



T-Cup Multi-Drug Urine Test Cup

Catalogue No. See Box Label

The SAFElife® T-Cup Multi-Drug Urine Test Cup contains competitive binding, lateral flow immunochemical/pharmacology assays for qualitative and simultaneous detection of 6-Monacoetylmorphine, Amphetamine, Sebacobarbitol, Buprenorphine, Oxycodone, Cocaine, Cotinine, 2-ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine (EDDP), Ethyl Glucuronide, Fentanyl, Gabapentin, Hydromorphone, Synthetic Cannabinoids, Ketamine, Kratom, Lysergic acid diethylamide, 25 Security Stems (100), Methamphetamine, Morphine, Methadone, Oxycodone, Phenylethylamine, Pregabalin, Propoxyphene, Nortriptyline, Cannabinoids, Tramadol and Alcohol in human urine with below cutoff concentrations and appropriate detection time:

Drug (Identifier)	Calibrator	Cut-off Level	Minimum Detection Time	Maximum Detection Time
6-Monacoetylmorphine (6-MAM)	6-Monacoetylmorphine	10 ng/mL	2 hours	8 hours
Amphetamine (AMP300)	d-Amphetamine	300 ng/mL	2-7 hours	1-2 days
Amphetamine (AMP500)	d-Amphetamine	500 ng/mL	2-7 hours	1-2 days
Sebacobarbitol (BAR)	d-Amphetamine	1000 ng/mL	2-7 hours	1-2 days
Buprenorphine (BUP)	Buprenorphine	5 ng/mL	2-4 hours	1-4 days
Buprenorphine (BUP10)	Buprenorphine	10 ng/mL	4 hours	1-3 days
Oxazepam (BZO100)	Oxazepam	100 ng/mL	2-7 hours	1-2 days
Oxazepam (BZO200)	Oxazepam	200 ng/mL	2-7 hours	1-2 days
Oxycodone (COX100)	Oxycodone	300 ng/mL	2-7 hours	1-2 days
Cocaine (COC100)	Benzoylecgonine	100 ng/mL	1-4 hours	2-4 days
Cocaine (COC300)	Benzoylecgonine	150 ng/mL	1-4 hours	2-4 days
Cocaine (COC100)	Benzoylecgonine	300 ng/mL	1-4 hours	2-4 days
Cocaine (COT)	Cotinine	200 ng/mL	2-8 hours	1-7 days
EDDP100	2-ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine	300 ng/mL	3-8 hours	1-3 days
EDDP300	2-ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine	300 ng/mL	3-8 hours	1-3 days
Ethyl Glucuronide (EG300)	Ethyl Glucuronide	300 ng/mL	1-2 hours	Up to 3+ days
Ethyl Glucuronide (EG500)	Ethyl Glucuronide	500 ng/mL	1-2 hours	Up to 3+ days
Fentanyl (FTY)	Nortriptyline	20 ng/mL	1-4 hours	1-3 days
Gabapentin (GAB)	Gabapentin	2000 ng/mL	5-7 hours	Up to 2 days
Hydromorphone (HMO)	Hydromorphone	300 ng/mL	4-6 hours	1-2 days
Synthetic Cannabinoid (K2)	JWH-073 Butanoic Acid	50 ng/mL	8-12 hours	Up to 5+ days
Ketamine (KET300)	Ketamine	300 ng/mL	2-4 hours	2-3 days
Ketamine (KET1000)	Ketamine	1000 ng/mL	2-4 hours	2-3 days
Kratom (KRA100)	Mitragynine	100 ng/mL	7 hours	5-6 days
Kratom (KRA300)	Mitragynine	300 ng/mL	7 hours	5-6 days
Lysergic acid diethylamide (LSD)	Lysergic acid diethylamide	20 ng/mL	2.5 hours	Up to 5+ days
Methylenedioxyamphetamphetamine (MDMA)	3,4-Methylenedioxyamphetamphetamine (MDMA)	500 ng/mL	2-7 hours	2-4 days
Methamphetamine (MET300/mAMP300)	Di(-)-Methamphetamine	300 ng/mL	2-7 hours	2-4 days
Methamphetamine (MET500/mAMP500)	Di(-)-Methamphetamine	500 ng/mL	2-7 hours	2-4 days
Methamphetamine (MET1000/mAMP1000)	Di(-)-Methamphetamine	1000 ng/mL	2-7 hours	2-4 days
Morphine (MOP100/OP100)	Morphine	300 ng/mL	2 hours	2-3 days
Morphine (MOP300/OP300)	Morphine	100 ng/mL	2 hours	2-3 days
Methadone (MTD200)	Methadone	200 ng/mL	3-8 hours	1-3 days
Methadone (MTD300)	Methadone	300 ng/mL	3-8 hours	1-3 days
Methaqualone (MQL)	Methaqualone	300 ng/mL	6-8 hours	Up to 7+ days
Opiate (OP1000)	Morphine	2000 ng/mL	2 hours	2-3 days
Oxycodone (OXY)	Oxycodone	100 ng/mL	4 hours	1-3 days
Phencyclidine (PCP)	Phencyclidine	25 ng/mL	4-6 hours	7-14 days
Pregabalin (PGB)	Pregabalin	500 ng/mL	6-8 hours	1-3 days
Propoxyphene (PPX)	Propoxyphene	300 ng/mL	2 hours	2-3 days
Nortriptyline (TCA)	Nortriptyline	1000 ng/mL	8-12 hours	2-7 days
Cannabinoids (THCS)	11-nor-Δ ⁹ -THC-9-COOH	15 ng/mL	2 hours	Up to 5+ days
Cannabinoids (THCS2)	11-nor-Δ ⁹ -THC-9-COOH	25 ng/mL	2 hours	Up to 5+ days
Cannabinoids (THCS4)	11-nor-Δ ⁹ -THC-9-COOH	40 ng/mL	2 hours	Up to 5+ days
Cannabinoids (THCS50)	11-nor-Δ ⁹ -THC-9-COOH	50 ng/mL	2 hours	Up to 5+ days
Tramadol (TRA 100)	Tramadol	100 ng/mL	8-12 hours	3-7 days
Tramadol (TRA 200)	Tramadol	200 ng/mL	8-12 hours	3-7 days
Tramadol (TRA 1000)	Tramadol	1000 ng/mL	8-12 hours	3-7 days
Alcohol (ETOH)	Alcohol	0.04 g/dL	-	-

