Multi-Drug

One Step

Screen Test Panel (Urine) Package Insert

English

Package insert for testing of any combination of the following drugs:

Amphetamine, Amphetamine 500, Amphetamine 300, Barbiturates, Benzodiazepines, Benzodiazepines 200, Buprenorphine, Cocaine, Cocaine 150, Marijuana, Methadone, Methamphetamine, Methamphetamine 500, Methamphetamine 300, Methylenedioxymethamphetamine, Morphine 300, Opiate 2000, Oxycodone, Phencyclidine, Propoxyphene and Tricyclic Antidepressants.

A rapid, one step screen test for the simultaneous, qualitative detection of multiple drugs and metabolites in human urine.

For medical and other professional in vitro diagnostic use only.

INTENDED USE & SUMMARY

Urine based screen tests for multiple drugs of abuse range from simple immunoassay tests to complex analytical procedures. The speed and sensitivity of immunoassays have made them the most widely accepted method to screen urine for multiple drugs of abuse.

The Multi-Drug One Step Screen Test Panel (Urine) is a lateral flow chromatographic immunoassay for the qualitative detection of following drugs without the need of instruments.¹

Test	Calibrator	Cut-off (ng/mL)
Amphetamine (AMP)	d-Amphetamine	1,000
Amphetamine (AMP 500)	d-Amphetamine	500
Amphetamine (AMP 300)	d-Amphetamine	300
Barbiturates (BAR)	Secobarbital	300
Benzodiazepines (BZO)	Oxazepam	300
Benzodiazepines (BZO 200)	Oxazepam	200
Buprenorphine (BUP)	Buprenorphine	10
Cocaine (COC)	Benzoylecgonine	300
Cocaine (COC 150)	Benzoylecgonine	150
Marijuana (THC)	11-nor-Δ ⁹ -THC-9 COOH	50
Methadone (MTD)	Methadone	300
Methamphetamine (MET)	d-Methamphetamine	1,000
Methamphetamine (MET 500)	d-Methamphetamine	500
Methamphetamine (MET 300)	d-Methamphetamine	300
Methylenedioxymethamphetamine (MDMA)	d,l-Methylenedioxymethamphetamine	500
Morphine (MOP 300)	Morphine	300
Opiate (OPI 2000)	Morphine	2,000
Oxycodone (OXY)	Oxycodone	100
Phencyclidine (PCP)	Phencyclidine	25
Propoxyphene (PPX)	Propoxyphene	300
Tricyclic Antidepressants (TCA)	Nortriptyline	1,000

This test will detect other related compounds, please refer to the Analytical Specificity table in this package insert.

This assay provides only a preliminary analytical test result. A more specific alternate chemical method must be used in order to obtain a confirmed analytical result. Gas chromatography/mass spectrometry (GC/MS) is the preferred confirmatory method. Clinical consideration and professional judgment should be applied to any drug of abuse test result, particularly when preliminary positive results are used.

PRINCIPLE

The Multi-Drug One Step Screen Test Panel (Urine) is an immunoassay based on the principle of competitive binding. Drugs which may be present in the urine specimen compete against their respective drug conjugate for binding sites on their specific antibody.

During testing, a urine specimen migrates upward by capillary action. A drug, if present in the urine specimen below its cut-off concentration, will not saturate the binding sites of its specific antibody coated on the particles. The antibody coated particles will then be captured by the immobilized drug conjugate and a visible colored line will show up in the test line region of the specific drug strip. The colored line will not form in the test line region if the drug level is above its cut-off concentration because it will saturate all the binding sites of the antibody coated on the particles.

A drug-positive urine specimen will not generate a colored line in the specific test line region of the strip because of drug competition, while a drug-negative urine specimen or a specimen containing a drug concentration less than the cut-off will generate a line in the test line region. To serve as a procedural control, a colored line will always appear at the control line region indicating that proper volume of specimen has been added and membrane wicking has occurred.

REAGENTS

Each test in the test panel contains mouse monoclonal antibody-coupled particles and corresponding drugprotein conjugates. A goat antibody is employed in each control line.

PRECAUTIONS

- For medical and other professional in vitro diagnostic use only. Do not use after the expiration date.
- · The test panel should remain in the sealed pouch until use.
- · All specimens should be considered potentially hazardous and handled in the same manner as an infectious agent
- The used test panel should be discarded according to local regulations.

STORAGE AND STABILITY

Store as packaged in the sealed pouch either at room temperature or refrigerated (2-30°C). The test panel is stable through the expiration date printed on the sealed pouch. The test panel must remain in the sealed pouch until use. DO NOT FREEZE. Do not use beyond the expiration date.

