

Test Updates

Immediate Action

Modified

Updated: Test Names for 1864B, 1864FL, 1864SP, 1864TI, 1864U

In our continuing effort to provide you with the highest quality toxicology laboratory services available, we have compiled important changes regarding a number of tests we perform. Listed below are the types of changes that may be included in this notification, effective Monday, August 15, 2022

Test Changes - Tests that have had changes to the method/ CPT code, units of measurement, scope of analysis, reference comments, or specimen requirements.

Discontinued Tests - Tests being discontinued with alternate testing suggestions.

Please use this information to update your computer systems/records. These changes are important to ensure standardization of our mutual laboratory databases.

If you have any questions about the information contained in this notification, please call our Client Support Department at (866) 522-2206. Thank you for your continued support of NMS Labs and your assistance in implementing these changes.

The CPT Codes provided in this document are based on AMA guidelines and are for informational purposes only. NMS Labs does not assume responsibility for billing errors due to reliance on the CPT Codes listed in this document.

Monday, August 15, 2022



Test Updates

Test Name	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
Amphetamines Confirmation (DUID/DRE), Blood				•	•			
Amphetamines Confirmation (Qualitative)					•			
Barbiturates Confirmation (DUID/DRE), Blood			•	•	•			
Barbiturates Confirmation (Qualitative) (DUID/DRE), Urine					•			
Bath Salts Confirmation, Blood								•
Bath Salts Confirmation, Serum/Plasma								•
Bath Salts Confirmation, Urine								•
Benzodiazepines Confirmation (DUID/DRE), Blood					•			
Benzodiazepines Confirmation (Qualitative) (DUID/DRE), Urine					•			
Cannabinoids Confirmation (Qualitative), Tissue	•							
DUID/DRE Designer Benzodiazepines					•			
DUID/DRE Designer Opioids					•			
DUID/DRE Expanded Drug Screen Add-			•		•	•		
DUID/DRE Expanded Drug Screen Add-		•			•	•		
DUID/DRE Panel (w/Alcohol)					•	•		
DUID/DRE Panel (w/Alcohol), Urine					•			
DUID/DRE Panel ProofPOSITIVE®,					•	•		
					•			
DUID/DRE Screen, Blood (Forensic)						•		
DUID/DRE Substituted Cathinone					•			
DUID/DRE Substituted Cathinone					•			
Designer Benzodiazepines Confirmation	•				•			
Designer Benzodiazepines Confirmation	<u> </u>				•			
Designer Benzodiazepines Confirmation	•				•			
Designer Benzodiazepines Confirmation	•				•			
Designer Benzodiazepines Confirmation	•				•			
Designer Benzodiazepines Confirmation					<u> </u>			•
	Amphetamines Confirmation (DUID/DRE), Blood Amphetamines Confirmation (Qualitative) (DUID/DRE), Urine Barbiturates Confirmation (Qualitative) (DUID/DRE), Urine Barbiturates Confirmation (Qualitative) (DUID/DRE), Urine Bath Salts Confirmation, Blood Bath Salts Confirmation, Serum/Plasma Bath Salts Confirmation, Urine Benzodiazepines Confirmation (DUID/DRE), Blood Benzodiazepines Confirmation (Qualitative) (DUID/DRE), Urine Cannabinoids Confirmation (Qualitative), Tissue DUID/DRE Designer Benzodiazepines Confirmation, Blood DUID/DRE Designer Opioids Confirmation, Blood DUID/DRE Expanded Drug Screen Add- On, Urine (Forensic) DUID/DRE Panel (w/Alcohol) ProofPOSITIVE®, Blood (Forensic) DUID/DRE Panel (w/Alcohol), Urine (Forensic) DUID/DRE Panel, Urine (Forensic) DUID/DRE Panel, Urine (Forensic) DUID/DRE Panel, Urine (Forensic) DUID/DRE Substituted Cathinone Confirmation, Blood DUID/DRE Substituted Cathinone Confirmatio	Name Amphetamines Confirmation (DUID/DRE), Blood (DUID/DRE), Urine Barbiturates Confirmation (DUID/DRE), Blood (DUID/DRE), Urine Barbiturates Confirmation (Qualitative) (DUID/DRE), Urine (DUID/DRE), Bath Salts Confirmation, Blood Bath Salts Confirmation, Serum/Plasma (Duid) Bath Salts Confirmation, Urine Benzodiazepines Confirmation (DUID/DRE), Blood (Duid) Benzodiazepines Confirmation (DUID/DRE), Blood Benzodiazepines Confirmation (Qualitative) (DUID/DRE), Urine • Cannabinoids Confirmation (Qualitative), Tissue • DUID/DRE Designer Benzodiazepines Confirmation, Blood • DUID/DRE Designer Opioids Confirmation, Blood • DUID/DRE Expanded Drug Screen Add- On ProofPOSITIVE®, Blood (Forensic) • DUID/DRE Expanded Drug Screen Add- On, Urine (Forensic) • DUID/DRE Panel (w/Alcohol) ProofPOSITIVE®, Blood (Forensic) DUID/DRE Panel (w/Alcohol), Urine (Forensic) • DUID/DRE Panel, Vrine (Forensic) • DUID/DRE Panel, Vrine (Forensic) • DUID/DRE Screen, Blood (Forensic) • DUID/DRE Substituted Cathinone Confirmation, Urine • Designer Benzodiazepines Confirmation (Qualitative) (DUID/DRE), U	NameCPT CodeAmphetamines Confirmation (DUID/DRE), Blood	NameCPT CodeReq.Amphetamines Confirmation (DUID/DRE), BloodAmphetamines Confirmation (Qualitative) (DUID/DRE), UrineBarbiturates Confirmation (DUID/DRE), BloodBarbiturates Confirmation, Qualitative) (DUID/DRE), UrineBath Salts Confirmation, BloodBath Salts Confirmation, Serum/PlasmaBath Salts Confirmation, Serum/PlasmaBath Salts Confirmation, UrineBenzodiazepines Confirmation (DUID/DRE), UrineCannabinoids Confirmation (Qualitative), TissueDUID/DRE Designer Benzodiazepines Confirmation, BloodDUID/DRE Designer Opioids Confirmation, BloodDUID/DRE Designer Opioids Confirmation, BloodDUID/DRE Papended Drug Screen Add- On ProofPOSITIVE®, Blood (Forensic)DUID/DRE Panel (w/Alcohol) ProofPOSITIVE®, Blood (Forensic)DUID/DRE Panel (w/Alcohol), Urine (Forensic)DUID/DRE Panel ProofPOSITIVE®, Blood (Forensic)DUID/DRE Substituted Cathinone Confirmation, BloodDUID/DRE Substituted Cathinone Confirmation, BloodDUID/DRE Substituted Cathinone Confirmation, BloodDUID/DRE Substituted Cathinone Confirmation, UrineDuID/DRE Substituted Cathinone Confirmation, Urine <td< td=""><td>Name CPT Code Req. Amphetamines Confirmation (DUID/DRE), Blood • • Amphetamines Confirmation (Qualitative) (DUID/DRE), Urine • • Barbiturates Confirmation (DUID/DRE), Blood • • Barbiturates Confirmation, Blood • • Bath Salts Confirmation, Serum/Plasma • • Bath Salts Confirmation, Vrine • • Benzodiazepines Confirmation • • Genzodiazepines Confirmation • • Gualitative) (DUID/DRE), Urine • • Benzodiazepines Confirmation • • Gualitative) (DUID/DRE), Urine • • Cannabinoids Confirmation (Qualitative), Tissue • • DUID/DRE Designer Benzodiazepines • • On ProofPOSITIVE®, Blood • • DUID/DRE Expanded Drug Screen Add- On, Urine (Forensic) • • DUID/DRE Panel (w/Alcohol) • • • DUID/DRE Panel (w/Alcohol), Urine (Forensic) • • • DU</td><td>NameCPT CodeReq.Amphetamines Confirmation(DUID/DRE), BloodAmphetamines Confirmation (Qualitative)(DUID/DRE), UrineBarbiturates Confirmation (Qualitative).BloodBarbiturates Confirmation (Qualitative).(DUID/DRE), UrineBarbiturates Confirmation, Blood.Bath Salts Confirmation, Serum/Plasma.Bath Salts Confirmation, Urine.Benzodiazepines Confirmation.(DUID/DRE), Blood.Benzodiazepines Confirmation.(Qualitative).Tissue.DUD/DRE Designer Benzodiazepines.Confirmation, Blood.DUID/DRE Designer BenzodiazepinesConfirmation, Blood.DUID/DRE Designer Depiolds.Confirmation, Blood.DUID/DRE Expanded Drug Screen AddOn ProofPOSITIVE®, Blood (Forensic).DUID/DRE Expanded Drug Screen AddDUID/DRE Panel (wAlcohol).ProofPOSITIVE®, Blood (Forensic).DUID/DRE Panel (wAlcohol), Urine.(Forensic).DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Panel (wAlcohol), Urine.(Forensic).DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Substit</td><td>NameCPT CodeReq.Amphetamines Confirmation</td><td>NameCPT CodeReq.CommentsAmphetamines ConfirmationAmphetamines Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation, Blood<td< td=""></td<></td></td<>	Name CPT Code Req. Amphetamines Confirmation (DUID/DRE), Blood • • Amphetamines Confirmation (Qualitative) (DUID/DRE), Urine • • Barbiturates Confirmation (DUID/DRE), Blood • • Barbiturates Confirmation, Blood • • Bath Salts Confirmation, Serum/Plasma • • Bath Salts Confirmation, Vrine • • Benzodiazepines Confirmation • • Genzodiazepines Confirmation • • Gualitative) (DUID/DRE), Urine • • Benzodiazepines Confirmation • • Gualitative) (DUID/DRE), Urine • • Cannabinoids Confirmation (Qualitative), Tissue • • DUID/DRE Designer Benzodiazepines • • On ProofPOSITIVE®, Blood • • DUID/DRE Expanded Drug Screen Add- On, Urine (Forensic) • • DUID/DRE Panel (w/Alcohol) • • • DUID/DRE Panel (w/Alcohol), Urine (Forensic) • • • DU	NameCPT CodeReq.Amphetamines Confirmation(DUID/DRE), BloodAmphetamines Confirmation (Qualitative)(DUID/DRE), UrineBarbiturates Confirmation (Qualitative).BloodBarbiturates Confirmation (Qualitative).(DUID/DRE), UrineBarbiturates Confirmation, Blood.Bath Salts Confirmation, Serum/Plasma.Bath Salts Confirmation, Urine.Benzodiazepines Confirmation.(DUID/DRE), Blood.Benzodiazepines Confirmation.(Qualitative).Tissue.DUD/DRE Designer Benzodiazepines.Confirmation, Blood.DUID/DRE Designer BenzodiazepinesConfirmation, Blood.DUID/DRE Designer Depiolds.Confirmation, Blood.DUID/DRE Expanded Drug Screen AddOn ProofPOSITIVE®, Blood (Forensic).DUID/DRE Expanded Drug Screen AddDUID/DRE Panel (wAlcohol).ProofPOSITIVE®, Blood (Forensic).DUID/DRE Panel (wAlcohol), Urine.(Forensic).DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Panel (wAlcohol), Urine.(Forensic).DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Panel (wAlcohol), Urine.Confirmation, Blood.DUID/DRE Substit	NameCPT CodeReq.Amphetamines Confirmation	NameCPT CodeReq.CommentsAmphetamines ConfirmationAmphetamines Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation (Qualitative)Barbiturates Confirmation, Blood <td< td=""></td<>

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Test	Test Name	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
52503B	Designer Benzodiazepines Confirmation 2, Blood								•
52503SP	Designer Benzodiazepines Confirmation 2, Serum/Plasma								•
52493B	Designer Benzodiazepines Confirmation, Blood					•			
52493SP	Designer Benzodiazepines Confirmation, Serum/Plasma					•			
52488U	Designer Opioids Confirmation (Qualitative), Urine					•			
52488B	Designer Opioids Confirmation, Blood					•			
52500B	Designer Opioids Confirmation, Blood					•			
52488SP	Serum/Plasma					•			
52500SP	Designer Opioids Confirmation, Serum/Plasma					•			
1480B	Designer Opioids, Blood					٠			
1480SP	Designer Opioids, Serum/Plasma					•			
8030B	Drug Facilitated Crime Panel, Blood (Forensic)						•		
8030SP	Drug Facilitated Crime Panel, Serum/Plasma (Forensic)						•		
8030U	Drug Facilitated Crime Panel, Urine (Forensic)						•		
8098B	Drug Screen (GC/MS), Blood					•	•		
8098SP	Drug Screen (GC/MS), Serum/Plasma			•		•	•		
8098U	Drug Screen (GC/MS), Urine					•	•		
1876B	Drug Screen - Expanded, Blood						•		
1876FL	Drug Screen - Expanded, Fluid						•		
1876SP	Drug Screen - Expanded, Serum/Plasma						•		
1876U	Drug Screen - Expanded, Urine						•		
1864U	Drugs of Abuse Screen (10 Panel), Urine	•	•			•			
1864B	Drugs of Abuse Screen (11 Panel), Blood	•				•			
1864FL	Drugs of Abuse Screen (11 Panel), Fluid	•				•			
1864SP	Drugs of Abuse Screen (11 Panel), Serum/Plasma	•				•			
1864TI	Drugs of Abuse Screen (11 Panel), Tissue	•				•			
90023B	Expanded Drug Screen (DUID/DRE), Blood (Forensic) (CSA)					•	•		
52486U	Fentanyl Panel Confirmation, Urine	•				•			
52486B	Fentanyl and 4-ANPP Confirmation, Blood	•				٠			

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Test Updates

Test	Test Name	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
52486SP	Fentanyl and 4-ANPP Confirmation, Serum/Plasma	•				•			
1860B	GC/MS Drug Screen (Acid/Neutral), Blood						•		
10053U	GC/MS Drug Screen, Urine (CSA)						•		
52320B	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Blood								•
52320SP	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Serum/Plasma								•
52320U	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Urine								•
52081B	Metoclopramide Confirmation, Blood								•
52081FL	Metoclopramide Confirmation, Fluid								•
52081SP	Metoclopramide Confirmation, Serum/Plasma								•
52081TI	Metoclopramide Confirmation, Tissue								•
52081U	Metoclopramide Confirmation, Urine								•
54342U	Mitragynine, Phenazepam Confirmation (Qualitative) (DUID/DRE), Urine								•
8756B	Novel Psychoactive Substances (NPS) Screen 1, Blood					•			
8756SP	Novel Psychoactive Substances (NPS) Screen 1, Serum/Plasma					•			
8756U	Novel Psychoactive Substances (NPS) Screen 1, Urine				•	•			
90036U	Opioids Panel, Urine (CSA)					•			
90035U	Opioids Screen, Urine (CSA)					•			
52326B	Piperazine Designer Drugs Confirmation, Blood								•
52326SP	Piperazine Designer Drugs Confirmation, Serum/Plasma								•
52326U	Piperazine Designer Drugs Confirmation, Urine								•
8155U	Postmortem Designer Opioids Add-On (Qualitative), Urine (Forensic)					•			
8155B	Postmortem Designer Opioids Add-On, Blood (Forensic)					•			
8155SP	Postmortem Designer Opioids Add-On, Serum/Plasma (Forensic)					•			
8063B	Postmortem, Basic to Expanded Upgrade, Blood (Forensic)					•	•		
8063FL	Postmortem, Basic to Expanded Upgrade, Fluid (Forensic)					•	•		
8063SP	Postmortem, Basic to Expanded Upgrade, Serum/Plasma (Forensic)					•	•		
8063TI	Postmortem, Basic to Expanded Upgrade, Tissue (Forensic)					•	•		

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Test Updates

Test	Test Name	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
8063U	Postmortem, Basic to Expanded Upgrade, Urine (Forensic)					•	•		
8084B	Postmortem, Expanded w/ Vitreous Alcohol and 6-MAM Confirmation, Blood (Forensic)					٠	•		
8042B	Postmortem, Expanded w/Vitreous Alcohol Confirmation, Blood (Forensic)					•	•		
10052B	Postmortem, Expanded w/Vitreous Alcohol Confirmation, Blood (Forensic) (CSA)					•	•		
8057B	Postmortem, Expanded w/Vitreous Alcohol Confirmation, Blood - University of MI (Forensic) (CSA)					•	•		
8062B	Postmortem, Expanded w/o Alcohol, Blood (Forensic)					•	•		
8062FL	Postmortem, Expanded w/o Alcohol, Fluid (Forensic)					•	•		
8062TI	Postmortem, Expanded w/o Alcohol, Tissue (Forensic)					•	•		
8062U	Postmortem, Expanded w/o Alcohol, Urine (Forensic)					•	•		
8054B	Postmortem, Expanded with NPS, Blood (Forensic)					•	•		
8052B	Postmortem, Expanded, Blood (Forensic)					٠	•		
90025B	Postmortem, Expanded, Blood (Forensic) (CSA)					•	•		
8052FL	Postmortem, Expanded, Fluid (Forensic)						•		
8052SP	Postmortem, Expanded, Serum/Plasma (Forensic)					•	•		
8052TI	Postmortem, Expanded, Tissue (Forensic)					•	•		
8052U	Postmortem, Expanded, Urine (Forensic)					٠	•		
39052B	Postmortem, Expanded-II, Blood (Forensic) (SSA)						•		
39042B	Postmortem, Expanded-II, with Vitreous Alcohol Confirmation, Blood (Forensic) (SSA)						•		
8043B	Postmortem, Expert w/Vitreous Alcohol Confirmation, Blood (Forensic)					•	•		
10092B	Postmortem, Expert w/Vitreous Alcohol Confirmation, Blood (Forensic) (CSA)					•	•		
10151B	Postmortem, Expert w/Vitreous Alcohol Confirmation, Blood (Forensic) (CSA)					•	•		
8092B	Postmortem, Expert, Blood (Forensic)					•	•		
8092FL	Postmortem, Expert, Fluid (Forensic)					•	•		
8092SP	Postmortem, Expert, Serum/Plasma (Forensic)			•		٠	•		
8092TI	Postmortem, Expert, Tissue (Forensic)					•	•		



