/min Temperature: Ambient Injection Volume: 10 µL Detection: ESI-Detection System: SCIEX API 5000



Table 1. Precision, accuracy and linear regression data for analytes

Analyte	Linear regression (R2)	QC1 (ng/mL)	%Accuracy (QC1)	%CV (QC1)	QC2 (ng/mL)	%Accuracy (QC2)	%CV (QC2)
6-MAM	0.9997	2.5	97.2	4.6	8	116.7	11.5
7-Aminoclonazepam	0.9999	2.5	103	2.2	80	116.5	5.1
$\alpha\text{-hydroxyalprazolam}$	0.9966	25	96.9	15.6	80	112	5.5
Alprazolam	0.9998	25	87.8	7.5	80	107.2	5.6
Amitriptyline	0.9991	12.5	101.3	10.8	40	113.1	5.5
Amphetamine	0.9994	25	84.3	14.1	80	116.7	4.2
Benzoylecgonine	0.9992	25	87.6	8.6	80	107.1	5.8
Buprenorphine	0.9994	5	113.8	4.9	16	105.2	8.9
Citalopram	0.997	12.5	100.6	7.9	40	110	10.7
Codeine	0.9995	25	90.8	8.9	80	98.1	3
Diazepam	0.9987	25	88.9	3.5	80	112.7	14.8
Fentanyl	0.9977	2.5	102.5	5.4	8	118.3	11.4
Fluoxetine	0.9956	25	98	11.3	80	104.6	15.2
Gabapentin	0.9967	25	94.9	14.3	80	108.1	7.7
Hydrocodone	0.9961	25	99	2.6	80	108.1	2.9
Hydromorphone	0.9955	25	98.7	7.9	80	109.9	10.3
Imipramine	0.9979	12.5	101.1	13.7	40	115.8	9.4
Lorazepam	0.9963	25	93	9.3	80	112.6	8.14
MDMA	0.9971	25	90.3	10	80	97.9	11.8
Meperidine	0.9982	25	90.4	6.8	80	113	6.9
Methadone	0.9998	25	102.2	9.6	80	82.3	10.8
Methamphetamine	0.9982	25	101.2	9.5	80	106.1	15.7
Methylphenidate	0.9994	2.5	95.3	5.8	8	113.6	4.8
Morphine	0.9988	25	101.7	10.9	80	107.4	7.7
Norbuprenorphine	0.9999	5	103.4	11.8	16	115	6.8
Nordiazepam	0.998	25	105.7	2.9	80	115.2	15.3
Norfentanyl	0.9993	2.5	92.2	9.1	8	114.8	4.2
Norhydrocodone	0.9982	25	91.4	13.1	80	96	3.7
Noroxycodone	0.9981	25	95	4.9	80	107.9	11.8
Normorphine	0.9956	25	95.1	7.9	80	99.5	4.2
Nortriptyline	0.9962	12.5	100.1	12.2	40	118.1	4.9
0-Desmethyltramadol	0.9975	25	114.1	2.2	80	94.5	8.6
Oxycodone	0.9932	25	104.3	2.1	80	115.9	11.3
Oxymorphone	0.9994	12.5	86.4	8.6	40	103.5	11.9
Paroxetine	0.9969	12.5	85.2	14.3	40	91.9	15.1
PCP	0.9997	12.5	96.1	12.3	40	115.3	12.5
Pregabalin	0.9983	25	102.7	9.9	80	112.7	13.4
Sertraline	0.9965	12.5	80.9	18.7	40	100.9	13.1
Zolpidem-carboxy	0.999	5	101.1	11.1	16	120	7.2
Tapentadol	0.997	25	102.3	5	80	107	15.2
Zolpidem	0.9999	5	101	12.2	16	118.1	4.9
Butalbital	0.9994	25	100.5	5.3	80	113.5	7.8
Phenobarbital	0.9979	25	82.8	5	80	89.1	5.9
Secobarbital	0.9997	25	93.2	5	80	96.8	8.3
THC-COOH	0.999	12.5	99	4.9	40	106.8	6.2





Figure 1.Calibration curve of extracted samples representing dynamic range of morphine (1-300 ng/mL); R=0.9988.

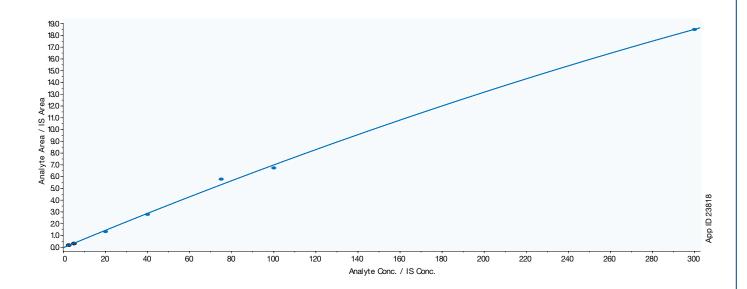


Figure 2.Calibration curve of extracted samples representing dynamic range of 6 MAM (0.1-30 ng/mL); R=0.9997.

