

Precision SalivaStik™

Multi-Saliva Drugs of Abuse Rapid Test Device

INTENDED USE

The Multi-Saliva Drugs of Abuse Rapid Test Device is a rapid visual immunoassay for the qualitative, presumptive detection of drugs of abuse in human oral fluid specimens. The test system consists of one or two membrane strips mounted in a plastic cassette. This test detects combinations of the following drugs at the concentrations listed below. Specific combinations will vary according to the test in question:

Test	Calibrator	Cut-off(ng/mL)
Amphetamine (AMP)	D-Amphetamine	50
Benzodiazepine (BZO)	Oxazepam	10
Buprenorphine(BUP)	Buprenorphine	5
Cocaine (COC)	Cocaine	20
Cocaine (COC)	Cocaine	50
Cotinine(COT)	Cotinine	50
EDDP(EDDP)	2-Ethyliden-1,5-Dimethyl-3,3-Diphenylpyrrolidine	20
Ketamine (KET)	Ketamine	50
Marijuana (THC)	11-nor- Δ^9 -THC-9 COOH	12
Metadone (MTD)	Metadone	30
Methamphetamine (MET)	D-Methamphetamine	50
Opiates (OPI)	Morphine	40
Opiates (OPI)	Morphine	50
Oxycodone(OXY)	Oxycodone	20
Phencyclidine (PCP)	Phencyclidine	10
Propoxyphene(PPX)	Propoxyphene	50
Barbiturate(BAR)	Barbiturate	50
Ecstasy (MDMA)	3,4-Methylenedioxymethamphetamine	50
FYL (Fentanyl)	Fentanyl	10

PRINCIPLE

The Multi-Saliva Drugs of Abuse Rapid Test Device is an immunoassay based on the principle of competitive binding. Drugs that may be present in the oral fluid specimen compete against their respective drug conjugate for binding sites on their specific antibody.

During testing, a portion of the oral fluid specimen migrates upward by capillary action. A drug, if present in the oral fluid specimen below its cut-off concentration, will not saturate the binding sites of its specific antibody. The antibody will then react with the drug-protein conjugate and a visible colored line will show up in the test line region of the specific drug strip. The presence of drug above the cut-off concentration in the oral fluid specimen will saturate all the binding sites of the antibody. Therefore, the colored line will not form in the test line region.

A drug-positive oral fluid specimen will not generate a colored line in the specific test line region of the strip because of drug competition, while a drug-negative oral fluid specimen will generate a line in the test line region because of the absence of drug competition. 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.

MATERIALS

Materials Provided

Individually packed test devices

Package insert

Materials Required but Not provided

Timer

Positive and negative controls

PRECAUTIONS

- For professional *in vitro* diagnostic use only.
- Do not use after the expiration date indicated on the package. Do not use the test if the foil pouch is damaged. Do not reuse tests.
- Read the entire procedure carefully prior to testing.
- Do not eat, drink or smoke in the area where specimens and kits are handled. Handle all specimens as if they contain infectious agents. Observe established precautions against microbiological hazards throughout the procedure and follow standard procedures for the proper disposal of specimens. Wear protective clothing such as laboratory coats, disposable gloves and eye protection when specimens are assayed.
- Humidity and temperature can adversely affect results.
- Used testing materials should be discarded in accordance with local regulations.

STORAGE AND STABILITY

- The kit should be stored at 2-30°C until the expiry date printed on the sealed pouch.
- The test must remain in the sealed pouch until use.
- Do not freeze.
- Kits should be kept out of direct sunlight.
- Care should be taken to protect the components of the kit from contamination. Do not use if there is evidence of microbial contamination or precipitation. Biological contamination of dispensing equipment, containers or reagents can lead to false results.