SPECIMEN COLLECTION AND PREPARATION

Urine Assay

The urine specimen must be collected in a clean and dry container. Urine collected at any time of the day may be used. Urine specimens exhibiting visible precipitates should be centrifuged, filtered, or allowed to settle to obtain a clear supernatant for testing.

Specimen Storage

Urine specimens may be stored at 2-8°C for up to 48 hours prior to assay. For prolonged storage, specimens may be frozen and stored below -20°C. Frozen specimens should be thawed and mixed before testing.

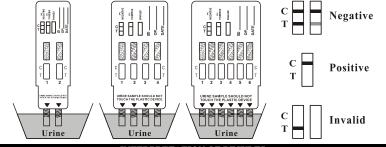
MATERIALS Materials Provided

· Package insert

DIRECTIONS FOR USE

Allow the test panel, urine specimen, and/or controls to equilibrate to room temperature (15-30°C) prior to testing.

- 1. Bring the pouch to room temperature before opening it. Remove the test panel from the sealed pouch and use it as soon as possible.
- 2. Take off the cap outside of the test end. With arrows pointing toward the urine specimen, immerse the test panel vertically into the urine specimen for at least 10-15 seconds. Immerse the test panel to at least the level of the wavy lines on the strip(s), do not pass the arrows on the test panel when immersing the panel. See the illustration below.
- 3. Place the test panel on a non-absorbent flat surface, start the timer and wait for the colored line(s) to appear. Read results at 5 minutes. Do not interpret results after 10 minutes.



INTERPRETATION OF RESULTS

(Please refer to the illustration above)

NEGATIVE:* A colored line in the control line region (C) and a colored line in the test line region (T) for a specific drug indicate a negative result. This indicates that the drug concentration in the urine specimen is below the designated cut-off level for that specific drug.

*NOTE: The shade of color in the test line region (T) may vary, but it should be considered negative whenever there is even a faint colored line.

POSITIVE: A colored line in the control line region (C) but no line in the test line region (T) for a specific drug indicates a positive result. This indicates that the drug concentration in the urine specimen exceeds the designated cut-off for that specific drug.

INVALID: Control line fails to appear. Insufficient specimen volume or incorrect procedural techniques are the most likely reasons for control line failure. Review the procedure and repeat the test using a new test panel. If the problem persists, discontinue using the lot immediately and contact your local distributor.

OUALITY CONTROL

A procedural control is included in the test. A colored line appearing in the control line region (C) is considered an internal procedural control. It confirms sufficient specimen volume, adequate membrane wicking and correct procedural technique.

Control standards are not supplied with this kit; however, it is recommended that positive and negative controls be tested as good laboratory practice to confirm the test procedure and to verify proper test performance.

LIMITATIONS

- 1. The Multi-Drug One Step Screen Test Panel (Urine) provides only a preliminary analytical result. A more specific chemical method must be used to obtain a confirmed result. Gas chromatography/mass spectrometry (GC/MS) is the preferred confirmatory method.^{2, 3}
- 2. It is possible that technical or procedural errors, as well as other interfering substances in the urine specimen may cause erroneous results.
- 3. Adulterants, such as bleach and/or alum, in urine specimens may produce erroneous results regardless of the analytical method used. If adulteration is suspected, the test should be repeated with another urine specimen.
- 4. A positive result indicates presence of the drug or its metabolites but does not indicate level of intoxication, administration route or concentration in urine.
- 5. A negative result may not necessarily indicate drug-free urine. Negative results can be obtained when drug is present but below the cut-off level of the test.
- 6. The test does not distinguish between drugs of abuse and certain medications.
- 7. A positive result might be obtained from certain foods or food supplements.

PERFORMANCE CHARACTERISTICS

Accuracy

A side-by-side comparison was conducted using the Multi-Drug One Step Screen Test Panel (Urine) and commercially available drug rapid tests. Testing was performed on approximately 300 specimens previously collected from subjects presented for Drug Screen Testing. Presumptive positive results were confirmed by GC/MS. The following results were tabulated:

% Agreement with Commercial Kit

Specimen	AMP	AMP 500	AMP 300	BAR	BZO	BZO 200	BUI) **	COC	CO 15	THC	MTD
Positive	97%	*	>99%	>99%	90%	*	88	%	95%	>99	% 98%	>99%
Negative	>99%	*	>99%	99%	97%	*	>99	9%	>99%	6 >99	% >99%	>99%
Total	98%	*	>99%	99%	94%	*	97	%	98%	>99	% 99%	>99%
Specimen	MET	MET 500	MET 300	MDM	IA MO		OPI 2000	(DXY	PCP	PPX	TCA
Positive	98%	>99%	*	>999	6 >99	9% >	-99%	Ģ	96%	98%	>99%	95%
Negative	>99%	80%	*	99%	>99	9% >	>99%	ģ	99%	>99%	>99%	>99%
Total	99%	87%	*	99%	>99	9% >	>99%	ģ	98%	99%	>99%	99%

*NOTE: Commercial kit unavailable for comparison testing.

