

# DrugCheck® Dip Drug Test

 $\in$ 

The **DrugCheck® Dip Drug Test** is an immunochromatographic assay for the qualitative detection of Ethyl Glucuronide in human urine at a cutoff concentration indicated in the table below.

The test may yield preliminary positive results when prescription drugs are ingested at prescribed doses. It is not intended to distinguish between prescription use and abuse of any drug. There are no uniformly recognized cutoff concentration levels for any drug in urine. The test provides only preliminary test results. A more specific alternative 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 exercised with any drug of abuse test result, particularly when the preliminary result is positive.

For in vitro diagnostic use and for professional testing use only.

### WHAT IS DRUGCHECK® DIP DRUG TEST?

The **DrugCheck® Dip Drug Test** is a rapid test for qualitative detection of Ethyl Glucuronide in human urine. The **DrugCheck® Dip Drug Test** yields a positive result when drug and/or its metabolite in urine is at or exceeds its cutoff concentration.

### WHAT IS THE CUT-OFF VALUE?

	Drug Test	Drug (Identifier)	Cut-off
I	Ethyl Glucuronide (ETG)	Ethyl Glucuronide	500 ng/mL

### **PRINCIPLE**

The **DrugCheck® Dip Drug Test** is an immunoassay. During testing, a urine specimen migrates upward on the test strip. A drug-positive urine specimen will not generate a colored line in the specific test line region of the strip, while a drug-negative urine specimen will generate a line in the test line region. A colored line will always appear at the control line region, indicating that proper volume of specimen has been added.

The test contains a membrane strip coated with drug-protein conjugates (purified bovine albumin) on the test line, a goat polyclonal antibody against gold-protein conjugate at the control line, and a dye pad which contains colloidal gold particles coated with mouse monoclonal antibody specific to individual drug on the list indicated in the table above.

### WARNINGS AND PRECAUTIONS

- 1. For in vitro diagnostic use and for professional testing use only.
- 2. For external use only.
- 3. For single use. Discard after first use.
- 4. Do not use the test if the pouch is punctured or not well sealed.
- 5. Do not use after expiration date
- 6. Keep out of the reach of children.
- The used dip test and urine specimen should be discarded according to federal, state and local regulations.

### CONTENT OF THE PACKAGE

Included in package:

- User Instruction

- Dip Test (inside foil pouch)

Not included in package:

- Watch, Timer or Clock

- Collection Cup

# STORAGE AND STABILITY

Store as packaged in the sealed pouch at 4°C - 30°C. The test is stable through the expiration date printed on the sealed pouch. The dip test must remain in the sealed pouch until use. Keep away from direct sunlight, moisture and heat. DO NOT FREEZE. Do not use beyond the expiration date.

## WHEN TO COLLECT URINE FOR THE TEST?

You can use urine from any time of the day. The minimum detection time varies for different drugs.

### **HOW TO COLLECT URINE?**

- 1. When you are ready to begin, remove the dip test from the sealed foil pouch.
- Notice the colored tape on each strip indicates the name of the drug you are testing for.
- 3. Fill the collection cup with a fresh urine sample. Do not over-fill.

### HOW TO DO THE TEST?

### For Single Dip Strip

- Insert the test strip into the urine sample for 10 to 15 seconds. DO NOT let the
  urine sample touch the conjugate pad on the strip, this could cause inconclusive
  drug test results. Place the test on a flat surface.
- Wait for 5 minutes (start timing immediately after dip is taken out of the urine sample). Read result at 5 minutes. DO NOT READ RESULT AFTER 5 MINUTES.

### For Single Dip Cassette

- Remove the cap from the dip test. Insert the exposed test strip into the urine sample for 10 to 15 seconds. DO NOT let the urine sample touch the plastic device, this could cause inconclusive drug test results.
- Insert the cap firmly back onto the dip test and lay it on a flat surface.
- Wait for 5 minutes (start timing immediately after dip test is taken out of the urine sample). Read result at 5 minutes. DO NOT READ RESULT AFTER 5 MINUTES.

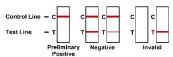
Note: Results after more than 5 minutes may not be accurate and should not be read

### READING THE RESULTS

<u>Preliminary Positive (+)</u>: If a line appears in the C - Control area but NO line appears in the T - Test area, then it indicates a Preliminary Positive result for the corresponding drug.

Negative (-): If a line appears in both the C - Control and T - Test area, then it indicates a Negative result for the corresponding drug regardless of how dark or light the line may appear.

<u>Invalid</u>: If at 5 minutes, NO line appears in the C - Control area, then the results are Invalid. In such case, retest with a new dip test.



**Note:** Each test strip needs to be looked at individually. Each line may vary in color and darkness or the rate at which the line appears. (DO NOT compare lines within the same test strip or between different test strips).