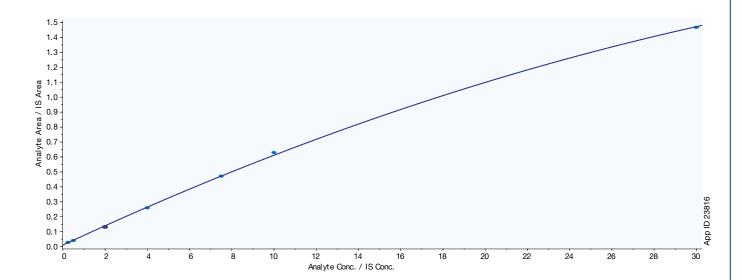




Figure 3.Calibration curve of extracted samples representing dynamic range of zolpidem (0.5-60 ng/mL); R=0.9999.

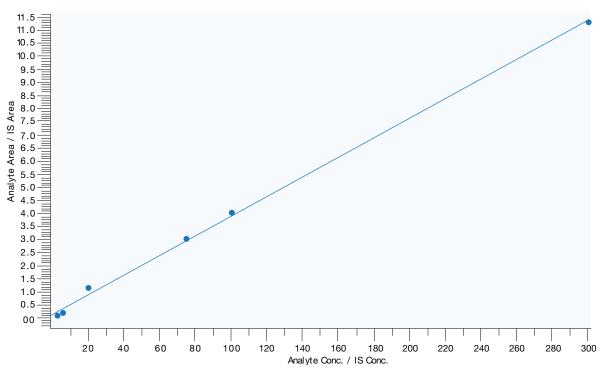


Figure 4. Calibration curve of extracted samples representing dynamic range of norfentanyl (0.25-30 ng/mL); R=0.9993.

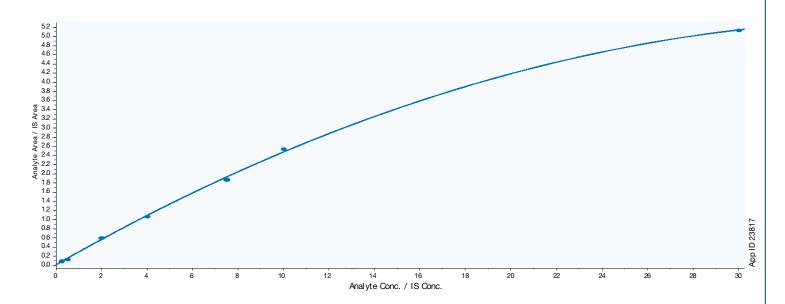




Figure 5.Calibration curve of extracted samples representing dynamic range of butalbital (1-300 ng/mL); R=0.9994.

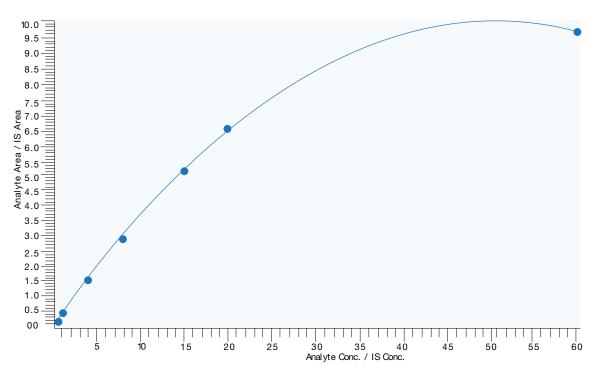
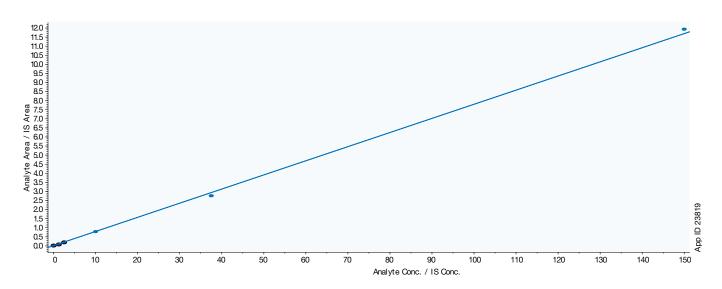


Figure 6.Calibration curve of extracted samples representing dynamic range of THC-COOH (0.5 ng/mL-150 ng/mL); R=0.9990.