**NOTE: BUP was compared to the self-reported use of Buprenorphine.

Specimen	AMP	AMP 500	AMP 300	BAR	BZO	BZO 200	BU	P*	COO	2	COC 150	THC	MTD
Positive	97%	95%	>99%	92%	97%	98%	989	%	96%	ó	99%	97%	99%
Negative	95%	>99%	99%	98%	95%	99%	999	%	90%	ó	>99%	88%	94%
Total	96%	98%	99%	95%	96%	99%	999	%	93%	ó	99%	91%	96%
Specimen	MET	MET 500	MET 300	MDMA	MOP 30	00 OPI	2000	C	DXY	I	РСР	PPX	TCA**
Positive	99%	>99%	97%	>99%	>99%	>9	9%	9	98%	>	99%	94%	>99%
Negative	94%	96%	>99%	98%	94%	90	1%	9	9%	Ģ	96%	99%	89%
Total	96%	98%	98%	99%	97%	95	i%	9	99%	9	97%	97%	91%

*NOTE: BUP was based on LC/MS data instead of GC/MS.

**NOTE: TCA was based on HPLC data instead of GC/MS.

Analytical Sensitivity

A drug-free urine pool was spiked with drugs to the concentrations at \pm 50% cut-off and \pm 25% cut-off. The results are summarized below.

Drug Conc.	n	A	мР	AMI	P 500	AMI	P 300	BA	R	BZ	zo	BZC	200	BU	JP
(Cut-off range)	п	-	+	-	+	•	+	•	+	•	+	-	+	•	+
0% Cut-off	30	30	0	30	0	30	0	30	0	30	0	90	0	90	0
-50% Cut-off	30	30	0	30	0	30	0	30	0	30	0	90	0	90	0
-25% Cut-off	30	22	8	24	6	27	3	27	3	27	3	81	9	75	15
Cut-off	30	12	18	16	14	13	17	22	8	11	19	55	35	60	30
+25% Cut-off	30	2	28	4	26	4	26	7	23	5	25	27	63	31	59
+50% Cut-off	30	0	30	0	30	0	30	2	28	0	30	0	90	0	90

· Test panels

• Timer · Specimen collection container

Materials Required But Not Provided

Drug Conc.	n	CO)C	COC	C 150	TI	IC	M	TD	M	ЕТ	MET	Г 500	MET	F 300
(Cut-off range)	п	•	+	•	+	•	+	•	+	•	+	•	+	•	+
0% Cut-off	30	30	0	30	0	30	0	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	29	1	30	0	30	0	30	0
-25% Cut-off	30	30	0	24	6	12	18	24	6	30	0	23	7	27	3
Cut-off	30	4	26	14	16	1	29	21	9	18	12	13	17	15	15
+25% Cut-off	30	0	30	7	23	1	29	2	28	1	29	8	22	4	26
+50% Cut-off	30	0	30	0	30	0	30	0	30	0	30	0	30	0	30

Drug Conc.		MD	MA	MOI	P 300	OPI	2000	02	XY	P	СР	PI	PX	TO	CA
(Cut-off range)	n	•	+	•	+	•	+	•	+	•	+	•	+	•	+
0% Cut-off	30	30	0	30	0	30	0	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	30	0	30	0	30	0	30	0
-25% Cut-off	30	26	4	25	5	30	0	30	0	19	11	24	6	22	8
Cut-off	30	17	13	17	13	13	17	18	12	16	14	17	13	17	13
+25% Cut-off	30	4	26	1	29	4	26	6	24	6	24	7	23	5	25
+50% Cut-off	30	0	30	0	30	0	30	0	30	0	30	0	30	0	30

Analytical Specificity

The following tables lists the concentration of compounds (ng/mL) that are detected positive in urine by the Multi-Drug One Step Screen Test Panel (Urine) at 5 minutes.