A positive test result does not always mean a person took illegal drugs and a negative test result does not always mean a person did not take illegal drugs. There are a number of factors that influence the reliability of drug tests. Certain drugs of abuse tests are more accurate than others

**IMPORTANT:** The result you obtained is called preliminary for a reason. The sample must be tested by a laboratory in order to determine if a drug of abuse is actually present.

### WHAT IS A FALSE POSITIVE TEST?

The definition of a false positive test would be an instance where the test result from the <code>DrugCheck</code> <code>Dip Drug Test</code> is positive, even though the initial target drug is not present in the sample. The most common causes of a false positive test are cross reactants. Certain foods and medicines, diet plan drugs and nutritional supplements may also cause a false positive test result with this product.

### WHAT IS A FALSE NEGATIVE TEST?

The definition of a false negative test is that the initial target drug is present but is not detected by the **DrugCheck® Dip Drug Test**. If the sample is diluted, or if the sample is tainted or contaminated with a substance this could cause false negative results

# TEST LIMITATIONS

- The DrugCheck® Dip Drug Test provides only a qualitative, preliminary analytical result. A secondary analytical method must be used to obtain a confirmed result. Gas chromatography/mass spectrometry (GC/MS) is the preferred confirmatory method.
- 2. There is a possibility that interfering substances in the urine specimen may cause erroneous results.
- 3. Substances, such as bleach and/or alum, in urine specimens may produce erroneous results.
- A positive result does not indicate intoxication, the concentration of drug in the urine, or the route of drug administration.
- A negative result may not necessarily indicate drug-free urine. Negative results can be obtained when drug is present but below the cutoff level of the test.
- 6. Test does not distinguish between drugs of abuse and certain medications.
- 7. A positive test result may be obtained from certain foods or food supplements.

### QUALITY CONTROL

If you work in a laboratory, you should perform quality control testing and you should read this section.

A procedural control is included in the test. A color line appearing in the control 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. Quality control testing should be done with each new lot and each new shipment. It should be done every thirty days to check storage. Please contact our Technical Support at 507-526-3951 for controls that work with the dip test.

### PERFORMANCE CHARACTERISTICS

### Accuracy

In the comparison study, the <code>DrugCheck®</code> <code>Dip Drug Test</code> was compared to a GC/MS reference method to determine its accuracy. Clinical urine samples were collected for Ethyl Glucuronide. Clinical specimens were quantified by GC/MS analysis before testing. The following results are tabulated from these clinical studies:

Analyte	Positive	Negative
Negative Samples	0	70
Near Cut-off Negative Samples [between 50% of cut-off and cut-off]	0	70
Near Cut-off Positive Samples [between cutoff and 150% of cut-off]	70	0
Positive Samples [>150% of cut-off]	70	0
Agreement with GC/MS	>99%	>99%

Overall Agreement with GC/MS is >99%

### Reproducibility

Reproducibility studies were carried out using commercially available stock solutions of the drug analyte listed. Dilutions were made from the stock solution of Ethyl Glucuronide to the concentrations specified in the following table. The results are listed in the following table.

Approximate Concentration of Sample (ng/mL)	Number of Determinations	Result	Precision
0	40	40 negative	>99%
250	40	40 negative	>99%
750	40	40 positive	>99%

### **Analytical Sensitivity**

A drug-free urine pool was spiked with drugs at concentrations listed. The results are summarized below.

		ET	rg .
Drug concentration Cut-off Range	n	-	+
0% Cut-off	30	30	0
-50% Cut-off	30	30	0
-25% Cut-off	30	30	0
Cut-off	30	3	27
+25% Cut-off	30	1	29
+50% Cut-off	30	0	30
2x Cut-off	30	0	30

### **Analytical Specificity**

The following table lists the concentration of compounds (ng/mL) that were detected positive in urine by the **DrugCheck® Dip Drug Test** at a read time of 5 minutes.

Ethyl Glucuronide (ETG)	Result	
(Ethyl-β-D-glucuronide, Cutoff = 500 ng/mL)	Positive at 500 ng/mL	

### **EFFECT OF URINARY SPECIFIC GRAVITY**

Urine samples of normal, high, and low specific gravity ranges (1.000-1.035) were spiked with drugs at 25% below and 25% above cut-off levels respectively. The <code>DrugCheck®Dip Drug Test</code> was tested using twelve drug-free urine and spiked urine samples. The results demonstrate that varying ranges of urinary specific gravity do not affect the test results.

### **EFFECT OF URINARY PH**

The pH of an aliquot of negative urine pool was adjusted to pH ranges of 4.0 - 9.0 and spiked with drugs at 25% below and 25% above cut-off levels. The spiked, pH-adjusted urine was tested with the **DrugCheck® Dip Drug Test**. The results demonstrate that varying ranges of pH do not interfere with the performance of the test.

