

**TEST CHANGE** 

## Drug Profile, Screen With Reflex to Quantitation, Serum or Plasma 0092420, DRUG SCRSP

Specimen Requirements:	
Patient Preparation:	
Collect:	Gray (sodium fluoride/potassium oxalate). Also acceptable: Plain red, green (sodium heparin), lavender (EDTA), or pink (K2EDTA).
Specimen Preparation:	Remove plasma from cells ASAP or within 2 hours of collection. Transfer 4 mL plasma to an ARUP standard transport tube. (Min: 3 mL) Also acceptable: Serum.
Transport Temperature:	Refrigerated.
Unacceptable Conditions:	Specimens exposed to repeated freeze/thaw cycles. Separator tubes. Plasma or whole blood collected in It. blue (sodium citrate). Hemolyzed specimens.
Remarks:	Cocaine and cocaethylene are more stable in fluoride-preserved plasma than serum.
Stability:	After separation from cells: Ambient: 1 week; Refrigerated: 2 weeks; Frozen: 3 years
Methodology:	Qualitative Enzyme-Linked Immunosorbent Assay (ELISA) / Quantitative Gas Chromatography-Mass Spectrometry (GC-MS) / Quantitative Liquid Chromatography-Tandem Mass Spectrometry
Note:	Screen-positive specimens are automatically confirmed by GC/MS and/or LC-MS/MS; additional charges may apply.
CPT Codes:	80307; if reflexed, add 80324; <u>80354;</u> 80345; 80346; 80348; 80349; 80353; 80358; 80359; 80361; 80365; 83992 (Reflexed Alt Code: G0480)
New York DOH Approval Status:	This test is New York DOH approved.

Effective Date: January 20, 2026

Interpretive Data:

Drugs/drug classes reported as "Positive" are automatically reflexed to mass spectrometry confirmation/quantitation. An unconfirmed positive immunoassay screen result may be useful for medical purposes but does not meet forensic standards. The absence of expected drug(s) and/or drug metabolite(s) may indicate noncompliance, inappropriate timing of specimen collection relative to drug administration, poor drug absorption, or limitations of testing. The concentration at which the screening test can detect a drug or metabolite varies within a drug class. Specimens for which drugs or drug classes are detected by the screen are automatically reflexed to a second, more specific technology (GC/MS and/or LC-MS/MS). The concentration value must be greater than or equal to the cutoff to be reported as positive. Interpretive questions should be directed to the laboratory.

For medical purposes only; not valid for forensic use.

Reference Interval:



Effective Date: January 20, 2026

<u>Test</u> <u>Number</u>	Components	Reference Interval
	Amphetamines Screen, S/P	Positive Cutoff: 20 ng/mL
	Barbiturates Screen, S/P	Positive Cutoff: 50 ng/mL
	Benzodiazepines Screen, S/P	Positive Cutoff: 50 ng/mL
	Buprenorphine Screen, S/P	Positive Cutoff: 1 ng/mL
	Carboxy-THC Screen, S/P	Positive Cutoff: 20 ng/mL
	Cocaine Screen, S/P	Positive Cutoff: 20 ng/mL
	Fentanyl Screen, S/P	Positive Cutoff: 1 ng/mL
	Methadone Screen, S/P	Positive Cutoff: 25 ng/mL
	Methamphetamine Screen, S/P	Positive Cutoff: 20 ng/mL
	Opiates Screen, S/P	Positive Cutoff: 20 ng/mL
	Oxycodone/Oxymorphone Screen, S/P	Positive Cutoff: 20 ng/mL
	Phencyclidine Screen, S/P	Positive Cutoff: 10 ng/mL

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Drugs Covered a	and Cutoff Con	<del>centrations</del>	
<del>Drugs/Drug</del> <del>Classes</del>	Screen		
Amphetamines	Effective August 17, 2020-20 ng/mL		
Methamphetamine	Effective August 17, 2020-20 ng/mL		
Barbiturates	Effective August 17, 2020-50 ng/mL		
Benzodiazepines	Effective August 17, 2020 50 ng/mL		
Buprenorphine	1 ng/mL		
<del>Cannabinoids</del>	Effective August 17, 2020-20 ng/mL		
<del>Cocaine</del>	Effective August 17, 2020-20 ng/mL		
<del>Methadone</del>	Effective August 17, 2020-25 ng/mL		
<del>Opiates</del>	Effective August 17, 2020 20 ng/mL		
<del>Oxycodone</del>	Effective August 17, 2020 20 ng/mL		
Phencyclidine Phencyclidine	Effective August 17, 2020-10 ng/mL		

HOTLINE NOTE: There is a component change associated with this test. One or more components



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have been added or removed. Refer to the Hotline Test Mix for interface build information.