Configurations of the SAFElife® T-Cup Multi-Drug Urine Test Cup can consist of any combination of the above listed drug analyses.

It is intended for forensic use only.

It is not intended to distinguish between prescription use or abuse of these drugs. Professional judgment should be applied to any drug of abuse test result, particularly in evaluating a preliminary positive result.

The tests provide only preliminary results. To obtain a confirmed analytical result, a more specific alternate chemical method must be used. Chromatography/Mass Spectrometry (GC/MS) or Liquid Chromatography/Tandem Mass Spectrometry (LC/MS-MS) is the recommended confirmatory method.

WARNINGS AND PRECAUTIONS

- The test kit is for external use only.
- Discard after first use. The test kit cannot be used more than once.
- Do not use the test kit beyond expiration date.
- Do not use the test kit if the pouch is punctured or not well sealed.
- Keep out of the reach of children.

Approximate Alcohol Concentration

0.0 mg/100 mL	40 mg/100 mL	80 mg/100 mL	200 mg/100 mL
	(0.04%)	(0.08%)	(0.2%)

Invalid

The test should be considered invalid if only the edge of the reaction pad turned color that might be ascribed to insufficient sampling. Another test should be run to re-evaluate the specimen. If test still fails, please contact the distributor, with the lot number.

What is the False Positive Test?

The definition of the false positive test would be an instance where a substance is identified incorrectly by the SAFElife® T-Cup Multi-Drug Urine Test Cup. The most common causes of the false positive test are cross reactants. Certain foods and medicines, diet plan drugs and nutritional supplements may cause the false positive test result.

What is the False Negative Test?

The definition of the false negative test is that the initial drug is present but isn't detected by the SAFElife® T-Cup Multi-Drug Urine Test Cup. If the specimen is diluted or adulterated, it may cause false negative result.

HOW TO COLLECT URINE?

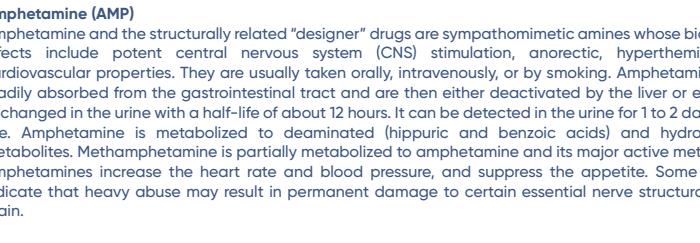
- Remove the test cup from the foil pouch by tearing at the notch. Use it as soon as possible. Instruct the donor to remove the test cup lid and void directly into the test cup until reach the Minimum Urine Level. After the urine has been collected, the test cup should be used immediately. If the specimen is not collected, instruct the donor to provide urine specimen again with another new test cup. Wipe off any splashes or spills that may be on the outside of the cup. It is recommended to wear gloves when handling the test cup with urine specimen.
- Observe the temperature strip affixed on the test cup between 2 to 4 minutes after urine is voided into the cup. The temperature between 90°F to 100°F (32°C-38°C) indicates the fresh uncontaminated specimen. If the temperature is out of this range, instruct the donor to provide urine specimen again with another new test cup.