### INTERFERENCE

A study was conducted to determine the cross-reactivity of the test with compounds in either drug-free urine or drug positive urine containing Ethyl Glucuronide. The following compounds show no cross-reactivity when tested with the  $\mathbf{DrugCheck}^{\otimes}$   $\mathbf{Dip}$   $\mathbf{DrugTest}$  at concentrations of 100  $\mu g/m$ L.

Acebutolol Hydrochloride	Gentisic acid	Pentazocine		
cepromazine-d6	D-Glucuronic acid	Perphenazine		
hydrochloride	Glutethimide	Penicillin G Sodium salt		
Acetaminophen	Guaifenesin	Phenelzine sulfate salt		
N-Acetylprocainamide	Hemoglobin porcine	Phenobarbital		
Acetophenetidin	Heroin hydrochloride	Phentermine HCL		
Alprazolam	Hippuric Acid	Phenylethylamine		
Alphenal	Hydralazine hydrochloride	L-phenylephrine		
Amoxicillin	Hydromorphone	Phenylpropanolamine		
Ampicillin	Hydrocodone	hydrochloride		
Amitriptyline	α-Hydroxyhippuric acid	Prednisolone		
Hydrochloride Tablets	21-Hydroxy progesterone	Prednisone Acetate		
S(+)Amphetamine	p-Hydroxymeth-	Tablets		
R(-)-Amphetamine	amphetamine	Procaine HCL		
Amobarbital	Hydrocortisone	Promazine hydrochloride		
(±)Amphetamine	Hydrochlorothiazide	Promethazine		
R-(-)-Apomorphine	-4-Hydroxyamphetamine	Propoxyphene,d-		
Hydrochloride	HCL	Propranolol Hydrochloride		
Aprobarbital	Ibuprofen	Pseudoephedrine		
Aspirin	Imipramine	Phendimetrazine		
Aspartame	Iprazid	Phenytoin		
L-Ascorbic Acid	Isoxsuprine hydrochloride	Quinine		
Atropine	Isoproterenol	Quinidine		
6-Acetylmorphine	Hydrochloride Injection	Quinacrine		
Acetylsalicylic acid	Ketamine hydrochloride	Ranitidine Hydrochloride		
Benzphetamine	Ketoprofen	Tablets		
Benzilic acid	Emetine dihydrochloride	Nortriptyline Hydrochloride		
Benzoylecgonine	hydrate	Salicylic Acid		
SS Benzoic Acid	Ephedrine-(+/-)	Secobarbital		
Bilirubin, Mixed Isomers	hydrochloride ´	Serotonin		
Brompheniramine maleate	(-)-Ephedrine HCL	Noroxymorphone HCL		
Buprenorphine	[1R,2S] (-) Ephedrine	Nylidrin hydrochloride		
Buspirone hydrochloride	Erythromycin	Norfentanyl		
Butalbital	Eserine	(±)-Octopamine HCL		
Butabarbital	Estazolam	Oxalic Acid		
Cannabidiol	β-Estradiol	Oxolinic Acid		

### BIBLIOGRAPHY OF SUGGESTED READING

- Stewart DJ, Inaba T, Lucassen M, Kalow W. Clin. Pharmacol. Ther. April 1979; 25 ed: 464, 264-8.
- Ambre J. J. Anal. Toxicol. 1985: 9:241.
- Hawks RL, CN Chiang. Urine Testing for Drugs of Abuse. National Institute for Drug Abuse (NIDA), Research Monograph 73, 1986.

### ADDITIONAL INFORMATION AND REFERENCES

The following list of organizations may be helpful to you for counseling support and resources. These groups also have an Internet address, which can be accessed for additional information.

National Clearinghouse for Alcohol and Drug Information www.health.org 1-800-729-6686

Center for Substance Abuse Treatment www.health.org 1-800-662-HELP

The National Council on Alcoholism and Drug Dependence <a href="www.ncadd.org">www.ncadd.org</a> 1-800-NCA-CALL

American Council for Drug Education (ACDE) www.acde.org 1-800-488-DRUG

### SYMBOLS

YMBOLS			
$\subseteq$	Use-By Date	<b></b>	Manufacturer
REF	Catalogue Number	1	Temperature Limit
	Do Not Use if Package is Damaged	8	Do Not Re-Use
漆	Keep Away from Sunlight	Ţį.	Consult Instructions for Use
<del>*</del>	Keep Dry	<u> </u>	Caution
LOT	Batch Code	IVD	In Vitro Diagnostic Medical Device
EC REP	Authorized representative in the European Community	Σ	Contains Sufficient for <n> Tests</n>



CareHeatlh America Corporation dba Express Diagnostics Int'l, Inc. 1550 Industrial Drive Blue Earth, MN 56013



CEpartner4U Esdoornlaan 13 3951 DB Maarn The Netherlands



DC4101-INT, Rev B 09/2021