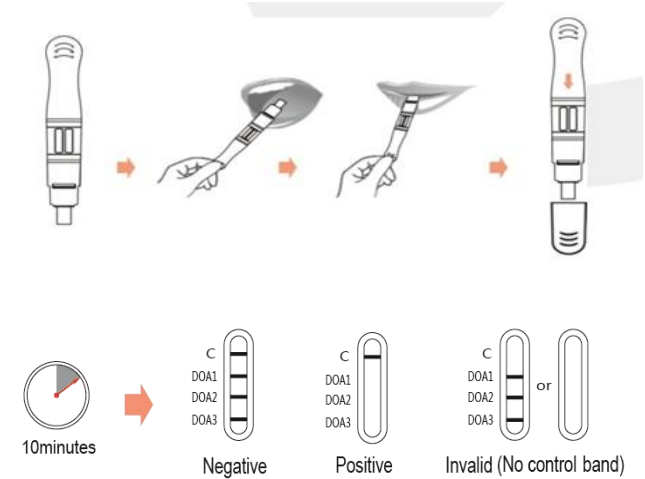
SPECIMEN COLLECTION AND STORAGE

- The Multi-Saliva Drugs of Abuse Rapid Test Device is intended for use with human oral fluid specimens only.
- Oral fluid specimens must be collected according to the directions in the Procedure section of this package insert.
- Perform testing immediately after specimen collection.
- If specimens are to be shipped, pack them in compliance with all applicable regulations for transportation of etiological agents.

PROCEDURE

Bring tests, specimens, and/or controls to room temperature (15-30°C) before use. Donors should avoid placing anything (including food, drink, gum and tobacco products) in their mouth for at least 10 minutes prior to specimen collection.

- Remove the cap by holding the sides and pulling gently. This will expose the collection pad.
- Place the collection pad underneath the tongue for approximately 30 seconds to collect saliva. Instruct the donor to hold the device in place with hand.
- Remove from mouth as soon as color move in both of the test windows. Re-cap the device.
- As the test begins to work, color will migrate across the membrane.**
Wait for the colored band(s) to appear. The result should be read at 10 minutes. Do not interpret the result after 20 minutes.



INTERPRETATION OF RESULTS

(See previous illustration)

POSITIVE: Only one colored band appears, in the control region (C). No colored band appears in the test region (T) for the drug in question. A positive result indicates that the drug concentration exceeds the detectable level.

NEGATIVE: Two colored bands appear on the membrane. One band appears in the control region (C) and another band appears in the test region (T) for the drug in question. A negative result indicates that the drug concentration is below the detectable level.

INVALID: Control band fails to appear. Results from any test which has not produced a control band at the specified read time must be discarded. Please review the procedure and repeat with a new test. If the problem persists, discontinue using the kit immediately and contact your local distributor.

NOTE:

- The intensity of color in the test region (T) may vary depending on the concentration of analytes

present in the specimen. Therefore, any shade of color in the test region (T) should be considered negative. Please note that this is a qualitative test only, and cannot determine the concentration of analytes in the specimen.

- Insufficient specimen volume, incorrect operating procedure or expired tests are the most likely reasons for control band failure.

QUALITY CONTROL

- Internal procedural controls are included in the test. A colored band appearing in the control region (C) is considered an internal positive procedural control, confirming sufficient specimen volume and correct procedural technique.
- External controls are not supplied with this kit. It is recommended that positive and negative controls be tested as a good laboratory practice to confirm the test procedure and to verify proper test performance.

LIMITATIONS OF THE TEST

- The Multi-Saliva Drugs of Abuse Rapid Test Device is for professional *in vitro* diagnostic use, and should be only used for the qualitative detection of drugs of abuse in oral fluid.
- This assay provides a preliminary analytical test result only. A more specific alternative chemical method must be used in order to obtain a confirmed analytical result. Gas chromatography/mass spectrometry (GC/MS) has been established as the preferred confirmatory method by the National Institute on Drug Abuse (NIDA). Clinical consideration and professional judgment should be applied to any test result, particularly when preliminary positive results are indicated.
- There is a possibility that technical or procedural errors as well as other substances and factors may interfere with the test and cause false results.
- A positive result indicates the presence of a drug/metabolite only, and does not indicate or measure intoxication.
- A negative result does not at any time rule out the presence of drugs/metabolites in urine, as they may be present below the minimum detection level of the test.
- This test does not distinguish between drugs of abuse and certain medications.

PERFORMANCE CHARACTERISTICS

A. Sensitivity

A phosphate-buffered saline (PBS) pool was spiked with drugs to target concentrations of \pm 50% cut-off and \pm 25% cut-off and tested with the Multi-Saliva Drugs of Abuse Rapid Test Device. The results are summarized below.