AMPHETAMINE		METHADONE					
d-Amphetamine	1,000	Methadone	300				
d,l-Amphetamine	3,000	Doxylamine	50,00				
l-Amphetamine	50,000	METHAMPHETAMINE					
Phentermine	3,000	d-Methamphetamine	1,00				
3,4-Methylendioxyamphetamine (MDA)	2,000	p-Hydroxymethamphetamine	30,00				
AMPHETAMINE 500		l-Methamphetamine	8,00				
d-Amphetamine	500	Mephentermine	50,00				
d,l-Amphetamine	1,500	3,4-Methylenedioxymethamphetamine (MDMA)	2,00				
β-Phenylethylamine	50,000	METHAMPHETAMINE 500					
3,4-Methylendioxyamphetamine (MDA)	800	d-Methamphetamine	500				
Phentermine	1,500	p-Hydroxymethamphetamine	15,00				
Tryptamine	50,000	l-Methamphetamine	4,000				
Tyramine	25,000	Mephentermine	25,00				
AMPHETAMINE 300	·	d,l-Amphetamine	75,00				
d-Amphetamine	300	(1R,2S)-(-)-Ephedrine	50,00				
d,l-Amphetamine	390	β-Phenylethylamine	75,00				
l-Amphetamine	50,000	3,4-Methylenedioxymethamphetamine (MDMA)	1,00				
3,4-Methylendioxyamphetamine (MDA)	1,560	d-Amphetamine	50,00				
p-Hydroxyamphetamine	1,560	Chloroquine	12,50				
β-Phenylethylamine	100,000	l-Phenylephrine	100,0				
Tyramine	100,000	METHAMPHETAMINE 300					
p-Hydroxynorephedrine	100,000	d-Methamphetamine	300				
Phenylpropanolamine (d,l-Norephedrine)	100,000	d,l-Amphetamine	100,0				
BARBITURATES		Chloroquine	25,00				
Secobarbital	300	p-Hydroxymethamphetamine	25,00				
Amobarbital	300	l-Methamphetamine	3,12				
Alphenol	150	3,4-Methylenedioxymethamphetamine (MDMA)	780				
Aprobarbital	200	Mephentermine	50,00				
Butabarbital	75	(1R,2S)-(-)-Ephedrine	100,0				
Butethal	100	l-Epinephrine	50,00				
Butalbital	2,500	Ephedrine	100,0				
Cyclopentobarbital	600	Fenfluramine	12,50				
Pentobarbital	300	Trimethobenzamide	25,00				
Phenobarbital	100	METHYLENEDIOXYMETHAMPHETAMI					
OXYCODONE	·	3,4-Methylenedioxymethamphetamine (MDMA)	500				
Oxycodone	100	3,4-Methylenedioxyamphetamine (MDA)	3,000				
Hydrocodone	6,250	3,4-Methylenedioxyethylamphetamine (MDEA)	300				
Hydromorphone	50,000	BUPRENORPHINE					
Levorphanol	50,000	Buprenorphine	10				
Naloxone	37,500	Norbuprenorphine	20				
Naltrexone	37,500	Buprenorphine 3-D-glucuronide	15				
Oxymorphone	200	Norbuprenorphine 3-D-glucuronide	200				