TEST PROCEDURE

Test should be performed at room temperature 65°F-86°F (18°C-30°C).

- After the urine has been collected, properly, tighten the lid and place the cup on a flat surface.
- Peel off the label from right to left.
- For the adulteration strip(s) if equipped, read results immediately, or at 30 seconds, or at 45 seconds and compare each adulterant pad to verify pad color is within acceptable range according to the Adulteration Color Comparison Chart. If the results indicate adulteration, do not read the drug test results. Instruct the donor to provide urine specimen again with another new test cup.
- For the alcohol test, read the alcohol test result at 2 minutes. Do not read results after 2 minutes.
- For the drug tests, read the drug test results at 5 minutes. Do not read results after 5 minutes.

- Remove the test cup from the foil pouch by tearing at the notch. Use it as soon as possible. Instruct the donor to remove the test cup lid and void directly into the test cup until reach the Minimum Urine Level. After the urine has been collected, the test cup should be used immediately. If the specimen is not collected, instruct the donor to provide urine specimen again with another new test cup. Wipe off any splashes or spills that may be on the outside of the cup. It is recommended to wear gloves when handling the test cup with urine specimen.
- Observe the temperature strip affixed on the test cup between 2 to 4 minutes after urine is voided into the cup. The temperature between 90°F to 100°F (32°C-38°C) indicates the fresh uncontaminated specimen. If the temperature is out of this range, instruct the donor to provide urine specimen again with another new test cup.



READING THE RESULTS

Barbiturates are a class of central nervous system depressants. They have a wide range of half-life of 2 to 40 hours and can be detected in the urine for 1 to 4 days after use. Phenobarbital is a long acting barbiturate derivative that has been used as a daytime sedative and very extensively as an anticonvulsant. Pentobarbital and sebacobarbitol are two examples of a short acting barbiturate sedative. Abuse of barbiturates can lead not only to impaired motor coordination and mental disorder, but also to respiratory collapse, coma, and even death. Barbiturates are taken orally and by intravenous injection. Pentobarbital and sebacobarbitol are generally excreted in urine as metabolites, while the long-acting barbiturates will primarily appear unchanged.

Buprenorphine (BUP)

Buprenorphine is a partial agonistic opioid used in the treatment of opioid addiction. The drug is sold under the trade names Subutex®, Buprenex®, Temgesic® and Suboxone®, all of which contain Buprenorphine HCl alone or in combination with Naloxone HCl. Therapeutically, Buprenorphine is used as a substitution treatment for opioid addicts. A substitution treatment is a form of medical care offered to opiate addicts (primarily heroin addicts) based on a similar or identical substance to the drug normally used in substitution therapy. Buprenorphine is as effective as Methadone but demonstrates a lower level of physical dependence. The plasma half-life of Buprenorphine is 2-4 hours. While complete elimination of a single-dose of the drug can take as long as 6 days, the detection window for the parent drug in urine is thought to be approximately 3 days.

Oxazepam (BZO)

Benzoizapines are the most widely used anxiolytic drugs. They are used extensively as anti-anxiety agents, hypnotics, muscle relaxants and sedatives. They are taken orally or sometimes by injection and have a wide range of half-life from 2 to 40 hours. They can generally be detected for 1 to 2 days after Benzoizapines use. Benzoizapines are metabolized in the liver. Some Benzoizapines and their metabolites are excreted in the urine. Their use can result in drowsiness and/or confusion. Benzoizapines potentiate alcohol and other CNS depressants. Psychological and physical dependence on benzoizapines can develop if high doses of the drug are given over a prolonged period.

Cocaine (COC)

Cocaine derived from leaves of coca plant, is a potent central nervous system stimulant and a local anesthetic. Among the psychological effects induced by using cocaine are euphoria, confidence and sense of increased energy accompanied by increased heart rate, dilation of the pupils, fever, tremors and sweating. Cocaine is excreted in urine primarily as benzoylecgonine in a short period of time.