Results and Discussion

Calibration curves for extracted samples in this study covered a range from 0.1 ng/mL to 300 ng/mL (Figures 1-6). At least five points per analyte were used for calibration curve. Beyond 300 ng/mL we encountered saturation of the MS for most of the analytes. Several analytes, zolpidem, norfentanyl, butalbital and THC-COOH (Figures 3-6), displayed non-linear curve. A quadratic calibration curve with 1/x weighting was applied to all analytes in this assay. Table 1 shows, the correlation coefficient value (R) in all cases were greater than 0.995. The precision and accuracy for low and high QCs spanned from 3-15% and 85-112% respectively for all four replicates at each concentration level.

Ordering Information

Kinetex®

Kinetex Core-Shell HPLC/UHPLC 2.6 µm Minibore Columns		SecurityGuard™ ULTRA Cartridges*	
Phase	50 x 3.0 mm	3/pk	
Biphenyl	00B-4622-Y0	AJ0-9208	

^{*}SecurityGuard ULTRA Cartridges require holder, Part No.: AJ0-9000

Strata®-X-C

Format	Sorbent Mass	Part Number	Unit	
Tube				
	30 mg	8B-S029-TAK**	1 mL (100/box)	
	30 mg	8B-S029-TBJ	3 mL (50/box)	
	60 mg	8B-S029-UBJ**	3 mL (50/box)	
	100 mg	8B-S029-EBJ	3 mL (50/box)	
	100 mg	8B-S029-ECH	6 mL (30/box)	
	200 mg	8B-S029-FBJ	3 mL (50/box)	
	200 mg	8B-S029-FCH	6 mL (30/box)	
	500 mg	8B-S029-HBJ	3 mL (50/box)	
	500 mg	8B-S029-HCH	6 mL (30/box)	
Giga™ Tube				
THE PERSON NAMED IN COLUMN TWO IS NOT THE PERSON NAMED IN COLUMN TWO IS NAM	500 mg	8B-S029-HDG	12 mL (20/box)	
Name and Address of the Owner, when the Owner, which the Owne	1 g	8B-S029-JDG	12 mL (20/box)	
	1 g	8B-S029-JEG	20 mL (20/box)	
	2 g	8B-S029-KEG	20 mL (20/box)	
	5 g	8B-S029-LFF	60 mL (16/box)	
96-Well Plate				
	10 mg	8E-S029-AGB	2 Plates/Box	
The same of	30 mg	8E-S029-TGB	2 Plates/Box	
12 2	60 mg	8E-S029-UGB	2 Plates/Box	
96-Well Microelution Plate				
100	2 mg	8M-S029-4GA	ea	

^{**}Tab-less tubes available. Contact Phenomenex for details.

Conclusion

Oral fluid is a complex matrix and LC/MS/MS analysis requires good recovery and a clean extract. Automated sample prep on a liquid handling robot helps rapidly increase throughput and reduce human error. This procedure provides good dynamic calibration range with good precision and accuracy with little human intervention.

References:

- S. Huq, S. Sadjadi, L. Snow; "A Superior Sample Preparation of Comprehensive Drug Panel Analytes from Oral Fluid Collection Devices; Phenomenex TN-0100
- S. Sadjadi, S. Huq, L. Snow; "An Investigation into Removing the Excipients from Select Oral Fluids Collection Devices by SPE and LC/MS Detection;" Mass Spec Application for Clinical Laboratory Conference, 2016
- 3. LC|GC Editors. "Overview of Sample Preparation" LC|GC November 01, 2015 Volume 33, Issue 11 (pg 46-51). Accessed on Sept 12, 2016 from http://www.chromatographyonline.com/overview-sample-preparation.

Strata-X-A

Format	Sorbent Mass	Part Number	Unit
Tube			
	30 mg	8B-S123-TAK**	1 mL (100/box)
	30 mg	8B-S123-TBJ	3 mL (50/box)
	60 mg	8B-S123-UBJ	3 mL (50/box)
	100 mg	8B-S123-EBJ	3 mL (50/box)
	100 mg	8B-S123-ECH	6 mL (30/box)
	200 mg	8B-S123-FBJ	3 mL (50/box)
	200 mg	8B-S123-FCH	6 mL (30/box)
	500 mg	8B-S123-HBJ	3 mL (50/box)
	500 mg	8B-S123-HCH	6 mL (30/box)
Giga™ Tube			
PRODUCT -	500 mg	8B-S123-HDG	12 mL (20/box)
Personal	1 g	8B-S123-JDG	12 mL (20/box)
	1 g	8B-S123-JEG	20 mL (20/box)
	2 g	8B-S123-KEG	20 mL (20/box)
	5 g	8B-S123-LFF	60 mL (16/box)
96-Well Plate			
	10 mg	8E-S123-AGB	2 Plates/Box
One I	30 mg	8E-S123-TGB	2 Plates/Box
1 4 4	60 mg	8E-S123-UGB	2 Plates/Box
96-Well Microelution	ı Plate		
1500	2 mg	8M-S123-4GA	ea



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