RapidTech Drug Screen MAMP

INTENDED USE

Genix RapidTech Drug Screen AMP test is an immunochromatography based one step test designed for the qualitative determination of methamphetamine metabolite in human urine specimens at the cut-off concentration of 1,000 ng/ml. For in-vitro diagnostic use only.

SUMMARY AND EXPLANATION

Methamphetamine is the most popular synthetic derivative of the amphetamines. It is a potent sympathomimetic agent with therapeutic applications. Acute large doses lead to enhanced stimulation of the central nervous system and induce euphoria, alertness, reduced appetite, and a sense of increased energy and power. More acute response produce anxiety, paranoia, psychotic behavior, and cardiac dysrhythmias. Methamphetamine is excreted in the urine as amphetamine and oxidized and deaminated derivatives. However, 10-40% of methamphetamine is generally detectable in the urine for 3 to 5 days after use.

PRINCIPLE

Each test strip is based on the principle of specific immunochemical reaction between antibodies and antigen to analyze particular compound in human urine specimen. The assay relies on the competition for binding antibody. When drug is present in the urine specimen, it competes with drug conjugate for the limited amount of antibody-dye conjugate. When the amount of drug is equal or more than the cut-off, it will prevent the binding of drug conjugate to the antibody. Therefore, a positive urine specimen will not show a coloured band on the test line zone, indicating a positive result, while the presence of a coloured band indicates a negative result.

A control line is present in the test window to work as procedural control. This coloured band should always appear on the control line zone if the test device is stored in good condition and the test is performed appropriately.

REAGENT AND MATERIALS SUPPLIED

- 1. Test strips.
- 2. Instruction for use.

MATERIAL REQUIRED BUT NOT SUPPLIED

- 1. Urine collection container.
- 2. Timer or clock.

STORAGE AND STABILITY

The test strip should be stored refrigerated or at room temperature (2° to 30° C) in the sealed pouch. The test strip is effective until the expiration date printed on the pouch.

PRECAUTIONS

- 1. For in vitro diagnostic and forensic use only.
- 2. Do not use the product beyond the expiration date.
- 3. Handle all specimens as potentially infectious.
- 4. Do not open foil pouch until it is ready to be tested.
- 5. Use a new urine specimen cup for each sample to avoid cross contamination.

SPECIMEN COLLECTION AND PREPARATION

Fresh urine does not require any special handling or pretreatment. If the assay is not performed immediately, urine specimen may be refrigerated at 2° to 8°C up to 7 days or frozen. Specimens should be brought to room temperature (18° to 25°C) before testing. Urine specimens exhibiting a large amount of precipitate or turbidity should be centrifuged or allowed to settle before testing.

QUALITY CONTROL

- 1. The control band is an internal reagent and procedural control. It will appear if the test has been performed correctly and the reagents are reactive.
- 2. Control standards can be used to validate reagent performance and establish test reliability. Controls that are not provided with this test are commercially available.

TEST PROCEDURE

1. Bring all materials and specimens to room temperature (18° to 25°C).

- 2. Open the sealed foil pouch and remove the test strip.
- 3. Label the strip with patient identification.
- 4. Dip the strip into the urine specimen with the arrow pointing toward the sample. The sample level should not be higher than the arrow pointed maximum line.
- 5. Hold the strip in the urine until a reddish colour appears at the test area (approximately 15 seconds).
- 6. Withdraw the strip and place it face up on a clean, non-absorbent surface or leave the strip in urine if the urine level is not higher than arrow pointed maximum line.
- 7. Read the result at 5 minutes.

Do not interpret the result after 10 minutes.

INTERPRETATION OF RESULTS

Negative:

Two coloured bands form. The appearance of two coloured bands, one in test (T) zone and the other in control (C) zone, indicates negative results. The negative result does not indicate the absence of drug in the specimen, it only indicates that the drug level in the specimen is less than cut-off level.

Positive:

One coloured band forms. One coloured band appears in control (C) zone. No coloured band is found in test (T) zone. This is an indication that the drug level in the specimen is above the cut-off level.

Invalid:

If there is no coloured band in control (C) zone, the test result is invalid. Repeat the test with a fresh test device.

Note: A very faint coloured band in test (T) zone indicates that the amount of drug in the sample is near the cut-off level. These specimens and any positive samples should be confirmed by and alternate method such as GC/MS.



EXPECTED RESULTS

The RapidTech Drug Screen MAMP Test is a qualitative assay. It identifies methamphetamine metabolite in human urine at its cut-off concentration or higher. The concentration of the drug cannot be determined by this assay. The test is intended to distinguish negative result from presumptive positive result. All positive results must be confirmed using an alternate method, preferably GC/MS.

PERFORMANCE CHARACTERISTICS Sensitivity

The following compounds were tests at the cut-off concentrations indicated. The cross reaction (calculated in percentage) is indicated in the following table.

Compounds tested	Cut-off	Cross Reactivity
	(ng/ml)	(%)
(+) Methamphetamine	1,000	100
(±) 3,4 MDMA	1,000	100

Interference testing

The following substances did not interfere with RapidTech Drug Screen MAMP Test at the concentrations tested.

Glucose	2000 mg/dl
Human albumin	2000 mg/dl
Human hemoglobin	10 mg/dl
Urea	4000 mg/dl
Uric acid	10 mg/dl

Specificity

The following compounds show no cross-reactivity at concentration up to 100 μ g/ml unless specified.

Acetaminophen Acetylsalicylic acid Amitriptvline Arterenol Ascorbic acid Caffeine Chloroquine Cortisone Dextromethorphan Digoxin Ecgonine Ephedrine Gentisic Histamine Homatrophine Ibuprofen Ketamine Meperidine Methaqualone Neomycin Perphenazine Phenylethylamine- α Promethazine Ouinine antidine Tetracvcline Theophyline Trifluoperazine Tyramine

4-Acetamidophenol Amikacin Amobarbital Aspartame Atrophine Camphor Chlopheniramine Deoxyephedrine Digitoxin Diphenhydramine Ecgonine methyl ester Epinephrine Guaiacol glycer ester Hydrochlorothiazide Imipramine Isoproterenol Lidocaine Methadone Methylphenidate Niacinamide Penicillin G Phenylpropanolamine Pseudoephedrine Salicylic acid Tetrahvdrozoline Thioridazine Tryptophan

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Revision A: January 2000.

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