Morphine (MOP/OP)

The opiates such as heroin, morphine, and codeine are derived from the resin of opium poppy. The principal metabolites of opiates are morphine, morphine-3-glucuronide normorphine and codeine with a half-life of about 3 hours. Heroin is quickly metabolized to morphine. Thus, morphine and morphine glucuronide might be found in the urine of a person who has taken only heroin. The body also changes codeine to morphine. Thus, the presence of morphine (for the metabolite, morphine glucuronide) in the urine indicates heroin, morphine and/or codeine use. The test for Morphine (MOP300/OP300) of the SAFElife® T-Cup Multi-Drug Urine Test Cup yields a positive result when the morphine in urine exceeds 300 ng/mL.

Methadone (MTD)

Methadone is a synthetic analgesic drug that is originally used in the treatment of narcotic addicts. Among the psychological effects induced by using methadone are analgesia, sedation and respiratory depression. Overdose of methadone may cause coma or even death. It is administered orally or intravenously and is metabolized in the liver and excreted in urine as methadone, EDDP, MDMA and methadone. The kidneys are a major route of methadone excretion. Methadone has a biological half-life of 15 to 60 hours.

ng/mL is expected to be up to 2-3 days after nicotine use.

EDDP

EDDP (2-ethylidene-1, 5-dimethyl-3, 3-diphenylpyrrolidine) is the primary metabolite of methadone. Methadone is a synthetic analgesic drug that is originally used in the treatment of narcotic addicts. The detection of EDDP is more beneficial than traditional methadone screening since EDDP exists only in urine after the individual has ingested methadone. The sampling of specimens with the test kit with methadone can be prevented. Secondly, renal clearance of EDDP is not affected by the pH of the urine, therefore the EDDP test provides a more accurate result of methadone ingestion than the methadone parent screening.

Ethyl Glucuronide (EtG)

Ethyl Glucuronide is a direct metabolite of alcohol. Presence in urine may be used to detect recent alcohol intake, even after alcohol is no longer measurable. Traditional laboratory methods detect the actual alcohol in the body, which reflects current intake within the past few hours (depending on how much was consumed). The presence of EtG in urine is a direct indicator that it can be detected in the urine for 2 to 4 days after drinking alcohol, even alcohol is eliminated from the body. Therefore, EtG is a more accurate indicator of the recent intake of alcohol than measuring for the presence of alcohol itself. The EtG test can aid in the diagnosis of drunk driving and alcoholism, which has important significance in the forensic identification and medical examination.

Fentanyl (FTY)

Fentanyl is a potent, synthetic narcotic analgesic with a rapid onset and short duration of action. It was first synthesized by Janssen Pharmaceutische (Belgium) in the late 1950s, and it is approximately 100 times more potent than morphine. Fentanyl is a strong agonist at the μ-opioid receptor. Historically it has been used to treat breakthrough pain and is commonly used in pre-procedures as a pain reliever as well as an anesthetic in combination with a benzodiazepine. Fentanyl is frequently given intrathecally as part of spinal anesthesia or epidurally for epidural anesthesia and analgesia.

Gabapentin (GAB)

Gabapentin (GAB), sold under the brand name Neurontin, is a medication used to treat epilepsy, neuropathic pain, hot flashes, and restless legs syndrome. In epilepsy, it may be used for those with partial motor (approximately 50% of cases), simple partial (approximately 10% of cases), and complex partial (approximately 10% of cases) seizures. It is also used in the treatment of postherpetic neuralgia, and central neuropathic pain. It is also used to relieve nerve pain following shingles (a painful rash due to herpes zoster infection) in adults. The most common side effects of gabapentin include dizziness, fatigue, weakness, ataxia, peripheral edema (swelling of extremities), myasthenia, and tremor. Serious side effects may include an increased risk of suicide, aggressive behavior, and drug reaction with eosinophilia and systemic symptoms.

Hydromorphone (HMO)

Hydromorphone, also known as dihydromorphone or dihydromorphine, is a semi-synthetic strong narcotic analgesic. It is structurally similar to morphine, but its pharmacological effects are more potent. Its side effects are lighter than morphine. It is mainly used for relieving medium-intensity pain caused by cancer, postoperative and soft tissue trauma.

Synthetic cannabinoids (K2)

Synthetic cannabinoids are psychoactive designer drugs derived of natural herbs sprayed with synthetic chemicals that, when consumed, allegedly mimic the effects of cannabis. It is best known by the brand names K2 and Spice. Synthetic cannabinoids act on the body in a similar way to cannabinoids naturally found in cannabis, such as THC. Although synthetic cannabinoids do not produce positive results in drug tests for cannabis, it is possible to detect its metabolites in human urine.