BENZODIAZEPINES	200	PROPOXYPHENE	200
Oxazepam	300	d-Propoxyphene	300
Alprazolam	196	d-Norpropoxyphene	300
α-Hydroxyalprazolam	1,262	BENZODIAZEPINES 200	
Bromazepam	1,562	Alprazolam	195
Chlordiazepoxide	1,562	α-Hydroxyalprazolam	1,562
Clobazam	98	Bromazepam	390
Clonazepam	781	Chlordiazepoxide	780
Clorazepate	195	Clobazam	390
Delorazepam	1,562	Clorazepate	1,562
Desalkylflurazepam	390	Desalkylflurazepam	1,000
Diazepam	195	Diazepam	200
Estazolam	2,500	Estazolam	780
Flunitrazepam	390	Flunitrazepam	12,500
d,l-Lorazepam	1,562	(+) Lorazepam	100,000
RS-Lorazepam glucuronide	1,562	Midazolam	6,250
Midazolam	12,500	Nitrazepam	100
Nitrazepam	98	Norchlordiazepoxide	3,125
Norchlordiazepoxide	195	Nordiazepam	780
Nordiazepam	390	Oxazepam	200
Temazepam	98	Sertraline	12,500
Triazolam	2,500	Temazepam	12,500
MORPHINE 300	2,500	Triazolam	50,000
Morphine	300	7-Aminoflunitrazepam	200
Codeine	300	7-Aminonitrazepam 7-Aminonitrazepam	5,000
	6,250	*	>100,00
Ethylmorphine Hydrocodone	50,000	7-Aminoclonazepam COCAINE	>100,00
	,		200
Hydromorphone	3,125	Benzoylecgonine	300
Levorphanol	1,500	Cocaine	780
6-Monoacetylmorphine (6-MAM)	400	Cocaethylene	12,500
Morphine 3-β-D-glucuronide	1,000	Ecgonine	32,000
Norcodeine	6,250	COCAINE 150	
Normorphine	100,000	Benzoylecgonine	150
Oxycodone	30,000	Cocaine	400
Oxymorphone	100,000	Cocaethylene	6,250
Procaine	15,000	Ecgonine	12,500
Thebaine	6,250	Ecgonine methylester	50,000
OPIATE 2000		MARIJUANA	
Morphine	2,000	11-nor-∆ ⁹ -THC-9 COOH	50
Codeine	2,000	Cannabinol	20,000
Ethylmorphine	5,000	11-nor-Δ ⁸ -THC-9 COOH	30
Hydrocodone	12,500	Δ ⁸ -THC	15,000
Hydromorphone	5,000	Δ ⁹ -THC	15,000
Levorphanol	75,000	TRICYCLIC ANTIDEPRESSANTS	
6-Monoacetylmorphine (6-MAM)	5,000	Nortriptyline	1,000
Morphine 3-β-D-glucuronide	2,000	Nordoxepin	1,000
Norcodeine	12,500	Trimipramine	3,000
Normorphine	50,000	Amitriptyline	1,500
Oxycodone	25,000	Promazine	1,500
Oxymorphone	25,000	Desigramine	200
Procaine	150,000	Imipramine	400
Thebaine	100,000	Clomipramine	12,500
PHENCYCLIDINE	100,000	Doxepin	2.000
Phencyclidine	25	Maprotiline	2,000
		Invian Office	2.000

Cross-Reactivity

A study was conducted to determine the cross-reactivity of the test with compounds in either drug-free urine or Amphetamine, Amphetamine 500, Amphetamine 300, Barbiturates, Benzodiazepines, Benzodiazepines 200, Buprenorphine, Cocaine, Cocaine 150, Marijuana, Methadone, Methamphetamine, Methamphetamine 500, Methamphetamine 300, Methylenedioxymethamphetamine, Morphine 300, Opiate 2000, Oxycodone, Phencyclidine, Propoxyphene, Tricyclic Antidepressants positive urine. The following compounds show no cross-reactivity when tested with the Multi-Drug One Step Screen Test Panel (Urine) at a concentration of 100 µg/mL.

	Non Cross-R	eacting Compounds	
Acetophenetidin	1-Cotinine	Ketamine	d-Pseudoephedrine
N-Acetylprocainamide	Creatinine	Ketoprofen	Quinidine
Acetylsalicylic acid	Deoxycorticosterone	Labetalol	Quinine
Aminopyrine	Dextromethorphan	Loperamide	Salicylic acid
Amoxicillin	Diclofenac	Meprobamate	Serotonin
Ampicillin	Diflunisal	Methoxyphenamine	Sulfamethazine
l-Ascorbic acid	Digoxin	Methylphenidate	Sulindac
Apomorphine	Diphenhydramine	Nalidixic acid	Tetracycline
Aspartame	Ethyl-p-aminobenzoate	Naproxen	Tetrahydrocortisone,
Atropine	β-Estradiol	Niacinamide	3-Acetate
Benzilic acid	Estrone-3-sulfate	Nifedipine	Tetrahydrocortisone
Benzoic acid	Erythromycin	Norethindrone	Tetrahydrozoline
Bilirubin	Fenoprofen	Noscapine	Thiamine
d,l-Brompheniramine	Furosemide	d,l-Octopamine	Thioridazine
Caffeine	Gentisic acid	Oxalic acid	d,l-Tyrosine
Cannabidiol	Hemoglobin	Oxolinic acid	Tolbutamide
Chloralhydrate	Hydralazine	Oxymetazoline	Triamterene
Chloramphenicol	Hydrochlorothiazide	Papaverine	Trifluoperazine
Chlorothiazide	Hydrocortisone	Penicillin-G	Trimethoprim
d,l-Chlorpheniramine	o-Hydroxyhippuric acid	Perphenazine	d,l-Tryptophan
Chlorpromazine	3-Hydroxytyramine	Phenelzine	Uric acid
Cholesterol	d,l-Isoproterenol	Prednisone	Verapamil
Clonidine	Isoxsuprine	d,l-Propanolol	Zomepirac
Cortisone			

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