Drug Conc. (Cut-off range)	n	AMP		BUP		BZO		COC	
		-	+	-	+	-	+	-	+
0% Cut-off	30	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	30	0
-25% Cut-off	30	30	0	28	2	30	0	29	1
Cut-off	30	18	12	11	19	14	16	12	18
+25% Cut-off	30	2	28	8	22	4	26	2	28
+50% Cut-off	30	0	30	0	30	0	30	0	30

Drug Conc. (Cut-off range)	n	COT		EDDP		KET		MET	
		-	+	-	+	-	+	-	+
0% Cut-off	30	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	30	0
-25% Cut-off	30	30	0	30	0	27	3	30	0
Cut-off	30	11	19	13	17	9	21	13	17
+25% Cut-off	30	1	29	2	28	3	27	3	27
+50% Cut-off	30	0	30	0	30	0	30	0	30

Drug Conc. (Cut-off range)	n	OPI		MTD		OXY		PCP	
		-	+	-	+	-	+	-	+
0% Cut-off	30	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	30	0
-25% Cut-off	30	28	2	30	0	28	2	28	2
Cut-off	30	10	20	10	20	10	20	11	19
+25% Cut-off	30	9	21	2	28	4	26	5	25
+50% Cut-off	30	0	30	0	30	0	30	0	30

Drug Conc. (Cut-off range)	n	THC		BAR		PPX	
		-	+	-	+	-	+
0% Cut-off	30	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0
-25% Cut-off	30	30	0	27	3	30	0
Cut-off	30	10	20	9	21	10	20
+25% Cut-off	30	5	25	3	27	4	26
+50% Cut-off	30	0	30	0	30	0	30

Drug Conc.	n	MDMA	COC 50	OPI 50	FYL 10
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(Cut-off range)		-	+	-	+	-	+	-	+
0% Cut-off	30	30	0	30	0	30	0	30	0
-50% Cut-off	30	30	0	30	0	30	0	30	0
-25% Cut-off	30	25	5	30	0	27	3	25	5
Cut-off	30	14	16	10	20	9	21	12	18
+25% Cut-off	30	4	26	4	26	3	27	8	22
+50% Cut-off	30	0	30	0	30	0	30	0	30

B. Specificity

The following table lists the concentrations of compounds (ng/mL) above which the Multi-Saliva Drugs of Abuse Rapid Test Device identified positive results at 10 minutes.

Amphetamine-Related Compounds		Δ ⁹ -Tetrahydrocannabinol	4,000
D-Amphetamine	50	11-hydroxy-Δ ⁹ -THC	300
L-Amphetamine	4,000	Propoxyphene -Related Compounds	
(+)-3,4-Methylenedioxyamphetamine	150	Propoxyphene (PPX)	50
Phentermine	40,000	D-Norpropoxyphene	200
PMA	125	Fentanyl -Related Compounds	
Tyramine	3,000	Fentanyl	10
Barbiturate -Related Compounds		Norfentanyl	200
Barbiturate (BAR)	50		
Allobarbitol	200	Methadone -Related Compounds	
Alphenal	100	Methadone	30
Amobarbital	100	Alpha-Methadol	125
Aprobarbital	30	Biperiden	80,000
Butabarbital	15	Doxylamine	12,500
Butalbital	400	2-Ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine	10,000
Butethal	30	Phencyclidine	12,500
Cyclopentobarbital	60	Pheniramine	25,000
Pentobarbital	150	Methamphetamine-Related Compounds	
Phenobarbital	300	D-Methamphetamine	50
Benzodiazepine-Related Compounds		Fenfluramine	3,000
Oxacepam	10	L-Methamphetamine	500
Alprazolam	15	L-Phenylephrine	2,500
Bromazepam	8	MDEA	400
Chlordiazepoxide	10	3,4-Methylenedioxymethamphetamine	75
Clonazepam	40	Mephentermine	200
Clorazepate	20	PMMA	50
Cibazam	6	Procaine	2,500
Diazepam	15	Ketamine-Related Compounds	
Estazolam	10	Ketamine(KET)	50
Desalkylflurazepam	8	Norketamine	50
Flunitrazepam	10	Dextromethorphan	25
Flurazepam	10	Hydrocodone	50
Lorazepam	20	Hydromorphone	100
Medazepam	10	Temazepam	8
Nitrazepam	10	Triazola	15
Nordiazepam	6	Opiates -Related Compounds	
Prazepam	20	Morphine	40
Temazepam	8	Codeine	10
Triazola	15	Diacetylmorphine (Heroin)	50
Buprenorphine -Related Compounds		Ethylmorphine	24
Buprenorphine	5	Hydrocodone	50
Nalorphine	10,000	Hydromorphone	100
Oxycodone	25,000	6-Monoacetylmorphine (6-MAM)	25
Oxymorphone	25,000	Morphine-3- β-d-glucuronide	50
Thebaine	5,000	Nalorphine	10,000
Cocaine-Related Compounds		Oxycodone	25,000
Cocaine	20	Oxymorphone	25,000
Benzoylcegonine	200	Thebaine	5,000
Ecgonine	100,000	Opiates -Related Compounds	
Ecgonine methyl ester	10,000	Morphine	50
Cocaine-Related Compounds		Codeine	15
Cocaine	50	Diacetylmorphine (Heroin)	60
Benzoylcegonine	500	Ethylmorphine	30