Test	Test Name	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
8092U	Postmortem, Expert, Urine (Forensic)					•	•		
4177B	Postmortem, SUIDS Screen, Blood (Forensic)						•		
52327B	Pyrrolidinophenone Confirmation, Blood								•
52327SP	Serum/Plasma								•
52494B	Substituted Cathinone Confirmation, Blood					•			
52494SP	Serum/Plasma					•			
52494U	Substituted Cathinone Confirmation, Urine					•			
52328B	Substituted Cathinone Panel Confirmation, Blood					•			
52328SP	Confirmation, Serum/Plasma					•			
52328U	Substituted Cathinone Panel Confirmation, Urine					•			
1021B	Substituted Cathinone Panel, Blood					•			
1021SP	Substituted Cathinone Panel, Serum/Plasma					•			
1021U	Substituted Cathinone Panel, Urine					•			
5970B	Synthetic Cannabinoids Confirmation (Qualitative), Blood					•			
9566B	Synthetic Cannabinoids Screen (Add- On), Blood					•			
9560B	Synthetic Cannabinoids Screen, Blood					•			



Test Changes

4000B	Amphetamines	s Confirmation (DUID/DRE), Blood
Sur	nmary of Changes:	Stability was changed. Scope of Analysis was changed. Ephedrine, Norpseudoephedrine, Phentermine, Phenylpropanolamine and Pseudoephedrine were removed.
N	Stability: Scope of Analysis: /ethod (CPT Code)	Room Temperature: 30 day(s) Refrigerated: 30 day(s) Frozen (-20 °C): 30 day(s) LC-MS/MS (80324, 80359): Amphetamine, Methamphetamine, MDA, MDMA, MDEA
54000U	Amphetamines	s Confirmation (Qualitative) (DUID/DRE), Urine
Sur	nmary of Changes:	Scope of Analysis was changed. Ephedrine, Norpseudoephedrine, Phentermine, Phenylpropanolamine and Pseudoephedrine were removed.
Ν	Scope of Analysis: lethod (CPT Code)	LC-MS/MS (80324, 80359): Amphetamine, Methamphetamine, MDA, MDMA, MDEA
54001B	Barbiturates C	Confirmation (DUID/DRE), Blood
Sur	nmary of Changes:	Specimen Requirements (Transport Temperature) were changed. Stability was changed. Scope of Analysis was changed. Amobarbital, Butabarbital, Pentobarbital and Secobarbital were removed.
Specir	nen Requirements:	2 mL Blood
Tran	sport Temperature:	Refrigerated
Sp	becimen Container:	Lavender top tube (EDTA)
	Light Protection:	Not Required
	Special Handling:	None
	Rejection Criteria:	None
	Stability:	Room Temperature: 14 day(s) Refrigerated: 14 day(s) Frozen (-20 °C): 12 month(s)
N	Scope of Analysis: lethod (CPT Code)	GC/MS (80345): Butalbital, Phenobarbital
	· · · · ·	
		Confirmation (Qualitative) (DUID/DRE), Urine
54001U		Confirmation (Qualitative) (DUID/DRE), Urine Scope of Analysis was changed. Amobarbital, Butabarbital, Pentobarbital and Secobarbital were removed.



54002B Benzodiaz	epines Confirmation (DUID/DI	RE), Blood			
Summary of Chang	Clobazam, Desalkylfluraze	anged. epam, Estazolam, Flurazepam, Hydroxytriazolam and Triazolam were removed.			
	de) Chlordiazepoxide, Lorazep	C-MS/MS (80347): Diazepam, Nordiazepam, Oxazepam, Temazepam, hlordiazepoxide, Lorazepam, Clonazepam, 7-Amino Clonazepam, Alprazolam, lpha-Hydroxyalprazolam, Midazolam			
54002U Benzodiaz	epines Confirmation (Qualitat	tive) (DUID/DRE), Urine			
Summary of Chang		epam, Estazolam, Hydroxyethylflurazepam and			
Scope of Analy Method (CPT Co	de) Chlordiazepoxide, Lorazep	-C-MS/MS (80339, 80347): Diazepam, Nordiazepam, Oxazepam, Temazepam, Chlordiazepoxide, Lorazepam, 7-Amino Clonazepam, Alprazolam, Alpha- Hydroxyalprazolam, 1-Hydroxymidazolam			
50013TI Cannabing	ids Confirmation (Qualitative), Tissue			
Summary of Chang	es: Test Name was changed.				
54456B DUID/DRE	Designer Benzodiazepines C	onfirmation, Blood			
Summary of Chang	es: Scope of Analysis was cha Diclazepam was removed				
Scope of Analy Method (CPT Co		azolam, Flubromazolam, Alpha-Hydroxyetizolam, Etizolam,			
54458B DUID/DRE	Designer Opioids Confirmation	on, Blood			
Summary of Chang	es: Scope of Analysis was cha Acrylfentanyl and Valerylfe Acryl Fentanyl and Valeryl	entanyl were added.			
Scope of Analy Method (CPT Co	de) Carfentanil, Butyrylfentany): 4-ANPP, Acrylfentanyl, 2-Furanylfentanyl, U-47700, /l, para-Fluoroisobutyrylfentanyl, cis-3-Methylfentanyl, -3-Methylfentanyl, Valerylfentanyl			
Analyte Name	Units	Reference Comment			
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.			
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.			



8152B DUID/DRE Exp	oanded Drug Screen Add-On ProofPOSITIVE®, Blood (Forensic)
Summary of Changes:	Specimen Requirements (Special Handling) were changed. Scope of Analysis was changed. Acrylfentanyl, Clobazam, Desalkylflurazepam, Ephedrine, Estazolam, Flurazepam, Hydroxyethylflurazepam, Hydroxytriazolam, Norpseudoephedrine, N-ethyl Pentylone, Phentermine, Phenylpropanolamine, Pseudoephedrine, Triazolam and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, Diclazepam, Maprotiline, Methaqualone, Mexiletine, N-Ethyl Pentylone, Trihexyphenidyl and Valeryl Fentanyl were removed.
Specimen Requirements:	10 mL Blood
Transport Temperature:	Frozen
Specimen Container:	Gray top tube (Sodium Fluoride / Potassium Oxalate), Lavender top tube (EDTA)
Light Protection:	Not Required
Special Handling:	Ensure that container remains tightly sealed.
Rejection Criteria:	Received Room Temperature. Received Refrigerated.
Scope of Analysis: Method (CPT Code)	ELISA (80307): Gabapentin ELISA (80307): Barbiturates LC/TOF-MS (80307): 2-Furanylfentanyl, 7-Amino Flunitrazepam, 9- Hydroxyrisperidone, 10-Hydroxycarbazepine, Acrylfentanyl, Alfentanil, Alpha- Hydroxyetizolam, Amitriptyline, Amoxapine, Aripiprazole, Brompheniramine, Bupropion, Buspirone, Butylone, Butyrylfentanyl, Caffeine, Carbamazepine, Carbamazepine-10,11-Epoxide, Carfentanil, Chlorpheniramine, Chlorpromazine, cis- 3-Methylfentanyl, Citalopram / Escitalopram, Clobazam, Clomipramine, Clonazolam, Clonidine, Clozapine, Cyclobenzaprine, Cyclopropylfentanyl, Delorazepam, Desalkylflurazepam, Desipramine, Desmethylclomipramine, Desmethyldoxepin, Desmethylsertraline, Desmethyltrimipramine, Dextro / Levo Methorphan, Dextrorphan / Levorphanol, Dicyclomine, Dibutylone, Diltiazem, Diphenhydramine, Doxepin, Doxylamine, Duloxetine, Ephedrine, Estazolam, Etizolam, Eszopiclone / Zopiclone, Flubromazolam, Flunitrazepam, Fluoxetine, Fluphenazine, Flurazepam, Fluvoxamine, Haloperidol, Hydroxybupropion, Hydroxyethylflurazepam, Hydroxytriazolam, Hydroxyzine, Iloperidone, Imipramine, Ketamine, Lamotrigine, Levetiracetam, Loxapine, LSD, mCPP, Meperidine, Mescaline, Mesoridazine, Metaxalone, Methocarbamol, Methylphenidate, Mirtazapine, Mitragynine, Norclozapine, Norflunitrazepam, Norfluoxetine, Norketamine, Normeperidine, Norpropoxyphene, Norpseudoephedrine, Nenterpilone, N-ethyl Pentylone, O- Desmethylvenlafaxine, Olanzapine, para-Fluoroisobutyrylfentanyl, Paroxetine, Perphenazine, Phenazepam, Pheniramine, Phentermine, Phenylpropanolamine, Phenytoin, Primidone, Promazine, Promethazine, Propoxyphene, Pseudoephedrine, Psilocin, Quetiapine, Risperidone, Sertraline, Sufentanil, Tapentadol, Thioridazine, Topiramate, trans-3-Methylfentanyl, Trazodone, Triazolam, Trifluoperazine, Trimipramine, U-47700, Valerylfentanyl, Venlafaxine, Verapamil, Xylazine, Zaleplon, Ziprasidone, Zonisamide, Scope Statement



Test Changes

Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
Clobazam Desalkylflurazepam Ephedrine Estazolam Flurazepam Hydroxyethylflurazepam Hydroxytriazolam mCPP	ng/mL ng/mL ng/mL ng/mL ng/mL ng/mL mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
Norpseudoephedrine N-ethyl Pentylone	ng/mL ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Phentermine Phenylpropanolamine Pseudoephedrine Triazolam Valerylfentanyl	ng/mL ng/mL ng/mL ng/mL ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.

8075U DUID/DRE Expanded Drug Screen Add-On, Urine (Forensic)

Summary of Changes:	Scope of Analysis was changed. Barbiturates, Acrylfentanyl, Clobazam, Desalkylflurazepam, Ephedrine, Estazolam, Hydroxyethylflurazepam, Hydroxytriazolam, Norpseudoephedrine, N-ethyl Pentylone, Phentermine, Phenylpropanolamine and Valerylfentanyl were added. Units were changed. Units were changed. Methods/CPT Codes were changed [EIA (80307)] Acryl Fentanyl, Bupivacaine, BZP, Diclazepam, Maprotiline, Methaqualone, Mexiletine, N-Ethyl Pentylone, Orphenadrine, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl and Zolazepam were removed.
	EIA (80307): Barbiturates LC/TOF-MS (80307): 2-Furanylfentanyl, 7-Amino Flunitrazepam, 9- Hydroxyrisperidone, 10-Hydroxycarbazepine, Acrylfentanyl, Alfentanil, Alpha- Hydroxyetizolam, Amitriptyline, Amoxapine, Atomoxetine, Benztropine, Bupropion, Brompheniramine, Butylone, Butyrylfentanyl, Buspirone, Caffeine, Carbamazepine, Carfentanil, Carbamazepine-10,11-Epoxide, Carisoprodol, cis-3-Methylfentanyl, Chlorpheniramine, Chlorpromazine, Citalopram / Escitalopram, Clobazam, Clomipramine, Clonidine, Clonazolam, Clozapine, Cyclobenzaprine,
MS Labs	



Test Updates

Test Changes

Cyclopropylfentanyl, Delorazepam, Desalkylflurazepam, Desipramine, Desmethylclomipramine, Desmethyldoxepin, Desmethyltrimipramine, Dextro / Levo Methorphan, Dextrorphan / Levorphanol, Dibutylone, Dicyclomine, Diltiazem, Diphenhydramine, Donepezil, Doxepin, Doxylamine, Ephedrine, Estazolam, Eszopiclone / Zopiclone, Etizolam, Flecainide, Flubromazolam, Flunitrazepam, Fluoxetine, Fluvoxamine, Guaifenesin, Hydroxybupropion, Hydroxyethylflurazepam, Hydroxytriazolam, Hydroxyzine, Imipramine, Ketamine, Lacosamide, Lamotrigine, Levetiracetam, LSD, mCPP, Memantine, Meperidine, Meprobamate, Mescaline, Mesoridazine, Metaxalone, Methylphenidate, Mirtazapine, Mitragynine, Norclozapine, Norflunitrazepam, Norfluoxetine, Norketamine, Normeperidine, Norpropoxyphene, Norpseudoephedrine, Nortriptyline, N-ethyl Pentylone, O-Desmethyltramadol, O-Desmethylvenlafaxine, Olanzapine, para-Fluoroisobutyrylfentanyl, Paroxetine, Phenazepam, Pheniramine, Phentermine, Phenylpropanolamine, Promazine, Promethazine, Propoxyphene, Pseudoephedrine, Psilocin, Quetiapine, Quinidine, Risperidone, Sertraline, Sildenafil, Sufentanil, Tapentadol, Tetrahydrozoline, Theophylline, Thioridazine, Topiramate, Tramadol, trans-3-Methylfentanyl, Trazodone, Trimipramine, Triprolidine, Valerylfentanyl, Venlafaxine, Verapamil, Xylazine, Yohimbine, Zaleplon, Zolpidem, Zonisamide, Scope Statement

	Scope Statement	
Analyte Name	Units	Reference Comment
Barbiturates	mcg/mL	
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Chlorpromazine	mcg/mL	
Clobazam	ng/mL	
Desalkylflurazepam	ng/mL	
Ephedrine	ng/mL	
Estazolam	ng/mL	
Hydroxyethylflurazepam	ng/mL	
Hydroxytriazolam	ng/mL	
mCPP	mcg/mL	
Norpseudoephedrine	ng/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Phentermine	ng/mL	
Phenylpropanolamine	ng/mL	
Valerylfentanyl	ng/mL	Valeryl fentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
151B DUID/DRE Par	nel (w/Alcohol) ProofPOSITIVE	®, Blood (Forensic)
Summary of Changes:	Scope of Analysis was change Units were changed. Barbiturates was removed	d.



Test Updates

Test Changes

Scope of Analysis: Method (CPT Code) Analyte Name	Amphetamines, Methadone Methamphetamine / MDMA, Fentanyl, Buprenorphine / M Headspace GC (80307): Eth Isopropanol, Acetone	caine / Metabolites, Benzodiazepines, Cannabinoids, / Metabolite, Phencyclidine, Carisoprodol / Metabolite, Oxycodone / Oxymorphone, Zolpidem, Fentanyl / Acetyl letabolite, Tramadol / Metabolite lanol, Blood Alcohol Concentration (BAC), Methanol, manol, Methanol, Isopropanol, Acetone Reference Comment				
Carisoprodol / Metabolite	mcg/mL					
8070U DUID/DRE Pa	nel (w/Alcohol), Urine (Foren	isic)				
Summary of Changes:	Scope of Analysis was changed. Barbiturates was removed.					
Scope of Analysis: Method (CPT Code)	EIA (80307): Opiates, Cocaine / Metabolites, Benzodiazepines, Cannabinoids, Methadone / Metabolite, Phencyclidine, Oxycodone / Oxymorphone Headspace GC (80307): Ethanol, Methanol, Isopropanol, Acetone Headspace GC (80320): Ethanol, Methanol, Isopropanol, Acetone EIA (80307): Amphetamines, MDMA, Buprenorphine / Metabolite, Fentanyl / Acetyl Fentanyl					
8150B DUID/DRE Pa	nel ProofPOSITIVE®, Blood	(Forensic)				
Summary of Changes:	Scope of Analysis was chang Units were changed. Barbiturates was removed.	ged.				
Scope of Analysis: Method (CPT Code)	Amphetamines, Methadone Methamphetamine / MDMA,	caine / Metabolites, Benzodiazepines, Cannabinoids, / Metabolite, Phencyclidine, Carisoprodol / Metabolite, Oxycodone / Oxymorphone, Zolpidem, Fentanyl / Acetyl letabolite, Tramadol / Metabolite				
Analyte Name	Units	Reference Comment				
Carisoprodol / Metabolite	mcg/mL					
8071U DUID/DRE Pa	nel, Urine (Forensic)					
Summary of Changes:	Scope of Analysis was chan Barbiturates was removed.	ged.				
	Methadone / Metabolite, Phe	ne / Metabolites, Benzodiazepines, Cannabinoids, encyclidine, Oxycodone / Oxymorphone , MDMA, Buprenorphine / Metabolite, Fentanyl / Acetyl				
90037B DUID/DRE Sc	reen, Blood (Forensic) (CSA)	- Maryland State Police				
Summary of Changes:	Units were changed.					



lest onanges		
Scope of Analy Method (CPT Co	ode) Amphetamines, Barbitu Metabolite, Methamphe	e, Cocaine / Metabolites, Benzodiazepines, Cannabinoids, rates, Methadone / Metabolite, Phencyclidine, Carisoprodol / tamine / MDMA, Oxycodone / Oxymorphone, Zolpidem, nyl, Buprenorphine / Metabolite, Tramadol / Metabolite
Analyte Name	Units	Reference Comment
Carisoprodol / Metaboli	ite mcg/mL	
54457B DUID/DRE	Substituted Cathinone Cor	ifirmation, Blood
Summary of Chang	ges: Scope of Analysis was on N-ethyl Pentylone was a	added.
Scope of Analy Method (CPT Co		ne, N-ethyl Pentylone, Dibutylone
Analyte Name	Units	Reference Comment
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
54457U DUID/DRE	Substituted Cathinone Cor	ifirmation, Urine
Summary of Chang	ges: Scope of Analysis was on N-ethyl Pentylone was a N-Ethyl Pentylone was a N-Ethyl Pentylone was a N-Ethyl Pentylone was a structure was a	added.
Scope of Analy Method (CPT Co		ne, N-ethyl Pentylone, Dibutylone
Analyte Name	Units	Reference Comment
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
54456U Designer E	Benzodiazepines Confirmat	ion (Qualitative) (DUID/DRE), Urine
Summary of Chang	ges: Test Name was change Scope of Analysis was o Diclazepam was remove	changed.
Scope of Analy Method (CPT Co		onazolam, Flubromazolam, Alpha-Hydroxyetizolam, Etizolam,
52493U Designer E	Benzodiazepines Confirmat	ion (Qualitative), Urine
Summary of Chang	ges: Scope of Analysis was o Diclazepam was remove	
		omazepam, Clonazolam, Flubromazolam, Alpha- am, Flubromazepam, Delorazepam
	, , , ,	



Test Changes

Summary of Changes:	Test Name was changed. Scope of Analysis was changed. Flualprazolam was added.	
		romazepam, Clonazolam, Flualprazolam, Flubromazolam, n, Etizolam, Flubromazepam, Delorazepam, Phenazepam,
Analyte Name	Units Reference Comment	
Flualprazolam	ng/mL	Flualprazolam is a benzodiazepine drug that is used as a novel psychoactive substance.