Ketamine (KET)

Ketamine is a sort of medical sedative. The principal metabolites are norketamine. Smoking, snorting, snuffing, and dissolving into drink or alcohol are the primary method of use of ketamine. Ketamine is usually administered in combination with heroin, marijuana, etc. for the relief of moderate to severe pain. Overdose may cause central nervous system effects, do harm to liver and kidney, and even cause death. Ketamine is metabolized in the liver. Over 70% ketamine metabolites and only 5% original drugs are excreted in the urine. They can generally be detected for 2 to 4 hours after ketamine use.

Kratom (KRA)

Kratom (Mitragyna speciosa) is a plant indigenous to Thailand and Southeast Asia. Kratom leaves produce a complex stimulant and opioid-like analgesic effect in Asia. It is often used to stave off withdrawal and manage pain, diarrhea, cough, and opioid withdrawal. Recently, kratom has become widely available in the United States and Europe by means of smoke shops and the Internet. The clinical manifestations of kratom are not well defined and studies are limited, but its safety profile has become a national health concern, primarily due to excessive consumption being linked to increase in hospital visits for septic injury, seizures, coma, and death. The main active ingredients include Mitragynine and 7-Hydroxymitragynine, which can be detected in urine up to 72 hrs¹⁻³.

Lysergic acid diethylamide (LSD)

Lysergic acid diethylamide (LSD) is a white powder or colorless liquid that is a strong semi-artificial hallucinogen. LSD is manufactured from Lysergic acid which occurs naturally in the ergot fungus that grows on wheat and rye. It is a schedule I controlled substance, available in liquid, powder, tablet (microdots), and capsule form. LSD is a non-selective 5-HTagonist, may exert its hallucinogenic effect by interacting with 5-HT_{2A} receptors as a partial agonist and modulating the 5-HT_{2A} receptors, producing a marked slowing of the firing rate of serotonergic neurons. LSD can cause the user's senses, feelings, memory, and self-awareness to intensify and change for 6 to 12 hours. In addition to causing mental confusion, LSD can also cause physical pain, with symptoms in the nervous system, cardiovascular, and digestive systems. Most LSD users use marijuana, heroin, or other drugs together.

Methylenedioxyamphetamphetamine (MDMA)

Methylenedioxyamphetamphetamine (ecstasy) is a designer drug first synthesized in 1914 by a German drug company for the treatment of obesity. Those who take the drug frequently report adverse effects, such as increased muscle tension and sweating. MDMA is not clearly a stimulant, although it has, in common with amphetamine drugs, a capacity to increase blood pressure and heart rate. MDMA does produce some perceptual changes in the form of increased sensitivity to light, difficulty in focusing, and blurred vision in some users. Its mechanism of action is thought to be via release of the neurotransmitter serotonin. MDMA may also release dopamine, although the general opinion is that it is a secondary effect of the drug (Nichols and Oberlander, 1990). The most pervasive effect of MDMA, occurring in virtually all people who take a reasonable dose of the drug, was to produce a clenching of the jaws.

Methamphetamine (MET/mAMP)

Methamphetamine is a potent sympathomimetic agent with therapeutic applications. Acute higher doses lead to enhanced stimulation of the central nervous system and induce euphoria, alertness, and a sense of increased energy and power. More acute responses produce anxiety, paranoia, psychotic behavior, and cardiac dysrhythmias. The pattern of psychomotor activity may appear alert for 15 hours and is excreted in urine as amphetamine and oxidized as desmethylamphetamine and desmethylamphetamine. However, 40% of methamphetamine is excreted unchanged. Thus the presence of the parent compound in the urine indicates methamphetamine use.

Morphine (MOP/OP)

The opiates such as heroin, morphine, and codeine are derived from the resin of opium poppy. The principal metabolites of opiates are morphine, morphine-3-glucuronide normorphine and codeine with a half-life of about 3 hours. Heroin is quickly metabolized to morphine. Thus, morphine and morphine glucuronide might be found in the urine of a person who has taken only heroin. The body also changes codeine to morphine. Thus, the presence of morphine (for the metabolite, morphine glucuronide) in the urine indicates heroin, morphine and/or codeine use. The test for Morphine (MOP300/OP300) of the SAFElife® T-Cup Multi-Drug Urine Test Cup yields a positive result when the morphine in urine exceeds 300 ng/mL.

Methadone (MTD)

Methadone is a synthetic analgesic drug that is originally used in the treatment of narcotic addicts. Among the psychological effects induced by using methadone are analgesia, sedation and respiratory depression. Overdose of methadone may cause coma or even death. It is administered orally or intravenously and is metabolized in the liver and excreted in urine as methadone, EDDP, MDMA and methadone. The kidneys are a major route of methadone excretion. Methadone has a biological half-life of 15 to 60 hours.