Ecgonine	>100,000	Hydrocodone	60
Ecgonine methyl ester	50,000	Hydromorphone	125
Cotinine-Related Compounds		6-Monoacetylmorphine (6-MAM)	60
Cotinine	50	Morphine-3- β-d-glucuronide	60
Buprenorphine	>100,000	Nalorphine	12,500
EDDP -Related Compounds		Oxycodone	31,250
EDDP	20	Oxymorphone	31,250
Meperidine	20,000	Thebaine	6,250
Methadone	20,000	Oxycodone-Related Compounds	
Norfentanyl	20,000	Oxycodone	20
Phencyclidine	20,000	Hydrocodone	1,000
Promazine	10,000	Hydromorphone	6,250
Promethazine	5,000	Naloxone	6,250
Prothipendyl	10,000	Oxymorphone	1,000
Prozine	2,500	Phencyclidine-Related Compounds	
Ecstasy -Related Compounds		Phencyclidine (PCP)	10
3,4-Methylenedioxymethamphetamine	50	Hydrocodone	2,000
3,4-Methylenedioxymphetamine	250	Hydromorphone	2,000
3,4-Methylenedioxyethylamphetamine	60	Morphine-3- β-d-glucuronide	20,000
Paramethoxyamphetamine	1,600	Nalorphine	10,000
Paramethoxymethamphetamine	160		
Marijuana -Related Compounds			
11-nor-Δ ⁹ -THC-9 COOH	12		
Δ ⁸ -Tetrahydrocannabinol	2,000		

A study was conducted to determine the cross-reactivity of the test with compounds spiked into drug-free PBS stock. The following compounds demonstrated no false positive results on the Multi-Saliva Drugs of Abuse Rapid Test Device when tested at concentrations up to 100 ug/mL.

(-) Ephedrine (Except MET)	Chlorpheniramine	Oxalic Acid
(+)-Naproxen	Creatine	Penicillin-G
(+/-) Ephedrine (Except MET)	Dextromethorphan (Except KET)	Pheniramine
4-Dimethylaminoantipyrene	Dextrorphan tartrate (Except KET)	Phenothiazine
Acetaminophen	Dopamine	Procaine
Acetone	Erythromycin	Protonix
Albumin	Ethanol	Pseudoephedrine
Amitriptyline	Furosemide	Quinidine
Ampicillin	Glucose	Ranitidine
Aspartame	Guaiacol Glyceryl Ether	Sertraline
Aspirin	Hemoglobin	Tyramine
Benzocaine	Imipramine	Trimeprazine
Bilirubin	(+/-)-Isoproterenol	Venlafaxine
b-Phenylethyl-amine	Methadone (Except MTD)	Ibuprofen
Caffeine	Vitamin C (Ascorbic Acid)	Lidocaine
Chloroquine (Except MET)		

LITERATURE REFERENCES

- Baselt RC. Disposition of Toxic Drugs and Chemicals in Man. 2nd ed. Davis: Biomedical Publications; 1982.
- Hawks RL, Chiang CN, eds. Urine Testing for Drugs of Abuse. Rockville: Department of Health and Human Services, National Institute on Drug Abuse; 1986.
- Substance Abuse and Mental Health Services Administration. Mandatory Guidelines for Federal Workplace Drug Testing Programs. 53 Federal Register; 1988.
- McBay AJ. Drug-analysis technology--pitfalls and problems of drug testing. Clin Chem. 1987 Oct; 33 (11 Suppl): 33B-40B.
- Gilman AG, Goodman LS, Gilman A, eds. Goodman and Gilman's The Pharmacological Basis of Therapeutics. 6th ed. New York: Macmillan; 1980.

GLOSSARY OF SYMBOLS

ρ	Catalog number	g	Temperature limitation
ι	Consult instructions for use	Λ	Batch code
I	<i>In vitro</i> diagnostic medical device	ε	Use by
μ	Manufacturer	σ	Do not reuse

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