52502B Designer Benzodiazepines Confirmation 1, Blood

Summary of Changes:	: Test Name was changed. Scope of Analysis was changed. Flualprazolam was added.	
	LC-MS/MS (80346): Bromazepam, Clonazolam, Flualprazolam, Flubromazolam, Alpha-Hydroxyetizolam, Etizolam, Flubromazepam, Delorazepam, Phenazepam, Diclazepam	
Analyte Name	Units	Reference Comment
Flualprazolam	ng/mL	Flualprazolam is a benzodiazepine drug that is used as a novel psychoactive substance.

52502SP Designer Benzodiazepines Confirmation 1, Serum/Plasma

Summary of Changes:	 s: Test Name was changed. Scope of Analysis was changed. Flualprazolam was added. s: LC-MS/MS (80346): Bromazepam, Clonazolam, Flualprazolam, Flubromazolam, e) Alpha-Hydroxyetizolam, Etizolam, Flubromazepam, Delorazepam, Phenazepam, Diclazepam 	
Analyte Name	Units	Reference Comment
Flualprazolam	ng/mL	Flualprazolam is a benzodiazepine drug that is used as a novel psychoactive substance.

52493B Designer Benzodiazepines Confirmation, Blood

,	e) Hydroxyetizolam, Etizolam, Flubromazepam, Delorazepam enzodiazepines Confirmation, Serum/Plasma
Scope of Analys	is: LC-MS/MS (80346): Bromazepam, Clonazolam, Flubromazolam, Alpha-
Summary of Chang	es: Scope of Analysis was changed. Diclazepam was removed.



Test Changes

	Scope of Analysis was changed. Diclazepam was removed.		
	LC-MS/MS (80346): Bromazepam, Clonazolam, Flubromazolam, Alpha- Hydroxyetizolam, Etizolam, Flubromazepam, Delorazepam		
2488U Designer Opi	bids Confirmation (Qualitative), Urine		
Summary of Changes	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. Acryl Fentanyl, meta-Methylmethoxyacetylfentanyl, para- Fluorobutyrylfentanyl, para-Methylmethoxyacetylfentanyl, THF-F, U-49900, U-51754 and Valeryl Fentanyl were removed.		
Scope of Analysis Method (CPT Code)			
Analyte Name	Units	Reference Comment	
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.	
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.	
	ng/mL oids Confirmation, Blood	synthetic opioid.	
	oids Confirmation, Blood Scope of Analysis was of Acrylfentanyl and Valery 4-ANPP, Acryl Fentanyl	synthetic opioid. d changed. /Ifentanyl were added. , meta-Methylmethoxyacetylfentanyl, para- ara-Methylmethoxyacetylfentanyl, THF-F, U-49900,	
2488B Designer Opi	oids Confirmation, Blood : Scope of Analysis was of Acrylfentanyl and Valery 4-ANPP, Acryl Fentanyl, Fluorobutyrylfentanyl, pro- U-51754 and Valeryl Fe : LC-MS/MS (80354, 803) Fluorofentanyl, ortho-Flucyclopropylfentanyl, tra	synthetic opioid. d changed. /Ifentanyl were added. , meta-Methylmethoxyacetylfentanyl, para- ara-Methylmethoxyacetylfentanyl, THF-F, U-49900,	
2488B Designer Opi Summary of Changes Scope of Analysis	oids Confirmation, Blood : Scope of Analysis was of Acrylfentanyl and Valery 4-ANPP, Acryl Fentanyl, Fluorobutyrylfentanyl, pro- U-51754 and Valeryl Fe : LC-MS/MS (80354, 803) Fluorofentanyl, ortho-Flucyclopropylfentanyl, tra	synthetic opioid. d changed. /lfentanyl were added. , meta-Methylmethoxyacetylfentanyl, para- ara-Methylmethoxyacetylfentanyl, THF-F, U-49900, ntanyl were removed. 64): Methoxyacetylfentanyl, Acrylfentanyl, para- uorofentanyl, 2-Furanylfentanyl, U-47700, Carfentanil, ns-3-Methylfentanyl, cis-3-Methylfentanyl, Isobutyrylfentanyl,	
2488B Designer Opi Summary of Changes Scope of Analysis Method (CPT Code)	oids Confirmation, Blood Scope of Analysis was of Acrylfentanyl and Valery 4-ANPP, Acryl Fentanyl, Fluorobutyrylfentanyl, pr U-51754 and Valeryl Fe LC-MS/MS (80354, 803 Fluorofentanyl, ortho-Flu Cyclopropylfentanyl, tra Butyrylfentanyl, para-Flu	synthetic opioid. d changed. //fentanyl were added. / meta-Methylmethoxyacetylfentanyl, para- ara-Methylmethoxyacetylfentanyl, THF-F, U-49900, ntanyl were removed. 64): Methoxyacetylfentanyl, Acrylfentanyl, para- uorofentanyl, 2-Furanylfentanyl, U-47700, Carfentanil, ns-3-Methylfentanyl, cis-3-Methylfentanyl, Isobutyrylfentanyl, uoroisobutyrylfentanyl, Valerylfentanyl	



Test Changes

Summary of Changes:	Scope of Analysis was of Acrylfentanyl and Valery Acryl Fentanyl and Vale	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
2488SP Designer Opio	ids Confirmation, Serui	m/Plasma
Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. 4-ANPP, Acryl Fentanyl, meta-Methylmethoxyacetylfentanyl, para- Fluorobutyrylfentanyl, para-Methylmethoxyacetylfentanyl, THF-F, U-49900, U-51754 and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80354, 80364): Methoxyacetylfentanyl, Acrylfentanyl, para- Fluorofentanyl, ortho-Fluorofentanyl, 2-Furanylfentanyl, U-47700, Carfentanil, Cyclopropylfentanyl, trans-3-Methylfentanyl, cis-3-Methylfentanyl, Isobutyrylfentanyl, Butyrylfentanyl, para-Fluoroisobutyrylfentanyl, Valerylfentanyl	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
2500SP Designer Opio	ids Confirmation, Serui	m/Plasma
Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. Acryl Fentanyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80354, 80364): Acrylfentanyl, 2-Furanylfentanyl, U-47700, Carfentanil, Butyrylfentanyl, para-Fluoroisobutyrylfentanyl, cis-3-Methylfentanyl, Cyclopropylfentanyl, trans-3-Methylfentanyl, Valerylfentanyl	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.



Test Changes

Analyte Name	Units	Reference Comment
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
480B Designer Opio	ids, Blood	
Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. Acryl Fentanyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80354, 80364): Methoxyacetylfentanyl, 4-ANPP, THF-F, meta- Methylmethoxyacetylfentanyl, para-Methylmethoxyacetylfentanyl, Acrylfentanyl, para-Fluorofentanyl, ortho-Fluorofentanyl, 2-Furanylfentanyl, U-47700, U-49900, U- 51754, Carfentanil, Cyclopropylfentanyl, trans-3-Methylfentanyl, cis-3-Methylfentanyl, Isobutyrylfentanyl, Butyrylfentanyl, para-Fluoroisobutyrylfentanyl, para- Fluorobutyrylfentanyl, Valerylfentanyl	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
480SP Designer Opio	ids, Serum/Plasma	
Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. Acryl Fentanyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80354, 80364): Methoxyacetylfentanyl, 4-ANPP, THF-F, meta-	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
030B Drug Facilitate	ed Crime Panel, Blood (F	Forensic)

Summary of Changes: Units were changed.



Scope of Analysis: Method (CPT Code)	ELISA (80307): Cannabinoids, Barbiturates Headspace GC (80307): Ethanol, Blood Alcohol Concentration (BAC), Methanol, Isopropanol, Acetone GC/MS (80307): Gamma-Hydroxybutyric Acid LC/TOF-MS (80307): 6-Monoacetylmorphine, 7-Amino Clonazepam, 7-Amino Flunitrazepam, Acetyl Fentanyl, Alpha-Hydroxyalprazolam, Alprazolam, Amitriptyline, Amphetamine, Benzoylecgonine, Brompheniramine, Buprenorphine, Carisoprodol, Chlordiazepoxide, Chlorpheniramine, Citalopram / Escitalopram, Clobazam, Clonazepam, Clonidine, Cocaethylene, Cocaine, Codeine, Cyclobenzaprine, Desalkylflurazepam, Desipramine, Desmethyldoxepin, Desmethylsertraline, Dextro / Levo Methorphan, Dextrorphan / Levorphanol, Diazepam, Dihydrocodeine / Hydrocodol, Diphenhydramine, Doxepin, Doxylamine, EDDP, Estazolam, Eszopiclone / Zopiclone, Fentanyl, Flunitrazepam, Fluoxetine, Hydrocodone, Hydromorphone, Hydroxyethylflurazepam, Hydroxytriazolam, Imipramine, Ketamine, Lidocaine, Lorazepam, MDA, MDMA, Meperidine, Meprobamate, Methadone, Methamphetamine, Midazolam, Monoethylglycinexylidide (MEGX), Morphine, Norbuprenorphine - Free, Nordiazepam, Norfentanyl, Norflunitrazepam, Norfluoxetine, Norketamine, Normeperidine, Norpropoxyphene, Nortriptyline, O- Desmethyltramadol, Oxazepam, Oxycodone, Oxymorphone, Paroxetine, Phencyclidine, Phenytoin, Propoxyphene, Scopolamine, Sertraline, Temazepam, Tetrahydrozoline, Tramadol, Triazolam, Zaleplon, Ziprasidone, Zolpidem, Scope
	Tetrahydrozoline, Tramadol, Triazolam, Zaleplon, Ziprasidone, Zolpidem, Scope Statement

Analyte Name	Units	Reference Comment
Carisoprodol	mcg/mL	Following a 350 mg oral dose of carisoprodol, peak plasma concentrations averaged 2.1 mcg/mL in 1 hour. Following a 700 mg oral dose of carisoprodol, peak plasma concentrations averaged 3.5 mcg/mL in 0.8 hour.
Lidocaine	mcg/mL	Lidocaine is an amide type of anesthetic that is used as a topical and injectable analgesic, antiarrhythmic, and in resuscitative efforts. It is also used as a 'cutting' agent in some drugs of abuse, especially cocaine.
Meperidine Meprobamate	mcg/mL mcg/mL	Usual therapeutic range: 10-30 mcg/mL.
Monoethylglycinexylidide (MEGX)	mcg/mL	MEGX (monoethylglycinexylidide) is an active metabolite of lidocaine.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Norpropoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 1.45 mcg Norpropoxyphene/mL.



Analyte Name	Units	Reference Comment
Phenytoin	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with phenytoin: 10-20 mcg/mL The blood to plasma ratio is approximately 0.5.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
030SP Drug Facilitate	d Crime Panel, Serum/P	lasma (Forensic)
Summary of Changes:	Units were changed.	
Scope of Analysis: Method (CPT Code)	GC/MS (80307): Gamma LC/TOF-MS (80307): 6-M Flunitrazepam, Acetyl Fe Amphetamine, Benzoyled Chlordiazepoxide, Chlorp Clonazepam, Clonidine, G Desalkylflurazepam, Des Levo Methorphan, Dextro Hydrocodol, Diphenhydra Eszopiclone / Zopiclone, Hydromorphone, Hydroxy Lidocaine, Lorazepam, M Methamphetamine, Mida Norbuprenorphine - Free Norfluoxetine, Norketami Desmethyltramadol, Oxa Phencyclidine, Phenytoin	Ethanol, Methanol, Isopropanol, Acetone
Analyte Name	Units	Reference Comment
Carisoprodol	mcg/mL	Following a 350 mg oral dose of carisoprodol, peak plasma concentrations averaged 2.1 mcg/mL in 1 hour. Following a 700 mg oral dose of carisoprodol, peak plasma concentrations averaged 3.5 mcg/mL in 0.8 hour.
Lidocaine	mcg/mL	Lidocaine is an amide type of anesthetic that is used as a topical and injectable analgesic, antiarrhythmic, and in resuscitative efforts. It is also used as a 'cutting' agent in some drugs of abuse, especially cocaine.
		Reported antiarrhythmic range: 2-5 mcg/mL.



Test Changes

Analyte Name	Units	Reference Comment
Meprobamate	mcg/mL	Usual therapeutic range: 10-30 mcg/mL.
Monoethylglycinexylidide (MEGX)	mcg/mL	MEGX (monoethylglycinexylidide) is an active metabolite of lidocaine.
		Following a Lidocaine I.V. infusion at rates varying between 20 and 50 mcg/min/kg, steady-state MEGX serum concentrations range from 0.2-5.2 mcg/mL.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Norpropoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 1.45 mcg Norpropoxyphene/mL.
Phenytoin	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with phenytoin: 10-20 mcg/mL
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.

8030U Drug Facilitated Crime Panel, Urine (Forensic)

Summary of Changes:	Units were changed. Norbuprenorphine - Total was removed.
	EIA (80307): Cannabinoids, Barbiturates Headspace GC (80307): Ethanol, Methanol, Isopropanol, Acetone GC/MS (80307): Gamma-Hydroxybutyric Acid LC/TOF-MS (80307): 1-Hydroxymidazolam, 6-Monoacetylmorphine, 7-Amino Clonazepam, 7-Amino Flunitrazepam, Acetyl Fentanyl, Alpha-Hydroxyalprazolam, Alprazolam, Amitriptyline, Amphetamine, Benzoylecgonine, Brompheniramine, Buprenorphine, Carisoprodol, Chlordiazepoxide, Chlorpheniramine, Citalopram / Escitalopram, Clobazam, Clonidine, Cocaethylene, Cocaine, Codeine, Cyclobenzaprine, Desalkylflurazepam, Desipramine, Desmethyldoxepin, Dextro / Levo Methorphan, Dextrorphan / Levorphanol, Diazepam, Dihydrocodeine / Hydrocodol, Diphenhydramine, Doxepin, Doxylamine, EDDP, Estazolam, Eszopiclone / Zopiclone, Fentanyl, Flunitrazepam, Fluoxetine, Hydrocodone, Hydromorphone, Hydroxyethylflurazepam, Hydroxytriazolam, Imipramine, Ketamine, Lidocaine, Lorazepam, MDA, MDMA, Meperidine, Meprobamate, Methadone, Methamphetamine, Nonoethylglycinexylidide (MEGX), Morphine, Norbuprenorphine, Nordiazepam, Norfentanyl, Norflunitrazepam, Norfluoxetine, Norketamine, Normeperidine, Norpropoxyphene, Nortriptyline, O-Desmethyltramadol, Oxazepam, Oxycodone, Oxymorphone, Paroxetine, Phencyclidine, Phenytoin, Propoxyphene, Scopolamine, Sertraline, Temazepam, Tetrahydrozoline, Tramadol, Zaleplon, Zolpidem, Scope Statement



Analyte Name	Units	Reference Comment
Carisoprodol	mcg/mL	
Meperidine	mcg/mL	
Meprobamate	mcg/mL	
Normeperidine	mcg/mL	
Norpropoxyphene	mcg/mL	
Phenytoin	mcg/mL	
Propoxyphene	mcg/mL	
098B Drug Screen (GC/MS), Blood	
Summary of Changes:	N-ethyl Pentylone was a Units were changed.	
		amine, Trihexyphenidyl and Zolazepam were
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL).
		The ratio of whole blood concentration to plasma concentration is 1.0.



Units	Reference Comment
mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole.
	The blood to plasma ratio is not known for this analyte.
mcg/mL	 Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively. Mean plasma concentrations following maintenance doses: 200 mg/day: 4.99 +/- 2.51 mcg/mL; 400 mg/day: 9.35 +/- 4.22 mcg/mL; 600 mg/day: 12.46 +/- 5.60 mcg/mL. The ratio of whole blood concentration to plasma concentration is 1.1
mcg/mL mcg/mL	Peak Serum Concentrations (Single Oral Dose): 250 mg: 5.1 mcg/mL 1000 mg: 20 mcg/mL
	Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
	The blood to plasma ratio for metronidazole is unknown.
ng/mL	Steady-state peak plasma levels following a daily regimen of Venlafaxine occur at approximately 2.5 hours for O-Desmethylvenlafaxine: 94-200 ng/mL (75 mg/day), 85-472 ng/mL (150 mg/day), 243-515 ng/mL (225 mg/day), 390-1096 ng/mL (450 mg/day). Steady-state trough plasma levels following a 150 mg per day regimen:
	mcg/mL mcg/mL mcg/mL mcg/mL



Analyte Name	Units	Reference Comment
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were produced 0.5-2.0 hours after a 100 mg oral dose and peak serum concentrations of 3 mcg/mL were produced 6 min. following a 100 mg IV dose. Potentially toxic at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with primidone: 10-40 mcg/mL. The blood to plasma ratio is approximately 0.8.
Clonidine	ng/mL	Immediate-release, oral: 0.50-2.0 ng/mL, 2 hours after administration; Sustained-release, patch: 0.20-2.0 ng/mL, at steady-state; Sustained-release, oral: 0.20-0.27 ng/mL, 6.8 +/- 3.6 hours after a 0.1 mg single dose in healthy fed adults; children receive higher doses on a mg/kg basis.
		The ratio of whole blood concentration to serum or plasma concentration is unknown for this analyte.
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg (approximately 1600 mg) daily rufinamide resulted in plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133 patients maintained on twice-daily 400 or 800 mg doses for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.
		The blood to plasma ratio of rufinamide is approximately 1.0
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced a mean peak blood concentration of 2.0 mcg/mL (range, 1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3 mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours. Potentially toxic at blood concentrations greater than 8 mcg/mL.
Xylazine EDDP	ng/mL ng/mL	



Analyte Name	Units	Reference Comment
Hydroxychloroquine	ng/mL	Peak plasma concentrations of 410 +/- 130 ng/mL were achieved 2.4 hours after a single oral dose of 400 mg hydroxychloroquine (n = 6). Two cases of hydroxychloroquine overdose (20 g each) were successfully treated throughout cardiovascular collapse and had serum concentrations of 14000 and 26000 ng/mL. The ratio of whole blood concentration to serum or plasma concentration is unknown for this analyte.
Levamisole	mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole. There is limited data available on therapeutic concentrations of levamisole and no data on levamisole concentrations encountered from tainted cocaine. The mean peak plasma concentration following a single 2.5 mg/kg dose was 0.48 +/- 0.22 mcg/mL. Following a single 50 mg dose the mean peak plasma concentration was 0.13 mcg/mL. The ratio of whole blood concentration to plasma concentration is unknown for this analyte.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
Mescaline N-Acetylprocainamide	mcg/mL mcg/mL	The normal therapeutic range for NAPA is 10 to 20 mcg/mL plasma. The blood to plasma ratio is not known for this analyte.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Phencyclidine Procyclidine	ng/mL mcg/mL	Steady-state concentrations following chronic oral 10 to 30 mg dose: 0.15-0.63 mcg/mL.