Methaqualone (MQL)

Methaqualone is a sedative that falls outside the benzodiazepine and barbiturate classes. It was once a popular pharmacological and recreational drug, but its current use is largely relegated to Africa, particularly South Africa. Because it faced few restrictions when it first entered the market, the drug was widely prescribed and perceived as uniquely safe. We now know methaqualone can be used recreationally and can cause physical dependence. A lot of lore exists around its effects. In reality, it's not a massively unique or especially dangerous drug. It can be compared to barbiturates, ethanol, carisoprodol, and benzodiazepines. Methaqualone is a sedative that increases the activity of the GABA receptors in the brain and nervous system. When GABA activity is increased, blood pressure drops and the breathing and pulse rates slow, leading to a state of deep relaxation. These properties explain why methaqualone was originally mainly prescribed for insomnia. Methaqualone peaks in the bloodstream within several hours, with a half-life of 20-40 hours. Regular users build up a physical tolerance, requiring larger doses for the same effect. Overdose can lead to nervous system shutdown, coma and death.

Opiate (OPI)

Opium refers to any drug that is derived from the opium poppy, including the natural products, morphine and codeine, and the semi-synthetic drugs such as heroin. Opioid is more general, referring to any drug that acts on the opioid receptor. Opioid analgesics comprise a large group of substances which control pain by depressing the central nervous system. Large doses of morphine can produce higher tolerance levels, physiological dependency in users, and may lead to substance abuse. Morphine is excreted unmetabolized, and is also the major metabolite product of codeine and heroin. Morphine is detectable in the urine for several days after an opiate dose. The test for Morphine 2000 (OPI) of the SAFElife® T-Cup Multi-Drug Urine Test Cup yields a positive result when the morphine in urine exceeds 2000 ng/mL.

Oxycodone (OXY)

Oxycodone is known as Oxycotin and Roxicodone. It is an ingredient of Percodan, Percocet, Roxicet and Tylox. Oxycodone is a semi-synthetic opiates derived from opium. Like other opiates, Oxycodone is characterized by its analgesic properties, and the tendency for users to form a physical dependency and develop tolerance with extended use. Oxycodone is usually administered in combination with non-opioid analgesics such as acetaminophen and salicylates for the relief of moderate to severe pain. Oxycodone is a central nervous system depressant that may cause drowsiness, dizziness, lethargy, weakness and confusion. Toxicity in an overdose of Oxycodone can lead to stupor, coma, muscle flaccidity, severe respiratory depression, hypotension, and cardiac arrest. Oxycodone is metabolized in the liver and excreted in urine as oxycodone, oxycodone-3-glucuronide, and oxycodone-5-glucuronide. Oxycodone is detectable in the urine for several days after an opiate dose. The test for Morphine 2000 (OPI) of the SAFElife® T-Cup Multi-Drug Urine Test Cup yields a positive result when the morphine in urine exceeds 2000 ng/mL.

Phencyclidine (PCP)

Phencyclidine is an arylcyclohexylamine that was originally used as an anesthetic agent and a veterinary tranquilizer. Its structure is similar to morphine, but its pharmacological effects are more potent. Its side effects are lighter than morphine. It is mainly used for relieving medium-intensity pain caused by cancer, postoperative and soft tissue trauma.

Pregabalin (PGB)

Pregabalin (PGB) is a synthetic analogue of γ-aminobutyric acid (GABA), which is similar to gabapentin in structure and action, and has antiepileptic, analgesic and anxiolytic activities. About 98% of pregabalin is recovered in the urine as the active drug after radiation labeling. Therefore, pregabalin abuse can be determined directly by measuring the amount of pregabalin in urine.

Propoxyphene (PPX)

Propoxyphene, a synthetic opiate agonist, is structurally similar to methadone. Propoxyphene is a narcotic analgesic used to relieve mild to moderate pain. The principal metabolites are norendropropoxyphene. The combination usage of propoxyphene, aspirin, acetaminophen or other sedatives can lead cooperative interaction. Abuse of propoxyphene can lead nausea, vomiting, constipation, illusion, heart failure, poisoning, lung dropsy and even death. Propoxyphene is metabolized in the liver and excreted in urine as norendropropoxyphene. Thus the presence of the propoxyphene or its metabolites in the urine indicates propoxyphene use.

Nortriptyline (TCA)

TCA (Tricyclic Antidepressants) are commonly used for the treatment of depressive disorders. TCA overdoses can result in profound central nervous system depression, cardiotoxicity and anticholinergic interactions. TCA overdose is the most common cause of death from prescription drugs. TCAs are taken orally or sometimes by injection. TCAs are metabolized in the liver. Both TCAs and their metabolites are excreted in urine mostly in the form of metabolites for up to ten days.