	Units	Reference Comment
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Pyrimethamine	mcg/mL	A single oral dose of 50 mg given to 5 subjects produced a peak plasma concentration of 0.21-0.43 mcg/mL in 2 to 4 hours following the dose.
Quinidine	ng/mL	For the treatment of arrhythmia, effective plasma concentrations typically range between 2000 and 5000 ng/mL. The blood/plasma ratio is not known for quinidine, but concentrations in red blood cells are usually lower than plasma.
Quinine	ng/mL	A single oral 648 mg antispasmodic dose produces average peak plasma concentrations of 2800 ng/mL 2 hr after administration. The blood/plasma ratio is not known for quinine, but concentrations in red blood cells are usually lower than plasma.
Tocainide	mcg/mL	
098SP Drug Screen (0	GC/MS), Serum/Plasma	
Summary of Changes:	Scope of Analysis was cha N-ethyl Pentylone was ad Units were changed.	
	Pentylone, TFMPP, Tiletar removed.	nine, Trihexyphenidyl and Zolazepam were
Specimen Requirements:	removed.	nine, Trihexyphenidyl and Zolazepam were
Specimen Requirements: Transport Temperature:	removed. 10 mL Serum or Plasma	nine, Trihexyphenidyl and Zolazepam were
Transport Temperature: Specimen Container:	removed. 10 mL Serum or Plasma Refrigerated Gray top tube (Sodium Flu Plastic container (preserva	uoride / Potassium Oxalate), Lavender top tube (EDTA),
Transport Temperature: Specimen Container: Light Protection:	removed. 10 mL Serum or Plasma Refrigerated Gray top tube (Sodium Flu Plastic container (preserva Not Required	uoride / Potassium Oxalate), Lavender top tube (EDTA), ative-free)
Transport Temperature: Specimen Container:	removed. 10 mL Serum or Plasma Refrigerated Gray top tube (Sodium Flu Plastic container (preserva Not Required Serum: Collect sample in Plasma: Collect sample in Peak serum levels are red the blood drops so rapidly peak occurs at 40 to 90 m or Plasma into a plastic so	uoride / Potassium Oxalate), Lavender top tube (EDTA), ative-free) Red top tube Lavender top tube (EDTA) or Pink top tube. commended when monitoring patients because the level in that many negative results are found at the trough. The inutes post dose. Promptly centrifuge and separate Serum crew capped vial using approved guidelines.
Transport Temperature: Specimen Container: Light Protection:	removed. 10 mL Serum or Plasma Refrigerated Gray top tube (Sodium Flu Plastic container (preserva Not Required Serum: Collect sample in Plasma: Collect sample in Peak serum levels are rec the blood drops so rapidly peak occurs at 40 to 90 m	uoride / Potassium Oxalate), Lavender top tube (EDTA), ative-free) Red top tube Lavender top tube (EDTA) or Pink top tube. commended when monitoring patients because the level in that many negative results are found at the trough. The inutes post dose. Promptly centrifuge and separate Serum crew capped vial using approved guidelines.



Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL).
Fluconazole	mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole.
Lacosamide	mcg/mL	Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.
		Mean plasma concentrations following maintenance doses: 200 mg/day: 4.99 +/- 2.51 mcg/mL; 400 mg/day: 9.35 +/- 4.22 mcg/mL; 600 mg/day: 12.46 +/- 5.60 mcg/mL.
		NMS Labs derived data: 5th - 95th Percentile Data: 1.8-13.0 mcg/mL Mean: 5.3 mcg/mL (N = 14900)
Metharbital	mcg/mL	



Analyte Name	Units	Reference Comment
Metronidazole	mcg/mL	Peak Serum Concentrations (Single Oral Dose): 250 mg: 5.1 mcg/mL 1000 mg: 20 mcg/mL
		Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
O-Desmethylvenlafaxine	ng/mL	Steady-state peak plasma levels following a daily regimen of Venlafaxine occur at approximately 2.5 hours for O-Desmethylvenlafaxine: 94-200 ng/mL (75 mg/day), 85-472 ng/mL (150 mg/day), 243-515 ng/mL (225 mg/day), 390-1096 ng/mL (450 mg/day).
		Steady-state trough plasma levels following a 150 mg per day regimen: 65-300 ng O-Desmethylvenlafaxine/mL.
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were produced 0.5-2.0 hours after a 100 mg oral dose and peak serum concentrations of 3 mcg/mL were produced 6 min. following a 100 mg IV dose. Potentially toxic at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with primidone: 10-40 mcg/mL.
Clonidine	ng/mL	Immediate-release, oral: 0.50-2.0 ng/mL, 2 hours after administration; Sustained-release, patch: 0.20-2.0 ng/mL, at steady-state; Sustained-release, oral: 0.20-0.27 ng/mL, 6.8 +/- 3.6 hours after a 0.1 mg single dose in healthy fed adults; children receive higher doses on a mg/kg basis.
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg (approximately 1600 mg) daily rufinamide resulted in plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133 patients maintained on twice-daily 400 or 800 mg doses for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.



Analyte Name	Units	Reference Comment
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced a mean peak blood concentration of 2.0 mcg/mL (range, 1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3 mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours. Potentially toxic at blood concentrations greater than 8 mcg/mL.
Xylazine EDDP Hydroxychloroquine	ng/mL ng/mL ng/mL	Peak plasma concentrations of 410 +/- 130 ng/mL were achieved 2.4 hours after a single oral dose of 400 mg hydroxychloroquine (n = 6). Two cases of hydroxychloroquine overdose (20 g each) were successfully treated throughout cardiovascular collapse and had serum concentrations of 14000 and 26000 ng/mL.
Levamisole	mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole. There is limited data available on therapeutic concentrations of levamisole and no data on levamisole concentrations encountered from tainted cocaine. The mean peak plasma concentration following a single 2.5 mg/kg dose was 0.48 +/- 0.22 mcg/mL. Following a single 50 mg dose the mean peak plasma concentration was 0.13 mcg/mL.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days.
Mescaline N-Acetylprocainamide	mcg/mL mcg/mL	The normal therapeutic range for NAPA is 10 to 20 mcg/mL plasma.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Phencyclidine	ng/mL	



Test Changes

Analyte Name	Units	Reference Comment
Procyclidine	mcg/mL	Steady-state concentrations following chronic oral 10 to 30 mg dose: 0.15-0.63 mcg/mL.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Pyrimethamine	mcg/mL	A single oral dose of 50 mg given to 5 subjects produced a peak plasma concentration of 0.21-0.43 mcg/mL in 2 to 4 hours following the dose.
Quinidine	ng/mL	For the treatment of arrhythmia, effective plasma concentrations typically range between 2000 and 5000 ng/mL.
Quinine	ng/mL	A single oral 648 mg antispasmodic dose produces average peak plasma concentrations of 2800 ng/mL 2 hr after administration.
Tocainide	mcg/mL	Reported antiarrhythmic concentration: 4-10 mcg/mL. Tocainide is an antiarrhythmic drug that is no longer available in the Unites States.

8098U Drug Screen (GC/MS), Urine

removed.	Summary of Changes:	Scope of Analysis was changed. N-ethyl Pentylone was added. Units were changed. BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed.
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Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	The disposition of butalbital has not been well studied in humans.
Fluconazole	mcg/mL	
Lacosamide	mcg/mL	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy treatment of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day.



Test Changes

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Analyte Name	Units	Reference Comment
		Single labeled oral or intravenous lacosamide doses in healthy subjects were eliminated in urine (95%) and feces (< 0.5%) over a 7 day interval. Urinary excretion products included parent drug (40% of the dose) and the pharmacologically inactive O-desmethyllacosamide.
Metharbital N-Acetylprocainamide	mcg/mL mcg/mL	N-acetylprocainamide is an antiarrhythmic drug and an active metabolite of procainamide. The reported qualitative result for this substance was based upon a single analysis only. If confirmation testing is required please contact the laboratory.
Chlorpromazine Pentobarbital	mcg/mL mcg/mL	Less than 1% of a dose is eliminated in the urine as unchanged drug.
Clonidine Phenobarbital	ng/mL mcg/mL	For patients on chronic therapy, a mean of 20% (range, 12 - 55%) of the dose is excreted unchanged in the 24 hour urine.
Procainamide Procyclidine	mcg/mL mcg/mL	Procyclidine is an anticholinergic drug that was previously used in the treatment of Parkinson's disease.
Secobarbital	mcg/mL	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
Xylazine EDDP	ng/mL ng/mL	In maintenance subjects: Up to 50000 ng of Methadone plus Methadone Metabolites/mL Urine.
Hydroxychloroquine Levamisole	ng/mL mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole.
mCPP Mescaline N-ethyl Pentylone	mcg/mL mcg/mL ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Normeperidine Phencyclidine	mcg/mL ng/mL mcg/mL	



Analyte Name	Units	Reference Comment
Pyrimethamine Quinidine	mcg/mL ng/mL	
Quinine	ng/mL	Quinine is derived from the bark of the cinchona tree. It has been used in the past as an antimalarial, but is more commonly used today to treat muscle cramps. It is also used as a flavoring agent in tonic water and as a cutting agent in illicit heroin. Adverse effects include gastrointestinal disturbances, tinnitus, dizziness, arrhythmias and hypotension.
Tocainide	mcg/mL	Tocainide is an antiarrhythmic drug that is no longer available in the Unites States.
876B Drug Screen -	Expanded, Blood	
Summary of Changes:	Units were changed. Acryl Fentanyl, Diclazepam, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, N-Ethyl Pentylone, Trihexyphenidyl, Valeryl Fentanyl and Voriconazole were removed.	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
876FL Drug Screen -	Expanded, Fluid	
Summary of Changes:	Units were changed. BZP, Laudanosine, Metoclopramide, Mexiletine, TFMPP and Trihexyphenidyl were removed.	
Scope of Analysis: Method (CPT Code)		
	Units	Reference Comment
Analyte Name		



Lacosamide	mcg/mL	Lacosamide is a functionalized amino acid specifically
		synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Pentobarbital Phenobarbital Rufinamide Secobarbital Chlorpromazine Clonidine	mcg/mL mcg/mL mcg/mL mcg/mL ng/mL	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed. While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Xylazine EDDP Hydroxychloroquine mCPP Mescaline Normeperidine Phencyclidine Propoxyphene Quinidine Quinine	ng/mL ng/mL ng/mL mcg/mL mcg/mL ng/mL ng/mL ng/mL ng/mL ng/mL ng/mL	



Test Updates

Summary of Changes:	Units were changed. Acryl Fentanyl, BZP, Diclazepam, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, N- Ethyl Pentylone, TFMPP, Trihexyphenidyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days.
Xylazine	ng/mL	
1876U Drug Screen -	Expanded, Urine	
Summary of Changes:	Units were changed. Acryl Fentanyl, BZP, Diclazepam, Glipizide, Laudanosine, Metoclopramide, Mexiletine, N-Ethyl Pentylone, Norbuprenorphine - Total, TFMPP, Trihexyphenidyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Chlorpromazine mCPP Xylazine	mcg/mL mcg/mL ng/mL	
1864U Drugs of Abus	se Screen (10 Panel), Urine	
Summary of Changes:	Test Name was changed. Scope of Analysis was change Fentanyl / Acetyl Fentanyl was Methods/CPT Codes were cha	added.
		 / Metabolites, Benzodiazepines, Cannabinoids, Methadone / Metabolite, Phencyclidine, Oxycodone / Fentanyl
Analyte Name	Units	Reference Comment
Fentanyl / Acetyl Fentanyl	ng/mL	
1864B Drugs of Abus	se Screen (11 Panel), Blood	



Summary of Changes:	Test Name was changed. Scope of Analysis was change Fentanyl / Acetyl Fentanyl was	
	Amphetamines, Barbiturates, I	ine / Metabolites, Benzodiazepines, Cannabinoids, Fentanyl / Acetyl Fentanyl, Methadone / Metabolite, nine / MDMA, Oxycodone / Oxymorphone
Analyte Name	Units	Reference Comment
Fentanyl / Acetyl Fentanyl	ng/mL	
1864FL Drugs of Abus	e Screen (11 Panel), Fluid	
Summary of Changes:	Test Name was changed. Scope of Analysis was change Fentanyl / Acetyl Fentanyl was	
Scope of Analysis: Method (CPT Code)	Amphetamines, Barbiturates, I	ine / Metabolites, Benzodiazepines, Cannabinoids, Fentanyl / Acetyl Fentanyl, Methadone / Metabolite, nine / MDMA, Oxycodone / Oxymorphone
Analyte Name	Units	Reference Comment
Fentanyl / Acetyl Fentanyl	ng/mL	
1864SP Drugs of Abus	e Screen (11 Panel), Serum/P	lasma
Summary of Changes:	Test Name was changed. Scope of Analysis was change Fentanyl / Acetyl Fentanyl was	
Scope of Analysis: Method (CPT Code)	Amphetamines, Barbiturates, I	ine / Metabolites, Benzodiazepines, Cannabinoids, Fentanyl / Acetyl Fentanyl, Methadone / Metabolite, nine / MDMA, Oxycodone / Oxymorphone
Analyte Name	Units	Reference Comment
Fentanyl / Acetyl Fentanyl	ng/mL	
1864TI Drugs of Abus	e Screen (11 Panel), Tissue	
Summary of Changes:	Test Name was changed. Scope of Analysis was change Fentanyl / Acetyl Fentanyl was	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Fentanyl / Acetyl Fentanyl	ng/g	
90023B Expanded Dru	g Screen (DUID/DRE), Blood (Forensic) (CSA)



Test Updates

Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, Butorphanol, BZP, Diclazepam, Maprotiline, Methaqualone, Mexiletine, Nalbuphine, N-Ethyl Pentylone, TFMPP, Trihexyphenidyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	LC/TOF-MS (80307): 2-Furanylfentanyl, 9-Hydroxyrisperidone, 10-	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
52486U Fentanyl Pane	I Confirmation, Urine	
Summary of Changes:	Test Name was changed. Scope of Analysis was changed Fentanyl and Norfentanyl were	



Test Changes

Scope of Analysis: LC-MS/MS (80362): Acetyl Fentanyl, Fentanyl, Norfentanyl

Method (CPT Code)

Analyte Name	Units	Reference Comment
Fentanyl	ng/mL	Approximately 6% of dose is excreted in the urine as unchanged drug in 3 to 4 days.
Norfentanyl	ng/mL	
52486B Fentanyl and 4	4-ANPP Confirmation, Blo	od
Summary of Changes:	Test Name was changed. Scope of Analysis was ch Fentanyl, Norfentanyl and	
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80362): Fenta	anyl, Norfentanyl, 4-ANPP, Acetyl Fentanyl
Analyte Name	Units	Reference Comment
Fentanyl	ng/mL	Immediately following a single 2 mcg/kg I.V. dose: Up to 11 ng/mL, declining to 1 ng/mL after one hour. Following the application of a 100 mcg/hour transdermal patch, serum levels (after an initial lag time of approximately six hours) of 0.8-2.6 ng/mL were maintained for more than 24 hours after application. Peak plasma levels following a single oral transmucosal dose (Fentanyl Oralet) of 15 mcg/kg to children: 2-4 ng/mL at 20 minutes.
Norfentanyl 4-ANPP	ng/mL ng/mL	4-ANPP (despropionylfentanyl) is a precursor chemical used in the production of fentanyl/fentanyl related analytes and is also a fentanyl metabolite and may be a metabolite of other fentanyl-related analytes. It is considered to be pharmacologically weak.
52486SP Fentanyl and 4	4-ANPP Confirmation, Ser	um/Plasma
Summary of Changes:	Test Name was changed. Scope of Analysis was cha Fentanyl, Norfentanyl and	

Scope of Analysis: LC-MS/MS (80362): Fentanyl, Norfentanyl, 4-ANPP, Acetyl Fentanyl Method (CPT Code)



Test Changes

Analyte Name	Units	Reference Comment
Fentanyl	ng/mL	Immediately following a single 2 mcg/kg I.V. dose: Up to 11 ng/mL, declining to 1 ng/mL after one hour. Following the application of a 100 mcg/hour transdermal patch, serum levels (after an initial lag time of approximately six hours) of 0.8-2.6 ng/mL were maintained for more than 24 hours after application. Peak plasma levels following a single oral transmucosal dose (Fentanyl Oralet) of 15 mcg/kg to children: 2-4 ng/mL at 20 minutes.
Norfentanyl 4-ANPP	ng/mL ng/mL	4-ANPP (despropionylfentanyl) is a precursor chemical used in the production of fentanyl/fentanyl related analytes and is also a fentanyl metabolite and may be a metabolite of other fentanyl-related analytes. It is considered to be pharmacologically weak.
860B GC/MS Drug S	creen (Acid/Neutral), Blo	ood
Summary of Changes:	Units were changed. Etomidate was removed	
Scope of Analysis: Method (CPT Code)	GC/MS (80307): Acetaminophen, Acetohexamide, Barbital, Butalbital, Caffeine,	
Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.



Analyte Name	Units	Reference Comment
Cotinine	ng/mL	Cotinine concentrations from use of tobacco products and/or nicotine replacement therapy: 100-1200 ng/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL). The ratio of whole blood concentration to plasma
		concentration is 1.0.
Fluconazole	mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole.
		The blood to plasma ratio is not known for this analyte.
Lacosamide	mcg/mL	Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.
		Mean plasma concentrations following maintenance doses: 200 mg/day: 4.99 +/- 2.51 mcg/mL; 400 mg/day: 9.35 +/- 4.22 mcg/mL; 600 mg/day: 12.46 +/- 5.60 mcg/mL.
		The ratio of whole blood concentration to plasma concentration is 1.1
Metharbital	mcg/mL	



Test Changes

Analyte Name	Units	Reference Comment
Metronidazole	mcg/mL	Peak Serum Concentrations (Single Oral Dose): 250 mg: 5.1 mcg/mL 1000 mg: 20 mcg/mL
		Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
		The blood to plasma ratio for metronidazole is unknown.
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were produced 0.5-2.0 hours after a 100 mg oral dose and peak serum concentrations of 3 mcg/mL were produced 6 min. following a 100 mg IV dose. Potentially toxic at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with primidone: 10-40 mcg/mL. The blood to plasma ratio is approximately 0.8.
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg (approximately 1600 mg) daily rufinamide resulted in plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133 patients maintained on twice-daily 400 or 800 mg doses for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.
		The blood to plasma ratio of rufinamide is approximately 1.0
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced a mean peak blood concentration of 2.0 mcg/mL (range, 1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3 mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours. Potentially toxic at blood concentrations greater than 8 mcg/mL.
Xylazine	ng/mL	
0053U GC/MS Drug S	creen, Urine (CSA)	
Summary of Changes:	Units were changed. Etomidate was removed.	
Scope of Analysis:		

Method (CPT Code)



Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	The disposition of butalbital has not been well studied in humans.
Fluconazole Chlorpromazine Methapyrilene Metharbital	mcg/mL mcg/mL mcg/mL mcg/mL	
N-Acetylprocainamide	mcg/mL	N-acetylprocainamide is an antiarrhythmic drug and an active metabolite of procainamide. The reported qualitative result for this substance was based upon a single analysis only. If confirmation testing is required please contact the laboratory.
Pentobarbital	mcg/mL	Less than 1% of a dose is eliminated in the urine as unchanged drug.
Phenobarbital	mcg/mL	For patients on chronic therapy, a mean of 20% (range, 12 - 55%) of the dose is excreted unchanged in the 24 hour urine.
Procainamide Procyclidine	mcg/mL mcg/mL	Procyclidine is an anticholinergic drug that was previously used in the treatment of Parkinson's disease.
EDDP	ng/mL	In maintenance subjects: Up to 50000 ng of Methadone plus Methadone Metabolites/mL Urine.
Secobarbital	mcg/mL	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
Xylazine Hydroxychloroquine Levamisole	ng/mL ng/mL mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole.
Mescaline Normeperidine Phencyclidine Propoxyphene Pyrimethamine Quinidine	mcg/mL mcg/mL ng/mL mcg/mL mcg/mL ng/mL	



Test Changes

Horsham, PA 19044-2208

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Analyte Name	Units	Reference Comment
Quinine	ng/mL	Quinine is derived from the bark of the cinchona tree. It has been used in the past as an antimalarial, but is more commonly used today to treat muscle cramps. It is also used as a flavoring agent in tonic water and as a cutting agent in illicit heroin. Adverse effects include gastrointestinal disturbances, tinnitus, dizziness, arrhythmias and hypotension.
Tiletamine Tocainide	mcg/mL mcg/mL	Tocainide is an antiarrhythmic drug that is no longer available in the Unites States.
Zolazepam	mcg/mL	
	ctive Substances (NPS	S) Screen 1, Blood
	alpha-PiHP, Benzylone, Flualprazolam, N-butyl Valerylfentanyl were ad 25B-NBOMe, 25C-NBC Fluorophenmetrazine, 4 Deschloroetizolam, MD Methylmethoxyacetylfer MPHP, N-Ethyl Pentylor Methylmethoxyacetylfer	nine, 3-hydroxy-PCP, Acrylfentanyl, alpha-PHP / , Deschloroketamine, Eutylone, Fentanyl, Pentylone, N-ethyl Pentylone, Norfentanyl and
Method (CPT Code)	3-MeO-PCP, 4-ANPP, A PHP / alpha-PiHP, alpha Carfentanil, cis-3-Methy Deschloroketamine, Dit Flualprazolam, Flubrom Methoxyacetylfentanyl, Norfentanyl, ortho-Fluor	fluoro Deschloroketamine, 2-Furanylfentanyl, 3-hydroxy-PCP, Acetyl Fentanyl, Acrylfentanyl, Alpha-Hydroxyetizolam, alpha- a-PVP, Benzylone, Bromazepam, Butylone, Butyrylfentanyl, vlfentanyl, Clonazolam, Cyclopropylfentanyl, Delorazepam, butylone, Diclazepam, Ethylone, Etizolam, Eutylone, Fentanyl, nazepam, Flubromazolam, Isobutyrylfentanyl, Mitragynine, N-butyl Pentylone, N-ethyl Pentylone, rofentanyl, para-Fluorofentanyl, para-Fluoroisobutyrylfentanyl, m, trans-3-Methylfentanyl, U-47700, Valerylfentanyl
Analyte Name	Units	Reference Comment
2-fluoro Deschloroketamine 3-hydroxy-PCP Acrylfentanyl	ng/mL ng/mL ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
	ng/mL	



Analyte Name	Units	Reference Comment
Eutylone	ng/mL	Eutylone is classified as a synthetic stimulant and belongs to the beta-keto methylenedioxyamphetamine subclass, which includes synthetic stimulants methylone, butylone, ethylone, and N-ethylpentylone.
Fentanyl Flualprazolam N-butyl Pentylone N-ethyl Pentylone	ng/mL ng/mL ng/mL ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norfentanyl Valerylfentanyl	ng/mL ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
756SP Novel Psychoa	active Substances (NPS	S) Screen 1, Serum/Plasma
Summary of Changes: Scope of Analysis: Method (CPT Code)	Eutylone, Fentanyl, Flua Norfentanyl, Valerylfent were added. 25B-NBOMe, 25C-NBC Fluorophenmetrazine, 4 Deschloroetizolam, MD Methylmethoxyacetylfer MPHP, N-Ethyl Pentylor Methylmethoxyacetylfer 49900, U-51754, Valery LC/TOF-MS (80307): 2- 3-MeO-PCP, 4-ANPP, A PHP / alpha-PiHP, alpha Carfentanil, cis-3-Methy Deschloroketamine, Dib Flualprazolam, Flubrom Methoxyacetylfentanyl,	changed. P / alpha-PiHP, Benzylone, Deschloroketamine, alprazolam, N-butyl Pentylone, N-ethyl Pentylone, anyl, 2-fluoro Deschloroketamine and 3-hydroxy-PCP Me, 25H-NBOMe, 25I-NBOMe, 3- H-MeO-PCP, Acryl Fentanyl, BZP, Clephedrone, PV, Meclonazepam, Mephedrone, meta- ntanyl, Methoxetamine, Methoxphenidine, Methylone, ne, para-Fluorobutyrylfentanyl, para- ntanyl, Pentedrone, Pyrazolam, TFMPP, THF-F, U- 1 Fentanyl were removed. fluoro Deschloroketamine, 2-Furanylfentanyl, 3-hydroxy-PCP, acetyl Fentanyl, Acrylfentanyl, Alpha-Hydroxyetizolam, alpha- a-PVP, Benzylone, Bromazepam, Butylone, Butyrylfentanyl, vifentanyl, Clonazolam, Cyclopropylfentanyl, Delorazepam, butylone, Diclazepam, Ethylone, Etizolam, Eutylone, Fentanyl, Mitragynine, N-butyl Pentylone, N-ethyl Pentylone, rofentanyl, para-Fluorofentanyl, para-Fluoroisobutyrylfentanyl,
	Pentylone, Phenazepar	n, trans-3-Methylfentanyl, U-47700, Valerylfentanyl
Analyte Name	Units	Reference Comment
2-fluoro Deschloroketamine 3-hydroxy-PCP Acrylfentanyl	ng/mL ng/mL ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
alpha-PHP / alpha-PiHP Benzylone Deschloroketamine	ng/mL ng/mL ng/mL	



Test Changes

Analyte Name	Units	Reference Comment
Eutylone	ng/mL Eutylone is classified as a synthetic stimulant and belongs to the beta-keto-methylenedioxyamphetamir subclass, which includes synthetic stimulants methylone, butylone, ethylone, and N-ethylpentylone	
Fentanyl	ng/mL	
Flualprazolam	ng/mL	
N-butyl Pentylone	ng/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norfentanyl	ng/mL	
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
756U Novel Psychoa	ctive Substances (NPS	S) Screen 1, Urine
Summary of Changes:	Scope of Analysis was of 2-fluoro Deschloroketar alpha-PiHP, Benzylone, Flualprazolam, N-butyl Valerylfentanyl were ad 25B-NBOMe, 25C-NBC Fluorophenmetrazine, 4 Deschloroetizolam, MD Methylmethoxyacetylfer MPHP, N-Ethyl Pentylon Methylmethoxyacetylfer	nine, 3-hydroxy-PCP, Acrylfentanyl, alpha-PHP / , Deschloroketamine, Eutylone, Fentanyl, Pentylone, N-ethyl Pentylone, Norfentanyl and
	3-MeO-PCP, 4-ANPP, A PHP / alpha-PiHP, alpha Carfentanil, cis-3-Methy Deschloroketamine, Dib Flualprazolam, Flubrom Methoxyacetylfentanyl, Norfentanyl, ortho-Fluor	
Analyte Name	Units	Reference Comment
2-fluoro Deschloroketamine	ng/mL	
3-hydroxy-PCP	ng/mL	
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription
-	-	synthetic opioid

Acrylfentanyl is a novel non-prescription synthetic opioid.



Test Changes

Analyte Name	Units	Reference Comment
alpha-PHP / alpha-PiHP	ng/mL	
Benzylone	ng/mL	
Deschloroketamine	ng/mL	
Eutylone	ng/mL	
Fentanyl	ng/mL	
Flualprazolam	ng/mL	
N-butyl Pentylone	ng/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norfentanyl	ng/mL	
Valerylfentanyl	ng/mL	Valeryl fentanyl is a novel non-prescription synthetic opioid.
90036U Opioids Panel	, Urine (CSA)	
Summary of Changes:	Scope of Analysis was changed. Norbuprenorphine was added. Norbuprenorphine - Total was removed.	
Scope of Analysis: Method (CPT Code)	Dihydrocodeine / Hydroc Meperidine, Methadone	Monoacetylmorphine, Buprenorphine, Butorphanol, Codeine, codol, EDDP, Fentanyl, Hydrocodone, Hydromorphone, , Morphine, Norbuprenorphine, Norfentanyl, Normeperidine, Oxycodone, Oxymorphone, Tapentadol, Tramadol
Analyte Name	Units	Reference Comment
Norbuprenorphine	ng/mL	
90035U Opioids Scree	n, Urine (CSA)	
Summary of Changes:	Scope of Analysis was c Norbuprenorphine was a Norbuprenorphine - Tota	added.
Scope of Analysis: Method (CPT Code)	Dihydrocodeine / Hydroc Meperidine, Methadone,	Monoacetylmorphine, Buprenorphine, Butorphanol, Codeine, codol, EDDP, Fentanyl, Hydrocodone, Hydromorphone, , Morphine, Norbuprenorphine, Norfentanyl, Normeperidine, Oxycodone, Oxymorphone, Tapentadol, Tramadol
Analyte Name	Units	Reference Comment
Norbuprenorphine	ng/mL	
8155U Postmortem D	esigner Opioids Add-Or	n (Qualitative), Urine (Forensic)
Summary of Changes:	Scope of Analysis was c Acrylfentanyl and Valery Acryl Fentanyl and Valer	

Effective Date: Monday, August 15, 2022



Scope of Analysis: Method (CPT Code)	Methylmethoxyacetylfenta para-Fluorofentanyl, ortho 51754, Carfentanil, Cyclo	4): Methoxyacetylfentanyl, 4-ANPP, THF-F, meta- anyl, para-Methylmethoxyacetylfentanyl, Acrylfentanyl, p-Fluorofentanyl, 2-Furanylfentanyl, U-47700, U-49900, U- propylfentanyl, trans-3-Methylfentanyl, cis-3-Methylfentanyl, lfentanyl, para-Fluoroisobutyrylfentanyl, para- erylfentanyl
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
8155B Postmortem D	esigner Opioids Add-On,	Blood (Forensic)
Summary of Changes:	Scope of Analysis was ch Acrylfentanyl and Valerylf Acryl Fentanyl and Valery	entanyl were added.
Scope of Analysis: Method (CPT Code)	LC-MS/MS (80354, 80364): Methoxyacetylfentanyl, 4-ANPP, THF-F, meta- Methylmethoxyacetylfentanyl, para-Methylmethoxyacetylfentanyl, Acrylfentanyl, para-Fluorofentanyl, ortho-Fluorofentanyl, 2-Furanylfentanyl, U-47700, U-49900, U- 51754, Carfentanil, Cyclopropylfentanyl, trans-3-Methylfentanyl, cis-3-Methylfentanyl, Isobutyrylfentanyl, Butyrylfentanyl, para-Fluoroisobutyrylfentanyl, para- Fluorobutyrylfentanyl, Valerylfentanyl	
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
8155SP Postmortem D	esigner Opioids Add-On,	Serum/Plasma (Forensic)
Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl and Valerylfentanyl were added. Acryl Fentanyl and Valeryl Fentanyl were removed.	
Scope of Analysis: Method (CPT Code)	Methylmethoxyacetylfenta para-Fluorofentanyl, ortho 51754, Carfentanil, Cyclo	4): Methoxyacetylfentanyl, 4-ANPP, THF-F, meta- anyl, para-Methylmethoxyacetylfentanyl, Acrylfentanyl, p-Fluorofentanyl, 2-Furanylfentanyl, U-47700, U-49900, U- propylfentanyl, trans-3-Methylfentanyl, cis-3-Methylfentanyl, lfentanyl, para-Fluoroisobutyrylfentanyl, para- erylfentanyl



Test Changes

Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
8063B Postmortem, E	Basic to Expanded Upgrad	le, Blood (Forensic)
Summary of Changes:	Norpropoxyphene, Propoxy Pentylone were added. Units were changed. Acryl Fentanyl, BZP, Diclaz Glyburide, Itraconazole, Ke Mexiletine, Monoethylglyci	unged. yphene, Valerylfentanyl, Acrylfentanyl and N-ethyl zepam, Etomidate, Glimepiride, Glipizide, etoconazole, Laudanosine, Metoclopramide, nexylidide (MEGX), N-Ethyl Pentylone, TFMPP, /l, Valeryl Fentanyl, Voriconazole and Zolazepam
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norpropoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 1.45 mcg Norpropoxyphene/mL.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.

8063FL Postmortem, Basic to Expanded Upgrade, Fluid (Forensic)



Test Changes

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Summary of Changes:	Scope of Analysis was changed.
	Units were changed.
	BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, TFMPP,
	Tiletamine, Trihexyphenidyl and Zolazepam were removed.

Scope of Analysis: Method (CPT Code)

mcg/mL mcg/mL mcg/mL mcg/mL	
	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The
	recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
mcg/mL mcg/mL ng/mL	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed.
	While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
ng/mL ng/mL mcg/mL mcg/mL mcg/mL mcg/mL ng/mL ng/mL	
	ng/mL ng/mL ng/mL ng/mL mcg/mL mcg/mL mcg/mL mcg/mL mcg/mL



lest changes		
8063SP Postmor	tem, Basic to Expanded Up	ograde, Serum/Plasma (Forensic)
Summary of Cha	Acrylfentanyl, N-ethyl Valerylfentanyl were a Units were changed. Acryl Fentanyl, BZP, I Glyburide, Itraconazo Mexiletine, Monoethy	Pentylone, Norpropoxyphene, Propoxyphene and
Scope of Ana Method (CPT (
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norpropoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 1.45 mcg Norpropoxyphene/mL.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
	tem, Basic to Expanded Up	ograde, Tissue (Forensic)
Summary of Cha	Units were changed. BZP, Etomidate, Lauc	s changed. danosine, Metoclopramide, Mexiletine, TFMPP, enidyl and Zolazepam were removed.
Scope of Ana Method (CPT (
Analyte Name	Units	Reference Comment
Ethinamate	mcg/g	
Felbamate	mcg/g	
Fluconazole	mcg/g	
NMS Labs 200 Welsh Rd.		



Analyte Name	Units	Reference Comment
Lacosamide	mcg/g	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Rufinamide	mcg/g	
Clonidine	ng/g	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed.
		While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Xylazine Hydroxychloroquine mCPP	ng/g ng/g mcg/g	
Mescaline Normeperidine	mcg/g mcg/g	
Propoxyphene	mcg/g	
Quinidine Quinine	ng/g ng/g	
Trimipramine	ng/g	
063U Postmortem, E	Basic to Expanded Upg	rade, Urine (Forensic)
Summary of Changes:	Valerylfentanyl were ad Units were changed. Acryl Fentanyl, BZP, Die Mexiletine, Monoethylgl	entylone, Norpropoxyphene, Propoxyphene and



Test Changes

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Chlorpromazine	mcg/mL	
mCPP	mcg/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norpropoxyphene	mcg/mL	
Propoxyphene	mcg/mL	
Valerylfentanyl	ng/mL	Valeryl fentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
084B Postmortem, E	Expanded w/ Vitreous A	Icohol and 6-MAM Confirmation, Blood (Forensic)
Summary of Changes:	Units were changed. Acryl Fentanyl, BZP, Dic Glyburide, Itraconazole, Mexiletine, Monoethylgh	changed. entylone and Valerylfentanyl were added. clazepam, Etomidate, Glimepiride, Glipizide, , Ketoconazole, Laudanosine, Metoclopramide, ycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, idyl, Valeryl Fentanyl, Voriconazole and Zolazepam
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in
		blood which may be dependent upon pH, collection tube,
		and storage temperature.
		Negative results should be interpreted with caution.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma
	5	overeged 0.02 meg/ml, et enprevimetely 8 heurs neet dese

	mog/me	averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.