Cannabinoids (THC)

Cannabinoids are hallucinogenic agents derived from the flowering portion of the hemp plant. The active ingredients in Cannabinoids, THC & Cannabinol can be metabolized and excreted as 11-nor-Δ⁹-tetrahydrocannabinol-9-carboxylic acid with a half-life of 24 hours. They can be detected for 1 to 5 days after use. Smoking is the primary method of use of Cannabinoids/cannabis. Higher doses used by abusers produce central nervous system effects altered mood and sensory perceptions, loss of coordination, impaired short-term memory, anxiety, paranoia, depression, confusion, hallucinations and increased heart rate. A tolerance to the cardiac and psychotropic effects can occur, and withdrawal syndrome produces restlessness, insomnia, anorexia and nausea.

Tramadol (TRA)

Tramadol (2-(dimethylamino)ethyl-1-[3-(methoxyphenyl) cyclohexan]ol) is used similarly to codeine, to treat moderate to moderate/severe pain. It is a synthetic analog of the phenanthrene alkaloid codeine and, as such, is an opioid and also a pro-drug (codeine is metabolized to morphine, tramadol is converted to O-desmethyl tramadol and sweating. MDMA is not clearly a stimulant, although it has, in common with amphetamine drugs, a capacity to increase blood pressure and heart rate. MDMA does produce some perceptual changes in the form of increased sensitivity to light, difficulty in focusing, and blurred vision in some users. Its mechanism of action is thought to be via release of the neurotransmitter serotonin. MDMA may also release dopamine, although the general opinion is that it is a secondary effect of the drug (Nichols and Oberlander, 1990). The most pervasive effect of MDMA, occurring in virtually all people who take a reasonable dose of the drug, was to produce a clenching of the jaws.

Alcohol (ETOH)

Alcohol Test is intended for use to detect the presence of alcohol in urine greater than 0.04%. Alcohol intoxication can lead to loss of alertness, coma, death and as well as birth defects. The BAC at which a person is considered to be intoxicated varies by jurisdiction. The United States Department of Transportation established a BAC of 0.02% (0.02 g/dL) as the cut-off level at which an individual is considered positive for the presence of alcohol. Since the urine alcohol concentration is normally higher than that in saliva and blood, the cutoff concentration for alcohol in urine was set at 0.04%. Normally, it will take at least 30 minutes for the alcohol to be detected in saliva, blood and urine after drinking.

PRINCIPLE

The SAFElife® T-Cup Multi-Drug Urine Test Cup is a competitive immunoassay that is used to screen for the presence of drugs of abuse in urine. It is a chromatographic absorbent device in which drugs in a sample competitively combine to a limited number of drug monoclonal antibody (mouse) conjugate binding sites. When the absorbent and is immersed into urine specimen, the urine is absorbed into the device by capillary action, mixes with the respective drug monoclonal antibody conjugate, and flows across the pre-coated membrane. When sample drug levels are zero or below the target cutoff (the detection sensitivity of the test), respective drug monoclonal antibody conjugate binds to the respective drug-protein conjugate immobilized in the Test Region (T) of the device. This produces a colored band in the Test Region (T) that, regardless of its intensity, indicates a negative result.

When sample drug levels are at or above the target cutoff, the free drug in the sample binds to the respective drug monoclonal antibody conjugate preventing the respective drug monoclonal antibody conjugate from binding to the respective drug-protein conjugate immobilized in the Test Region (T) of the device. This prevents the development of a distinct colored band in the Test Region (T), indicating a potentially positive result.

To serve as a procedure control, a colored band will appear at the Control Region (C), where the Goat anti mouse

THC (50)	Viewer C	+	0	0	2	20	20	100% (91.2% - 100%)	
	Viewer B	-	10	13	15	0	0	95% (83.5% - 98.6%)	
	Viewer A	+	0	0	1	18	22	100% (84.5% - 100%)	
	Viewer B	+	0	0	1	18	22	100% (84.5% - 100%)	
	Viewer C	-	10	12	17	0	0	97.5% (82% - 100%)	
	Viewer A	+	0	0	2	19	21	100% (84.5% - 100%)	
TRA (100)	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer A	+	0	0	1	18	20	95% (84.5% - 100%)	
	Viewer C	-	10	20	9	2	0	97.5% (82% - 100%)	
	Viewer A	+	0	0	2	19	21	100% (84.5% - 100%)	
	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer C	-	10	20	9	1	18	20	95% (84.5% - 100%)
TRA (200)	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer A	+	0	0	2	19	21	100% (84.5% - 100%)	
	Viewer C	-	10	20	9	2	0	97.5% (82% - 100%)	
	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer A	+	0	0	1	18	20	95% (84.5% - 100%)	
	Viewer C	-	10	20	9	1	18	20	95% (84.5% - 100%)
TRA (1000)	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer A	+	0	0	1	18	20	95% (84.5% - 100%)	
	Viewer C	-	10	20	9	1	18	20	95% (84.5% - 100%)
	Viewer B	-	10	20	8	0	0	95% (79.5% - 100%)	
	Viewer A	+	0	0	1	18	20	95% (84.5% - 100%)	
	Viewer C	-	10	20	9	1	18	20	95% (84.5% - 100%)