10052B Postmortem, Expanded w/Vitreous Alcohol Confirmation, Blood (Forensic) (CSA)



Test Changes

Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole, Zolazepam and were removed.
Scope of Analysis:	

Method (CPT Code) **Analyte Name** Units **Reference Comment** Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown. N-ethyl Pentylone ng/mL N-ethyl Pentylone is a novel psychoactive stimulant. Valerylfentanyl ng/mL Valerylfentanyl is a novel non-prescription synthetic opioid. 8042B Postmortem, Expanded w/Vitreous Alcohol Confirmation, Blood (Forensic) Seene of Analysis was shanged

Summar	ry of Changes:	Scope of Analysis was changed.
		Valerylfentanyl, Acrylfentanyl and N-ethyl Pentylone were added.
		Units were changed.
		Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide,
		Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide,
		Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP,
		Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam
		were removed.
Sco	pe of Analysis:	

Method (CPT Code)

Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.



Test Changes

Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
8057B Postmortem, B (CSA)	Expanded w/Vitreous Alc	ohol Confirmation, Blood - University of MI (Forensic)
Summary of Changes:	Acrylfentanyl, N-ethyl Per Units were changed. Acryl Fentanyl, BZP, Dicla Glyburide, Itraconazole, H Mexiletine, Monoethylglyd	nanged. ntylone and Valerylfentanyl were added. azepam, Etomidate, Glimepiride, Glipizide, Ketoconazole, Laudanosine, Metoclopramide, cinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, dyl, Valeryl Fentanyl, Voriconazole and Zolazepam
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
8062B Postmortom	Expanded w/o Alcohol B	lood (Earoncia)

8062B Postmortem, Expanded w/o Alcohol, Blood (Forensic)



Scope of Analysis:

Test Updates

Test Changes

Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam
	were removed.

Method (CPT Code) **Analyte Name** Units **Reference Comment** Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown. N-ethyl Pentylone ng/mL N-ethyl Pentylone is a novel psychoactive stimulant. Valerylfentanyl ng/mL Valerylfentanyl is a novel non-prescription synthetic opioid.

8062FL Postmortem, Expanded w/o Alcohol, Fluid (Forensic)

Summary of Changes: Scope of Analysis was changed. Units were changed. BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed.

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment	
Butalbital	mcg/mL		
Ethinamate	mcg/mL		
Felbamate	mcg/mL		
Fluconazole	mcg/mL		



Synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to maintenance dose of 200 to 400 mg/day. Patients ex to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.Pentobarbitalmcg/mL mcg/mLPentobarbitalmcg/mL mcg/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLClonidineng/mLKylazineng/mLKylazineng/mLPPPng/mLHydroxychloroquineng/mLNormeperidineng/mLPropoxyhenemcg/mLPropoxyhenemcg/mL	Analyte Name	Units	Reference Comment
Phenobarbital mcg/mL Rufinamide mcg/mL Secobarbital mcg/mL Clonidine ng/mL Secobarbital Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommend daily oral dose ranges from a minimum of 0.2 mg to maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed. While overdoses from clonidine are common, deaths rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somolence, and hypoventilation Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death. Xylazine ng/mL Hydroxychloroquine ng/mL McScaline mcg/mL Mescaline mcg/mL Normeperidine ng/mL Propoxyphene mcg/mL	Lacosamide	mcg/mL	of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended
EDDPng/mLHydroxychloroquineng/mLmCPPmcg/mLMescalinemcg/mLNormeperidinemcg/mLPhencyclidineng/mLPropoxyphenemcg/mL	Phenobarbital Rufinamide Secobarbital Chlorpromazine	mcg/mL mcg/mL mcg/mL mcg/mL	 activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed. While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma
Quinidine ng/mL Quinine ng/mL	EDDP Hydroxychloroquin mCPP Mescaline Normeperidine Phencyclidine Propoxyphene Quinidine	ng/mL e ng/mL mcg/mL mcg/mL mcg/mL ng/mL mcg/mL ng/mL	



Test Changes

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Summary of Changes:	Scope of Analysis was changed. Units were changed.
	BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed.

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment
Butalbital Ethinamate Felbamate Fluconazole Lacosamide	mcg/g mcg/g mcg/g mcg/g	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Pentobarbital Phenobarbital Rufinamide Secobarbital	mcg/g mcg/g mcg/g mcg/g	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
Clonidine	ng/g	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed.
		While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Xylazine EDDP	ng/g ng/g	
S Labs Welsh Rd. sham, PA 19044-2208		Dego 55 of 97



Test Changes

Analyte Name	Units	Reference Comment
Hydroxychloroquine	ng/g	
mCPP	mcg/g	
Mescaline	mcg/g	
Normeperidine	mcg/g	
Phencyclidine	ng/g	
Propoxyphene	mcg/g	
Quinidine	ng/g	
Quinine	ng/g	
Trimipramine	ng/g	
062U Postmortem, E	Expanded w/o Alcohol,	Urine (Forensic)
Summary of Changes:	Scope of Analysis was	changed.
		Pentylone, Norbuprenorphine and Valerylfentanyl were
	added.	
	Units were changed.	
		iclazepam, Glipizide, Laudanosine, Metoclopramide,
		lycinexylidide (MEGX), N-Ethyl Pentylone,
		tal, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl
	Fentanyl and Zolazepa	
Scope of Analysis:	· · · · ·	
Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Chlorpromazine	mcg/mL	
mCPP	mcg/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
	-	
Norbuprenorphine	ng/mL	
Valerylfentanyl	ng/mL	Valeryl fentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
•	Expanded with NPS, Bl	ood (Forensic)
,	,	
Summary of Changes:	2-fluoro Deschloroketa alpha-PiHP, Benzylone Pentylone, N-ethyl Pen Units were changed.	mine, 3-hydroxy-PCP, Acrylfentanyl, alpha-PHP / e, Deschloroketamine, Eutylone, Flualprazolam, N-butyl htylone and Valerylfentanyl were added.
	-	OMe, 25H-NBOMe, 25I-NBOMe, 3-

- Fluorophenmetrazine, 4-MeO-PCP, 5-fluoro-MDMB-PICA, Acryl Fentanyl,
- BZP, Clephedrone, Deschloroetizolam, Etomidate, Glimepiride, Glipizide,
- Glyburide, Itraconazole, Ketoconazole, Laudanosine, MDPV, Meclonazepam,



Scope of Analysis:

Test Updates

Test Changes

Mephedrone, meta-Methylmethoxyacetylfentanyl, Methoxetamine, Methoxphenidine, Methylone, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), MPHP, N-Ethyl Pentylone, para-Fluorobutyrylfentanyl, para-Methylmethoxyacetylfentanyl, Pentedrone, Pyrazolam, TFMPP, THF-F, Tiletamine, Trihexyphenidyl, U-49900, U-51754, Valeryl Fentanyl, Voriconazole and Zolazepam were removed.

Method (CPT Code)		
Analyte Name	Units	Reference Comment
2-fluoro Deschloroketamine 3-hydroxy-PCP Acrylfentanyl	ng/mL ng/mL ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
alpha-PHP / alpha-PiHP Benzylone Deschloroketamine Eutylone	ng/mL ng/mL ng/mL ng/mL	Eutylone is classified as a synthetic stimulant and belongs to the beta-keto methylenedioxyamphetamine subclass, which includes synthetic stimulants methylone, butylone, ethylone, and N-ethylpentylone.
Flualprazolam mCPP	ng/mL mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-butyl Pentylone N-ethyl Pentylone	ng/mL ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.

90025B Postmortem, Expanded, Blood (Forensic) (CSA)

Summary of Changes:	Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed.
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Test Changes

Scope of Analysis: Method (CPT Code)

blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/. 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown. N-ethyl Pentylone ng/mL N-ethyl Pentylone is a novel psychoactive stimulant. Valerylfentanyl ng/mL Valerylfentanyl is a novel non-prescription synthetic opioid. S2E Postmortem, Expanded, Blood (Forensic) Summary of Changes: Scope of Analysis was changed. Acrylfentanyl, N=ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexyllidide (MEGX), N=Ethyl Pentylone, TFMPP, Titletamine, Trihexyphenidyl, Valeryl Fentanyl, Noriconazole and Zolazepam were removed. Scope of Analysis: Method (CPT Code) Monte Vision Reference Comment Acrylfentanyl Analyte Name Units Reference Comment Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/-0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	Analyte Name	Units	Reference Comment
Averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown. N-ethyl Pentylone ng/mL N-ethyl Pentylone is a novel psychoactive stimulant. Valerylfentanyl ng/mL Valerylfentanyl is a novel non-prescription synthetic opioid. D52B Postmortem, Expanded, Blood (Forensic) Valerylfentanyl is a novel non-prescription synthetic opioid. D52B Postmortem, Expanded, Blood (Forensic) Valerylfentanyl is a novel non-prescription synthetic opioid. D52B Postmortem, Expanded, Blood (Forensic) Valerylfentanyl is a novel non-prescription synthetic opioid. D52B Postmortem, Expanded, Blood (Forensic) Valerylfentanyl were added. Units were changed. Acryl Fentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazeparm, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazeparm were removed. Scope of Analysis: Method (CPT Code) Analyte Name Units Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose and 0.03 +/- 0.	Acrylfentanyl	ng/mL	blood which may be dependent upon pH, collection tube, and storage temperature.
Valerylfentanyl ng/mL Valerylfentanyl is a novel non-prescription synthetic opioid. V52B Postmortem, Expanded, Blood (Forensic) Summary of Changes: Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletarmine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed. Scope of Analysis: Method (CPT Code) Reference Comment Analyte Name Units Reference Comment Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/-0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	mCPP	mcg/mL	averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone
Synthetic opioid. D52B Postmortem, Expanded, Blood (Forensic) Summary of Changes: Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed. Scope of Analysis: Method (CPT Code) Reference Comment Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Summary of Changes: Scope of Analysis was changed. Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed. Scope of Analysis: Method (CPT Code) Reference Comment Analyte Name Units Reference Comment Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	Valerylfentanyl	ng/mL	
Acrylfentanyl, N-ethyl Pentylone and Valerylfentanyl were added. Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed. Scope of Analysis: Method (CPT Code) Reference Comment Acrylfentanyl ng/mL Acrylfentanyl Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL mcCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	052B Postmortem, E	Expanded, Blood (Forens	sic)
Method (CPT Code) Image: Method (CPT Code) Analyte Name Units Reference Comment Acrylfentanyl ng/mL Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution. mCPP mcg/mL Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	Soone of Analysia	Units were changed. Acryl Fentanyl, BZP, Dicl Glyburide, Itraconazole, Mexiletine, Monoethylgly Tiletamine, Trihexypheni	lazepam, Etomidate, Glimepiride, Glipizide, Ketoconazole, Laudanosine, Metoclopramide, /cinexylidide (MEGX), N-Ethyl Pentylone, TFMPP,
Acrylfentanylng/mLAcrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.mCPPmcg/mLPeak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.			
blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.mCPPmcg/mLPeak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	Analyte Name	Units	Reference Comment
averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	Acrylfentanyl	ng/mL	blood which may be dependent upon pH, collection tube, and storage temperature.
N-ethyl Pentylone ng/mL N-ethyl Pentylone is a novel psychoactive stimulant.	mCPP	mcg/mL	averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone
	N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.



Analyte Name	Units	Reference Comment
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
052FL Postmortem, I	Expanded, Fluid (Forens	ic)
Summary of Changes:	BZP, Etomidate, Laudan	osine, Metoclopramide, Mexiletine, TFMPP, idyl and Zolazepam were removed.
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Butalbital Ethinamate Felbamate Fluconazole Lacosamide	mcg/mL mcg/mL mcg/mL mcg/mL mcg/mL	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Pentobarbital Phenobarbital Rufinamide Secobarbital Chlorpromazine Clonidine	mcg/mL mcg/mL mcg/mL mcg/mL ng/mL	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed. While overdoses from clonidine are common, deaths are



Analyte Name	Units	Reference Comment
		rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Xylazine EDDP Hydroxychloroquine mCPP Mescaline Normeperidine Phencyclidine Propoxyphene Quinidine Quinine 8052SP Postmortem, E	ng/mL ng/mL ng/mL mcg/mL mcg/mL mcg/mL mcg/mL mcg/mL ng/mL ng/mL	na (Forensic)
Summary of Changes:	Valerylfentanyl, Acrylfen Units were changed. Acryl Fentanyl, BZP, Dic Glyburide, Itraconazole, Mexiletine, Monoethylgh	changed. tanyl and N-ethyl Pentylone were added. clazepam, Etomidate, Glimepiride, Glipizide, Ketoconazole, Laudanosine, Metoclopramide, ycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, idyl, Valeryl Fentanyl, Voriconazole and Zolazepam
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
8052TI Postmortem, E	Expanded, Tissue (Forei	nsic)



Test Changes

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Summary of Changes:	Scope of Analysis was changed. Units were changed.
	BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed.

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment
Butalbital	mcg/g	
Ethinamate	mcg/g	
Felbamate	mcg/g	
Fluconazole	mcg/g	
Lacosamide	mcg/g	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Pentobarbital	mcg/g	
Phenobarbital	mcg/g	
Rufinamide	mcg/g	
Secobarbital	mcg/g	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
Clonidine	ng/g	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed.
		While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Xylazine EDDP	ng/g ng/g	
S Labs Welsh Rd.		
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Analyte Name	Units	Reference Comment
Hydroxychloroquine	ng/g	
mCPP	mcg/g	
Mescaline	mcg/g	
Normeperidine	mcg/g	
Phencyclidine	ng/g	
Propoxyphene	mcg/g	
Quinidine	ng/g	
Quinine	ng/g	
Trimipramine	ng/g	
052U Postmortem, E	Expanded, Urine (Forer	isic)
Summary of Changes:		changed. Pentylone, Norbuprenorphine and Valerylfentanyl were
	added.	
	Units were changed.	alazonom Olinizida Laudanagina Mataglanzamida
		clazepam, Glipizide, Laudanosine, Metoclopramide, lycinexylidide (MEGX), N-Ethyl Pentylone,
		al, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl
	Fentanyl and Zolazepa	
Coope of Arrely -'-		
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Acrylfentanyl	ng/mL	Acrylfentanyl is a novel non-prescription synthetic opioid.
Chlorpromazine	mcg/mL	
mCPP	mcg/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Norbuprenorphine	ng/mL	
Valerylfentanyl	ng/mL	Valeryl fentanyl is a novel non-prescription synthetic opioid.
Xylazine	ng/mL	
•	Expanded-II, Blood (For	rensic) (SSA)
Summary of Changes:	Glyburide, Itraconazole Mexiletine, Monoethylg Tiletamine, Trihexyphe	clazepam, Etomidate, Glimepiride, Glipizide, , Ketoconazole, Laudanosine, Metoclopramide, lycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, nidyl, Valeryl Fentanyl, Voriconazole and Zolazepam
	were remvoved.	



Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
9042B Postmortem, E	Expanded-II, with Vitreou	us Alcohol Confirmation, Blood (Forensic) (SSA)
Summary of Changes:	Glyburide, Itraconazole, Mexiletine, Monoethylgly	lazepam, Etomidate, Glimepiride, Glipizide, Ketoconazole, Laudanosine, Metoclopramide, ycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, idyl, Valeryl Fentanyl, Voriconazole and Zolazepam
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
0092B Postmortem, E	Expert w/Vitreous Alcoh	ol Confirmation, Blood (Forensic) (CSA)
Summary of Changes:	EMB-PICA, Acrylfentany Deschloroketamine, Eut Pentylone and Valerylfer Units were changed. 25B-NBOMe, 25C-NBOI Fluorophenmetrazine, 4- BZP, Clephedrone, Desc Glyburide, Itraconazole, Mephedrone, meta-Meth Methoxphenidine, Methy Monoethylglycinexylidide Fluorobutyrylfentanyl, pa Pyrazolam, TFMPP, THF	nine, 3-hydroxy-PCP, 5-fluoro-MDMB-PICA / 5-fluoro- /l, alpha-PHP / alpha-PiHP, Benzylone, ylone, Flualprazolam, N-butyl Pentylone, N-ethyl
	Valeryl Fentanyl, Voricoi	azolo ana zolazopam nolo fomotoal
Scope of Analysis: Method (CPT Code) Analyte Name	Units	Reference Comment



Test Changes

Analyte Name	Units	Reference Comment
3-hydroxy-PCP	ng/mL	
5-fluoro-MDMB-PICA / 5-fluoro- EMB-PICA	ng/mL	
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
alpha-PHP / alpha-PiHP	ng/mL	
Benzylone	ng/mL	
Deschloroketamine	ng/mL	Futulence is close ified as a synthetic stimulant and
Eutylone	ng/mL	Eutylone is classified as a synthetic stimulant and belongs to the beta-keto methylenedioxyamphetamine subclass, which includes synthetic stimulants methylone, butylone, ethylone, and N-ethylpentylone.
Flualprazolam	ng/mL	
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-butyl Pentylone	ng/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Valerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
0151B Postmortem, Expert	w/Vitreous Alcoh	ol Confirmation, Blood (Forensic) (CSA)

Summary of Changes: Scope of Analysis was changed

Summary of Changes:	Scope of Analysis was changed.
	3-hydroxy-PCP, 5-fluoro-MDMB-PICA / 5-fluoro-EMB-PICA, Acrylfentanyl, 2-
	fluoro Deschloroketamine, alpha-PHP / alpha-PiHP, Benzylone,
	Deschloroketamine, Eutylone, Flualprazolam, N-butyl Pentylone, N-ethyl
	Pentylone and Valerylfentanyl were added.
	Units were changed.
	25B-NBOMe, 25C-NBOMe, 25H-NBOMe, 25I-NBOMe, 3-
	Fluorophenmetrazine, 4-MeO-PCP, 5-fluoro-MDMB-PICA, Acryl Fentanyl,
	BZP, Clephedrone, Deschloroetizolam, Etomidate, Glimepiride, Glipizide,
	Glyburide, Itraconazole, Ketoconazole, Laudanosine, MDPV, Meclonazepam,
	Mephedrone, meta-Methylmethoxyacetylfentanyl, Methoxetamine,
	Methoxphenidine, Methylone, Metoclopramide, Mexiletine,
	Monoethylglycinexylidide (MEGX), MPHP, N-Ethyl Pentylone, para-
	Fluorobutyrylfentanyl, para-Methylmethoxyacetylfentanyl, Pentedrone,
	Pyrazolam, TFMPP, THF-F, Tiletamine, Trihexyphenidyl, U-49900, U-51754,
	Valeryl Fentanyl, Voriconazole and Zolazepam were removed.