Precision and Sensitivity

To investigate the precision and sensitivity each drug sample was analyzed at the following concentrations: cutoff -100%, cutoff -75%, cutoff -50%, cutoff -25%, cutoff -10%, cutoff -50%, cutoff -75% and the cutoff +100%. All concentrations were confirmed with GC-MS. The study was performed 2 runs /day and lasted 25 days using three different lots of the corresponding drug test. Totally 3 operators participated in the study of the corresponding drug test. Each of the 3 operators tests 2 aliquots at each concentration for each lot per day (2 runs/day), for a total of 50 determinations per concentration per lot of the corresponding drug test.

Drug Test	Approximate Concentration of Sample (ng/mL)	Number of Determinations per Lot	Results			
			Negative		Positive	
6-MAM	0	50	50/0	50/0	50/0	50/0
	2.5	50	50/0	50/0	50/0	50/0
	7.5	50	47/3	48/2	47/3	
	10	50	4/46	5/45	6/44	
	12.5	50	3/47	2/48	2/48	
	15	50	0/50	0/50	0/50	
	17.5	50	0/50	0/50	0/50	
	20	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	7.5	50	50/0	50/0	50/0	
AMP (300)	150	50	50/0	50/0	50/0	
	225	50	50/0	50/0	50/0	
	300	50	5/45	5/45	4/46	
	375	50	0/50	0/50	0/50	
	450	50	0/50	0/50	0/50	
	525	50	0/50	0/50	0/50	
	600	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	125	50	50/0	50/0	50/0	
	250	50	50/0	50/0	50/0	
AMP (500)	375	50	50/0	50/0	50/0	
	500	50	6/44	7/43	6/44	
	625	50	0/50	0/50	0/50	
	750	50	0/50	0/50	0/50	
	875	50	0/50	0/50	0/50	
	1000	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	250	50	50/0	50/0	50/0	
	500	50	50/0	50/0	50/0	
	750	50	50/0	50/0	50/0	
AMP (1000)	1000	50	5/45	6/44	6/44	
	1250	50	0/50	0/50	0/50	
	1500	50	0/50	0/50	0/50	
	1750	50	0/50	0/50	0/50	
	2000	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	75	50	50/0	50/0	50/0	
	150	50	50/0	50/0	50/0	
	225	50	50/0	50/0	50/0	
	300	50	5/45	5/45	6/44	
BAR	375	50	0/50	0/50	0/50	
	450	50	0/50	0/50	0/50	
	525	50	0/50	0/50	0/50	
	600	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	1.25	50	50/0	50/0	50/0	
	2.5	50	50/0	50/0	50/0	
	3.75	50	50/0	50/0	50/0	
	5.0	50	5/45	5/45	6/44	
	6.25	50	0/50	0/50	0/50	
BUP (5)	7.5	50	0/50	0/50	0/50	
	8.75	50	0/50	0/50	0/50	
	10	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
	2.5	50	50/0	50/0	50/0	
	5.0	50	5/45	5/45	6/44	
	6.25	50	0/50	0/50	0/50	
	7.5	50	0/50	0/50	0/50	
	8.75	50	0/50	0/50	0/50	
	10	50	0/50	0/50	0/50	
BUP (10)	0	50	50/0	50/0	50/0	
	2.5	50	50/0	50/0	50/0	
	5.0	50	50/0	50/0	50/0	
	7.5	50	50/0	50/0	50/0	
	10.0	50	5/45	5/45	6/44	
	12.5	50	0/50	0/50	0/50	
	15.0	50	0/50	0/50	0/50	
	17.5	50	0/50	0/50	0/50	
	20.0	50	0/50	0/50	0/50	
	0	50	50/0	50/0	50/0	
BZO (100)	25	50	50/0	50/0	50/0	
	50	50	50/0	50/0	50/0	
	75	50	46/4	46/4	47/3	
	100	50	4/46	4/46	4/46	
	125	50	3/47	3/47	2/48	