Test Changes

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment
2-fluoro Deschloroketamine	ng/mL	
3-hydroxy-PCP	ng/mL	
5-fluoro-MDMB-PICA / 5-fluoro- EMB-PICA	ng/mL	
Acrylfentanyl	ng/mL	Acrylfentanyl is known to have limited stability in blood which may be dependent upon pH, collection tube, and storage temperature. Negative results should be interpreted with caution.
alpha-PHP / alpha-PiHP	ng/mL	
Benzylone	ng/mL	
Deschloroketamine	ng/mL	
Eutylone	ng/mL	Eutylone is classified as a synthetic stimulant and belongs to the beta-keto methylenedioxyamphetamine subclass, which includes synthetic stimulants methylone, butylone, ethylone, and N-ethylpentylone.
Flualprazolam nCPP	ng/mL mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
N-butyl Pentylone	ng/mL	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
/alerylfentanyl	ng/mL	Valerylfentanyl is a novel non-prescription synthetic opioid.
43B Postmortem, Experi	w/Vitreous Alcoh	ol Confirmation, Blood (Forensic)

Summary of Changes:	Scope of Analysis was changed.
	N-ethyl Pentylone was added.
	Units were changed.
	BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, N-Ethyl
	Pentylone, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were
	removed,

Scope of Analysis: Method (CPT Code)



Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL). The ratio of whole blood concentration to plasma concentration is 1.0.
Fluconazole	mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole.
		The blood to plasma ratio is not known for this analyte.
Lacosamide	mcg/mL	Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.
		Mean plasma concentrations following maintenance doses 200 mg/day: 4.99 +/- 2.51 mcg/mL; 400 mg/day: 9.35 +/- 4.22 mcg/mL; 600 mg/day: 12.46 +/- 5.60 mcg/mL.
		The ratio of whole blood concentration to plasma concentration is 1.1



Analyte Name	Units	Reference Comment
Metronidazole	mcg/mL	Peak Serum Concentrations (Single Oral Dose): 250 mg: 5.1 mcg/mL 1000 mg: 20 mcg/mL
		Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
		The blood to plasma ratio for metronidazole is unknown.
O-Desmethylvenlafaxine	ng/mL	Steady-state peak plasma levels following a daily regimen of Venlafaxine occur at approximately 2.5 hours for O-Desmethylvenlafaxine: 94-200 ng/mL (75 mg/day), 85-472 ng/mL (150 mg/day), 243-515 ng/mL (225 mg/day), 390-1096 ng/mL (450 mg/day). Steady-state trough plasma levels following a 150 mg per day regimen: 65-300 ng O-Desmethylvenlafaxine/mL.
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were produced 0.5-2.0 hours after a 100 mg oral dose and peak serum concentrations of 3 mcg/mL were produced 6 min. following a 100 mg IV dose. Potentially toxic at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with primidone: 10-40 mcg/mL. The blood to plasma ratio is approximately 0.8.
Clonidine	ng/mL	Immediate-release, oral: 0.50-2.0 ng/mL, 2 hours after administration; Sustained-release, patch: 0.20-2.0 ng/mL, at steady-state; Sustained-release, oral: 0.20-0.27 ng/mL, 6.8 +/- 3.6 hours after a 0.1 mg single dose in healthy fed adults; children receive higher doses on a mg/kg basis. The ratio of whole blood concentration to serum or plasma concentration is unknown for this analyte.



Analyte Name	Units	Reference Comment
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg (approximately 1600 mg) daily rufinamide resulted in plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133 patients maintained on twice-daily 400 or 800 mg doses for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.
		The blood to plasma ratio of rufinamide is approximately 1.0
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced a mean peak blood concentration of 2.0 mcg/mL (range, 1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3 mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours. Potentially toxic at blood concentrations greater than 8 mcg/mL.
Xylazine	ng/mL	
EDDP Hydroxychloroquine	ng/mL ng/mL	Peak plasma concentrations of 410 +/- 130 ng/mL were achieved 2.4 hours after a single oral dose of 400 mg hydroxychloroquine (n = 6). Two cases of hydroxychloroquine overdose (20 g each) were successfully treated throughout cardiovascular collapse and had serum concentrations of 14000 and 26000 ng/mL. The ratio of whole blood concentration to serum or plasma concentration is unknown for this analyte.
Levamisole	mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole. There is limited data available on therapeutic concentrations of levamisole and no data on levamisole concentrations encountered from tainted cocaine. The mean peak plasma concentration following a single 2.5 mg/kg dose was 0.48 +/- 0.22 mcg/mL. Following a single 50 mg dose the mean peak plasma concentration was 0.13 mcg/mL. The ratio of whole blood concentration to plasma concentration is unknown for this analyte.



Analyte Name	Units	Reference Comment
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
Mescaline N-Acetylprocainamide	mcg/mL mcg/mL	The normal therapeutic range for NAPA is 10 to 20 mcg/mL plasma. The blood to plasma ratio is not known for this analyte.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Phencyclidine Procyclidine	ng/mL mcg/mL	Steady-state concentrations following chronic oral 10 to 30 mg dose: 0.15-0.63 mcg/mL.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Pyrimethamine	mcg/mL	A single oral dose of 50 mg given to 5 subjects produced a peak plasma concentration of 0.21-0.43 mcg/mL in 2 to 4 hours following the dose.
Quinidine	ng/mL	For the treatment of arrhythmia, effective plasma concentrations typically range between 2000 and 5000 ng/mL. The blood/plasma ratio is not known for quinidine, but concentrations in red blood cells are usually lower than plasma.
Quinine	ng/mL	A single oral 648 mg antispasmodic dose produces average peak plasma concentrations of 2800 ng/mL 2 hr after administration. The blood/plasma ratio is not known for quinine, but concentrations in red blood cells are usually lower than plasma.
Tocainide	mcg/mL	
092B Postmortem, Ex	xpert, Blood (Forensic)	



Summary of Changes:	Scope of Analysis was changed. N-ethyl Pentylone was added. Units were changed. BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed	
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL). The ratio of whole blood concentration to plasma concentration is 1.0.
Fluconazole	mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole. The blood to plasma ratio is not known for this analyte.
Lacosamide	mcg/mL	Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.



Analyte Name	Units	Reference Comment
		200 mg/day: 4.99 +/- 2.51 mcg/mL;
		400 mg/day: 9.35 +/- 4.22 mcg/mL;
		600 mg/day: 12.46 +/- 5.60 mcg/mL.
		The ratio of whole blood concentration to plasma concentration is 1.1
Metharbital	mcg/mL	
Metronidazole	mcg/mL	Peak Serum Concentrations (Single Oral Dose):
		250 mg: 5.1 mcg/mL
		1000 mg: 20 mcg/mL
		Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
		The blood to plasma ratio for metronidazole is unknown.
O-Desmethylvenlafaxine	ng/mL	Steady-state peak plasma levels following a daily
		regimen of Venlafaxine occur at approximately
		2.5 hours for O-Desmethylvenlafaxine:
		94-200 ng/mL (75 mg/day),
		85-472 ng/mL (150 mg/day),
		243-515 ng/mL (225 mg/day),
		390-1096 ng/mL (450 mg/day).
		Steady-state trough plasma levels following a
		150 mg per day regimen:
		65-300 ng O-Desmethylvenlafaxine/mL.
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were
	Ū	produced 0.5-2.0 hours after a 100 mg oral dose and
		peak serum concentrations of 3 mcg/mL were produced
		6 min. following a 100 mg IV dose. Potentially toxic
		at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during
	-	anticonvulsant therapy with primidone: 10-40 mcg/mL.
		The blood to plasma ratio is approximately 0.8.



Analyte Name	Units	Reference Comment
Clonidine	ng/mL	Immediate-release, oral: 0.50-2.0 ng/mL, 2 hours after administration:
		Sustained-release, patch: 0.20-2.0 ng/mL, at
		steady-state;
		Sustained-release, oral: 0.20-0.27 ng/mL,
		6.8 +/- 3.6 hours after a 0.1 mg single dose in
		healthy fed adults;
		children receive higher doses on a mg/kg basis.
		The ratio of whole blood concentration to serum or
		plasma concentration is unknown for this analyte.
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg
		(approximately 1600 mg) daily rufinamide resulted in
		plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133
		patients maintained on twice-daily 400 or 800 mg doses
		for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.
		The blood to plasma ratio of rufinamide is
		approximately 1.0
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced
		a mean peak blood concentration of 2.0 mcg/mL (range,
		1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3
		mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours.
		Potentially toxic at blood concentrations greater than 8 mcg/mL.
Vulazina	n n/ml	
Xylazine EDDP	ng/mL ng/mL	
Hydroxychloroquine	ng/mL	Peak plasma concentrations of 410 +/- 130 ng/mL were
nyaroxyonioroquine	ng/me	achieved 2.4 hours after a single oral dose of 400 mg
		hydroxychloroquine (n = 6). Two cases of
		hydroxychloroquine overdose (20 g each) were
		successfully treated throughout cardiovascular collapse
		and had serum concentrations of
		14000 and 26000 ng/mL.
		The ratio of whole blood concentration to serum or
		plasma concentration is unknown for this analyte.



Analyte Name	Units	Reference Comment
Levamisole	mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole. There is limited data available on therapeutic concentrations of levamisole and no data on levamisole concentrations encountered from tainted cocaine. The mean peak plasma concentration following a single 2.5 mg/kg dose was 0.48 +/- 0.22 mcg/mL. Following a single 50 mg dose the mean peak plasma concentration was 0.13 mcg/mL. The ratio of whole blood concentration to plasma concentration is unknown for this analyte.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.
Mescaline N-Acetylprocainamide	mcg/mL mcg/mL	The normal therapeutic range for NAPA is 10 to 20 mcg/mL plasma. The blood to plasma ratio is not known for this analyte.
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.
Phencyclidine Procyclidine	ng/mL mcg/mL	Steady-state concentrations following chronic oral 10 to 30 mg dose: 0.15-0.63 mcg/mL.
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.
Pyrimethamine	mcg/mL	A single oral dose of 50 mg given to 5 subjects produced a peak plasma concentration of 0.21-0.43 mcg/mL in 2 to 4 hours following the dose.



Test Changes

Analyte Name	Units	Reference Comment
Quinidine	ng/mL	For the treatment of arrhythmia, effective plasma concentrations typically range between 2000 and 5000 ng/mL. The blood/plasma ratio is not known for quinidine, but concentrations in red blood cells are usually lower than plasma.
Quinine	ng/mL	A single oral 648 mg antispasmodic dose produces average peak plasma concentrations of 2800 ng/mL 2 hr after administration. The blood/plasma ratio is not known for quinine, but concentrations in red blood cells are usually lower than plasma.
Tocainide	mcg/mL	
	xpert, Fluid (Forensic)	
Summary of Changes:		nged. ne, Metoclopramide, Mexiletine, TFMPP, and Zolazepam were removed.
Scope of Analysis: Method (CPT Code)		
Analyte Name	Units	Reference Comment
Butalbital Ethinamate Felbamate Fluconazole Lacosamide	mcg/mL mcg/mL mcg/mL mcg/mL	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Metharbital Pentobarbital Chlorpromazine	mcg/mL mcg/mL mcg/mL	



Test Changes

Analyte Name	Units	Reference Comment
Clonidine	ng/mL	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed. While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Procyclidine Rufinamide Secobarbital Xylazine EDDP Hydroxychloroquine mCPP Mescaline Normeperidine Phencyclidine Propoxyphene Pyrimethamine Quinidine Quinine Tocainide	mcg/mL mcg/mL ng/mL ng/mL ng/mL mcg/mL mcg/mL mcg/mL mcg/mL mcg/mL ng/mL ng/mL ng/mL ng/mL ng/mL	
092SP Postmortem Summary of Change	Scope of Analysis was ch N-ethyl Pentylone was ac Units were changed. BZP, Etomidate, Laudand	(Light Protection) were changed. nanged.

removed.



Specimen Requirements:	10 mL Serum or Plasma
Transport Temperature:	Refrigerated
Specimen Container:	Gray top tube (Sodium Fluoride / Potassium Oxalate), Lavender top tube (EDTA), Plastic container (preservative-free)
Light Protection:	Not Required
Special Handling:	Serum: Collect sample in Red top tube Plasma: Collect sample in Gray top tube (Sodium Fluoride / Potassium Oxalate). Collect sample using alcohol free skin preparation. Promptly centrifuge and separate Serum or Plasma into a plastic screw capped vial using approved guidelines.
Rejection Criteria:	Polymer gel separation tube (SST or PST).
Scope of Analysis: Method (CPT Code)	

Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	A single oral 100 mg dose resulted in a mean peak blood concentration of 2.1 mcg/mL (range, 1.7-2.6 mcg/mL) at 2 hours, with a decline to 1.5 mcg/mL (range, 1.3-1.7 mcg/mL) by 24 hours. Potentially toxic at plasma concentrations greater than 10 mcg/mL.
Ethinamate	mcg/mL	Usual hypnotic range: 5-10 mcg/mL
Felbamate	mcg/mL	Fifty-six adult patients receiving an average daily oral dose of 2300 mg had steady-state trough plasma concentrations averaging 33 mcg/mL (range, 18-52 mcg/mL). Twenty-six patients ages 10-69 years receiving an average daily dose of 2685 mg had serum concentrations averaging 69 mcg/mL (range, 16-165 mcg/mL).
Fluconazole	mcg/mL	Single oral doses of 50 or 150 mg fluconazole resulted in peak plasma concentrations of 0.93 +/- 0.13 mcg/mL and 2.7 +/- 0.4 mcg/mL respectively. Peak plasma concentrations were 6.7 mcg/mL (range 4.1-8.1 mcg/mL) approximately 1 to 2 hours after a single 400 mg oral dose of fluconazole.
Lacosamide	mcg/mL	Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.
		Mean plasma concentrations following maintenance doses: 200 mg/day: 4.99 +/- 2.51 mcg/mL;
S Labs Welsh Rd.		



Analyte Name	Units	Reference Comment
		400 mg/day: 9.35 +/- 4.22 mcg/mL; 600 mg/day: 12.46 +/- 5.60 mcg/mL.
		NMS Labs derived data: 5th - 95th Percentile Data: 1.8-13.0 mcg/mL Mean: 5.3 mcg/mL (N = 14900)
Metharbital Metronidazole	mcg/mL mcg/mL	Peak Serum Concentrations (Single Oral Dose): 250 mg: 5.1 mcg/mL 1000 mg: 20 mcg/mL
		Plasma Steady-State (500 mg, IV, every 8 h): 22 mcg/mL
O-Desmethylvenlafaxine	ng/mL	Steady-state peak plasma levels following a daily regimen of Venlafaxine occur at approximately 2.5 hours for O-Desmethylvenlafaxine: 94-200 ng/mL (75 mg/day), 85-472 ng/mL (150 mg/day), 243-515 ng/mL (225 mg/day), 390-1096 ng/mL (450 mg/day).
		Steady-state trough plasma levels following a 150 mg per day regimen: 65-300 ng O-Desmethylvenlafaxine/mL.
Pentobarbital	mcg/mL	Peak serum concentrations of 1.2-3.1 mcg/mL were produced 0.5-2.0 hours after a 100 mg oral dose and peak serum concentrations of 3 mcg/mL were produced 6 min. following a 100 mg IV dose. Potentially toxic at blood concentrations greater than 10 mcg/mL.
Phenobarbital	mcg/mL	Recommended serum concentration range during anticonvulsant therapy with primidone: 10-40 mcg/mL.
Clonidine	ng/mL	Immediate-release, oral: 0.50-2.0 ng/mL, 2 hours after administration; Sustained-release, patch: 0.20-2.0 ng/mL, at steady-state; Sustained-release, oral: 0.20-0.27 ng/mL, 6.8 +/- 3.6 hours after a 0.1 mg single dose in healthy fed adults; children receive higher doses on a mg/kg basis.



Analyte Name	Units	Reference Comment
Rufinamide	mcg/mL	Maintenance therapy with 45 mg/kg (approximately 1600 mg) daily rufinamide resulted in plasma concentrations ranging from 5.0-48 mcg/mL (n = 74).
		Trough plasma concentrations in groups of 129-133 patients maintained on twice-daily 400 or 800 mg doses for 3 months averaged 2.6 or 4.7 mcg/mL, respectively.
Secobarbital	mcg/mL	A 3.3 mg/kg oral dose (approx. 230 mg/70 kg) produced a mean peak blood concentration of 2.0 mcg/mL (range, 1.8-2.2 mcg/mL) at 3 hours, diminishing to 1.3 mcg/mL by 20 hours and 0.8 mcg/mL by 40 hours. Potentially toxic at blood concentrations greater than 8 mcg/mL.
Xylazine EDDP Hydroxychloroquine	ng/mL ng/mL ng/mL	Peak plasma concentrations of 410 +/- 130 ng/mL were achieved 2.4 hours after a single oral dose of 400 mg hydroxychloroquine (n = 6). Two cases of hydroxychloroquine overdose (20 g each) were successfully treated throughout cardiovascular collapse and had serum concentrations of 14000 and 26000 ng/mL.
Levamisole	mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole. There is limited data available on therapeutic concentrations of levamisole and no data on levamisole concentrations encountered from tainted cocaine. The mean peak plasma concentration following a single 2.5 mg/kg dose was 0.48 +/- 0.22 mcg/mL. Following a single 50 mg dose the mean peak plasma concentration was 0.13 mcg/mL.
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days.
Mescaline N-Acetylprocainamide	mcg/mL mcg/mL	The normal therapeutic range for NAPA is 10 to 20 mcg/mL plasma.



Test Changes

Analyte Name	Units	Reference Comment	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.	
Normeperidine	mcg/mL	Expected analgesic range: 0.1-0.6 mcg Meperidine/mL. Normeperidine concentrations: Up to 0.5 mcg/mL.	
Phencyclidine Procyclidine	ng/mL mcg/mL	Steady-state concentrations following chronic oral 10 to 30 mg dose: 0.15-0.63 mcg/mL.	
Propoxyphene	mcg/mL	Average serum concentrations following a daily regimen of 195 mg Propoxyphene: 0.42 mcg Propoxyphene/mL.	
Pyrimethamine	mcg/mL	A single oral dose of 50 mg given to 5 subjects produced a peak plasma concentration of 0.21-0.43 mcg/mL in 2 to 4 hours following the dose.	
Quinidine	ng/mL	For the treatment of arrhythmia, effective plasma concentrations typically range between 2000 and 5000 ng/mL.	
Quinine	ng/mL	A single oral 648 mg antispasmodic dose produces average peak plasma concentrations of 2800 ng/mL 2 hr after administration.	
Tocainide	mcg/mL	Reported antiarrhythmic concentration: 4-10 mcg/mL. Tocainide is an antiarrhythmic drug that is no longer available in the Unites States.	

8092TI Postmortem, Expert, Tissue (Forensic)

Summary of Changes: Scope of Analysis was changed. Units were changed. BZP, Etomidate, Laudanosine, Metoclopramide, Mexiletine, TFMPP, Tiletamine, Trihexyphenidyl and Zolazepam were removed.

Scope of Analysis: Method (CPT Code)

Analyte Name	Units	Reference Comment	
Butalbital	mcg/g		
Ethinamate	mcg/g		
Felbamate	mcg/g		
Fluconazole	mcg/g		



Analyte Name	Units	Reference Comment
Lacosamide	mcg/g	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day. Patients exposed to supratherapeutic doses during clinical trials had adverse events that were not clinically different than those reported in patients receiving recommended doses of lacosamide.
Metharbital Pentobarbital Phenobarbital Clonidine	mcg/g mcg/g mcg/g ng/g	Clonidine is a potent antihypertensive agent that has both alpha-adrenergic agonist and antagonist activities. It is normally supplied in tablets containing 0.1, 0.2, or 0.3 mg each. The recommended daily oral dose ranges from a minimum of 0.2 mg to a maximum of 2.4 mg per day, although the dose at the high end of the range is rarely employed.
		While overdoses from clonidine are common, deaths are rare. Clinical sequelae of clonidine toxicity may include hypertension or hypotension, bradycardia, lethargy, weakness, somnolence, and hypoventilation. Severe overdoses may result in cardiac conduction defects, arrhythmias, apnea, convulsions, coma and death.
Procyclidine Rufinamide Secobarbital	mcg/g mcg/g mcg/g	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
Xylazine EDDP Hydroxychloroquine mCPP Mescaline Normeperidine Phencyclidine Propoxyphene Pyrimethamine Quinidine	ng/g ng/g mcg/g mcg/g mcg/g ng/g mcg/g mcg/g mcg/g ng/g	



Analyte Name	Units	Reference Comment
Tocainide	mcg/g	
Trimipramine	ng/g	
8092U Postmortem	, Expert, Urine (Forensic)	
Summary of Change	N-ethyl Pentylone was add Units were changed. BZP, Etomidate, Laudanos	
Scope of Analysi Method (CPT Code		
Analyte Name	Units	Reference Comment
Butalbital	mcg/mL	The disposition of butalbital has not been well studied in humans.
Fluconazole	mcg/mL	
Lacosamide	mcg/mL	Lacosamide is a functionalized amino acid specifically synthesized as an anticonvulsant drug. In addition to being approved for use as an adjunctive therapy treatment of partial-onset seizures it has been investigated as a treatment for diabetic neuropathic pain. Lacosamide can be administered either orally or intravenously. The recommended initial dose is 50 mg/twice a day, up to a maintenance dose of 200 to 400 mg/day.
		Single labeled oral or intravenous lacosamide doses in healthy subjects were eliminated in urine (95%) and feces (< 0.5%) over a 7 day interval. Urinary excretion products included parent drug (40% of the dose) and the pharmacologically inactive O-desmethyllacosamide.
Metharbital	mcg/mL	
N-Acetylprocainamide	mcg/mL	N-acetylprocainamide is an antiarrhythmic drug and an active metabolite of procainamide. The reported qualitative result for this substance was based upon a single analysis only. If confirmation testing is required please contact the laboratory.
Chlorpromazine Pentobarbital	mcg/mL mcg/mL	Less than 1% of a dose is eliminated in the urine as unchanged drug.



Test Changes

Units	Reference Comment
mcg/mL	For patients on chronic therapy, a mean of 20% (range, 12 - 55%) of the dose is excreted unchanged in the 24 hour urine.
ng/mL	
mcg/mL	
mcg/mL	Procyclidine is an anticholinergic drug that was previously used in the treatment of Parkinson's disease.
mcg/mL	Secobarbital undergoes extensive biotransformation. However, approximately 5% of secobarbital is excreted unchanged in the urine within 2 days.
na/mL	
ng/mL	In maintenance subjects: Up to 50000 ng of Methadone plus Methadone Metabolites/mL Urine.
ng/mL	
mcg/mL	Levamisole is used as a veterinary antihelminthic (worming agent) in animals. It is no longer available in North America for human use. However, from July- September 2008 approximately 30% of cocaine seized by the DEA was contaminated with levamisole.
mca/ml	
ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
mca/ml	
mcg/mL	
mcg/mL	
	Quinine is derived from the bark of the cinchona tree.
ng/me	It has been used in the past as an antimalarial, but is
	more commonly used today to treat muscle cramps.
	It is also used as a flavoring agent in tonic water and
	as a cutting agent in illicit heroin. Adverse effects
	include gastrointestinal disturbances, tinnitus, dizziness, arrhythmias and hypotension.
mag/ml	
mcg/mL	Tocainide is an antiarrhythmic drug that is no longer
	mcg/mL ng/mL mcg/mL mcg/mL ng/mL ng/mL ng/mL mcg/mL mcg/mL ng/mL ng/mL ng/mL ng/mL ng/mL ng/mL ng/mL

4177B Postmortem, SUIDS Screen, Blood (Forensic)

Effective Date: Monday, August 15, 2022



Test Updates

Test Changes

	Units were changed. Acryl Fentanyl, BZP, Diclazepam, Etomidate, Glimepiride, Glipizide, Glyburide, Itraconazole, Ketoconazole, Laudanosine, Metoclopramide, Mexiletine, Monoethylglycinexylidide (MEGX), N-Ethyl Pentylone, TFMPP, Tiletamine, Trihexyphenidyl, Valeryl Fentanyl, Voriconazole and Zolazepam were removed.		
Scope of Analysis: Method (CPT Code)			
Analyte Name	Units	Reference Comment	
mCPP	mcg/mL	Peak steady-state concentrations of mCPP in plasma averaged 0.03 mcg/mL at approximately 8 hours post dose following 300 mg normal release trazodone for 7 days and 0.03 +/- 0.01 mcg/mL following 200 mg nefazodone for 8 days. The blood to plasma ratio is unknown.	
52494B Substituted C	athinone Confirmation, Blood		
Summary of Changes:	Scope of Analysis was changed. N-ethyl Pentylone was added. N-Ethyl Pentylone was removed.		
Scope of Analysis: Method (CPT Code)		hyl Pentylone, Dibutylone	
Analyta Nama	· · ·		
Analyte Name	Units	Reference Comment	
N-ethyl Pentylone	Units ng/mL	Reference Comment N-ethyl Pentylone is a novel psychoactive stimulant.	
N-ethyl Pentylone		N-ethyl Pentylone is a novel psychoactive stimulant.	
N-ethyl Pentylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma	
N-ethyl Pentylone 52494SP Substituted C	ng/mL athinone Confirmation, Serum/ Scope of Analysis was changed N-ethyl Pentylone was added. N-Ethyl Pentylone was removed GC/MS (80371): Butylone, N-et	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma I.	
N-ethyl Pentylone 52494SP Substituted C Summary of Changes: Scope of Analysis:	ng/mL athinone Confirmation, Serum/ Scope of Analysis was changed N-ethyl Pentylone was added. N-Ethyl Pentylone was removed GC/MS (80371): Butylone, N-et	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma I.	
N-ethyl Pentylone 52494SP Substituted C Summary of Changes: Scope of Analysis: Method (CPT Code)	ng/mL sathinone Confirmation, Serum/ Scope of Analysis was changed N-ethyl Pentylone was added. N-Ethyl Pentylone was removed GC/MS (80371): Butylone, N-et	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma d. hyl Pentylone, Dibutylone	
N-ethyl Pentylone 52494SP Substituted C Summary of Changes: Scope of Analysis: Method (CPT Code) Analyte Name N-ethyl Pentylone	ng/mL Eathinone Confirmation, Serum/ Scope of Analysis was changed N-ethyl Pentylone was added. N-Ethyl Pentylone was removed GC/MS (80371): Butylone, N-ethyl Units	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma I. d. hyl Pentylone, Dibutylone Reference Comment	
N-ethyl Pentylone 52494SP Substituted C Summary of Changes: Scope of Analysis: Method (CPT Code) Analyte Name N-ethyl Pentylone	ng/mL athinone Confirmation, Serum/ Scope of Analysis was changed N-ethyl Pentylone was added. N-Ethyl Pentylone was removed GC/MS (80371): Butylone, N-ether Units ng/mL athinone Confirmation, Urine	N-ethyl Pentylone is a novel psychoactive stimulant. Plasma I. J. Hyl Pentylone, Dibutylone Reference Comment N-ethyl Pentylone is a novel psychoactive stimulant. I.	



Analyte N	Name	Units	Reference Comment	
N-ethyl Pe	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.	
52328B	Substituted Ca	athinone Panel Confirmation, Blood		
Sumn	nary of Changes:	Scope of Analysis was changed. N-ethyl Pentylone was added. N-Ethyl Pentylone was removed.		
	cope of Analysis: thod (CPT Code)	GC/MS (80371): Ethylone, But	tylone, Pentylone, N-ethyl Pentylone, Dibutylone	
Analyte N	Name	Units	Reference Comment	
N-ethyl Pe	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.	
52328SP	Substituted Ca	athinone Panel Confirmation,	Serum/Plasma	
Sumn	nary of Changes:	Scope of Analysis was change N-ethyl Pentylone was added. N-Ethyl Pentylone was remove		
	cope of Analysis: thod (CPT Code)	GC/MS (80371): Ethylone, But	tylone, Pentylone, N-ethyl Pentylone, Dibutylone	
Analyte N	Name	Units	Reference Comment	
N-ethyl Pe	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.	
52328U	Substituted Co	athinone Panel Confirmation,	Urine	
525200	Substituted Ca			
	nary of Changes:	Scope of Analysis was change N-ethyl Pentylone was added. N-Ethyl Pentylone was remove		
Sumn		N-ethyl Pentylone was added. N-Ethyl Pentylone was remove		
Sumn	nary of Changes: cope of Analysis: thod (CPT Code)	N-ethyl Pentylone was added. N-Ethyl Pentylone was remove	ed.	
Sumn S Me	nary of Changes: cope of Analysis: thod (CPT Code) Name	N-ethyl Pentylone was added. N-Ethyl Pentylone was remove GC/MS (80371): Ethylone, But	ed. tylone, Pentylone, N-ethyl Pentylone, Dibutylone	
Sumn S Me Analyte N N-ethyl Pe	nary of Changes: cope of Analysis: thod (CPT Code) Name entylone	N-ethyl Pentylone was added. N-Ethyl Pentylone was remove GC/MS (80371): Ethylone, But Units	ed. tylone, Pentylone, N-ethyl Pentylone, Dibutylone Reference Comment	
Sumn S Mer Analyte N N-ethyl Pe 1021B	nary of Changes: cope of Analysis: thod (CPT Code) Name entylone	N-ethyl Pentylone was added. N-Ethyl Pentylone was remove GC/MS (80371): Ethylone, But Units ng/mL athinone Panel, Blood	ed. tylone, Pentylone, N-ethyl Pentylone, Dibutylone Reference Comment N-ethyl Pentylone is a novel psychoactive stimulant.	



Analyte	Name	Units	Reference Comment
N-ethyl P	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
1021SP	Substituted Ca	athinone Panel, Serum	ı/Plasma
Sumr	mary of Changes:	Scope of Analysis was N-ethyl Pentylone was N-Ethyl Pentylone was	s added.
	Scope of Analysis: ethod (CPT Code)		
Analyte	Name	Units	Reference Comment
N-ethyl P	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
1021U	Substituted Ca	athinone Panel, Urine	
Sumr	mary of Changes:	Scope of Analysis was N-ethyl Pentylone was N-Ethyl Pentylone was	s added.
	Cope of Analysis: hod (CPT Code)	GC/MS (80371): Penty	ylone, Ethylone, Butylone, Dibutylone, N-ethyl Pentylone
Analyte	Name	Units	Reference Comment
N-ethyl P	entylone	ng/mL	N-ethyl Pentylone is a novel psychoactive stimulant.
5970B	Synthetic Can	nabinoids Confirmatio	on (Qualitative), Blood
Sumr	mary of Changes:	Scope of Analysis was 5-fluoro-MDMB-PICA 5-fluoro-MDMB-PICA	/ 5-fluoro-EMB-PICA was added.
	Cope of Analysis: hthod (CPT Code)		
Analyte	Name	Units	Reference Comment
5-fluoro-N EMB-PIC	MDMB-PICA / 5-flu A	oro- ng/mL	
9566B	Synthetic Can	nabinoids Screen (Ade	d-On), Blood
Sumr	mary of Changes:	Scope of Analysis was 5-fluoro-MDMB-PICA / 5-fluoro-MDMB-PICA \	/ 5-fluoro-EMB-PICA was added.



	LC-MS/MS (80307): ADMB-FUBINACA, 5-fluoro-PICA 3,3-dimethylbutanoic acid, 5- fluoro-PINACA 3-methylbutanoic acid, 4-fluoro-BINACA 3,3-dimethylbutanoic acid, FUBINACA 3-methylbutanoic acid, 5-fluoro-PINACA 3,3-dimethylbutanoic acid, FUBINACA 3,3-dimethylbutanoic acid, APP-BINACA, 5-fluoro-MDMB-PICA / 5- fluoro-EMB-PICA, MMB-FUBINACA, 5-fluoro-MDMB-PINACA / 5-fluoro-EMB- PINACA, MDMB-4en-PINACA, ADMB-CHMINACA, 4-fluoro-MDMB-BINACA			
Analyte Name	Units	Reference Comment		
5-fluoro-MDMB-PICA / 5-flu EMB-PICA	ioro- ng/mL			
9560B Synthetic Can	nabinoids Screen, Blo	bod		
Summary of Changes:		/ 5-fluoro-EMB-PICA was added.		
	LC-MS/MS (80307): ADMB-FUBINACA, 5-fluoro-PICA 3,3-dimethylbutanoic acid, 5- fluoro-PINACA 3-methylbutanoic acid, 4-fluoro-BINACA 3,3-dimethylbutanoic acid, FUBINACA 3-methylbutanoic acid, 5-fluoro-PINACA 3,3-dimethylbutanoic acid, FUBINACA 3,3-dimethylbutanoic acid, APP-BINACA, 5-fluoro-MDMB-PICA / 5- fluoro-EMB-PICA, MMB-FUBINACA, 5-fluoro-MDMB-PINACA / 5-fluoro-EMB- PINACA, MDMB-4en-PINACA, ADMB-CHMINACA, 4-fluoro-MDMB-BINACA			
Analyte Name	Units	Reference Comment		
5-fluoro-MDMB-PICA / 5-flu EMB-PICA	ioro- ng/mL			



Discontinued Tests

Test	Test Name	Alternative Test
52366B	Bath Salts Confirmation, Blood	No Alternate Tests Available
52366SP	Bath Salts Confirmation, Serum/Plasma	No Alternate Tests Available
52366U	Bath Salts Confirmation, Urine	No Alternate Tests Available
52503U	Designer Benzodiazepines Confirmation 2 (Qualitative), Urine	No Alternate Tests Available
52503B	Designer Benzodiazepines Confirmation 2, Blood	No Alternate Tests Available
52503SP	Designer Benzodiazepines Confirmation 2, Serum/Plasma	No Alternate Tests Available
52320B	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Blood	No Alternate Tests Available
52320SP	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Serum/Plasma	No Alternate Tests Available
52320U	Hallucinogens and Stimulants Confirmation 2 (Qualitative), Urine	No Alternate Tests Available
52081B	Metoclopramide Confirmation, Blood	No Alternate Tests Available
52081FL	Metoclopramide Confirmation, Fluid	No Alternate Tests Available
52081SP	Metoclopramide Confirmation, Serum/Plasma	No Alternate Tests Available
52081TI	Metoclopramide Confirmation, Tissue	No Alternate Tests Available
52081U	Metoclopramide Confirmation, Urine	No Alternate Tests Available
54342U	Mitragynine, Phenazepam Confirmation (Qualitative) (DUID/DRE), Urine	No Alternate Tests Available
52326B	Piperazine Designer Drugs Confirmation, Blood	No Alternate Tests Available
52326SP	Piperazine Designer Drugs Confirmation, Serum/Plasma	No Alternate Tests Available
52326U	Piperazine Designer Drugs Confirmation, Urine	No Alternate Tests Available
52327B	Pyrrolidinophenone Confirmation, Blood	No Alternate Tests Available
52327SP	Pyrrolidinophenone Confirmation, Serum/Plasma	No Alternate Tests Available