

### **AMPHETAMINE (AMP) DRUG SCREEN**

Accuracy: The performance of the the iCup amphetamine drug screen was compared to the laboratory initial screen and a leading commercially available amphetamine rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between -25% or +25% of the cut-off concentration of 1,000 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 500 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	142	0	142
Test Device	Negative	9	149	158
Total Results		151	149	300

Relative Sensitivity: 94% Relative Specificity: >99% Accuracy: 97%

**Note:** The FDA requires on-site test devices to be accurate within +25% of the drug screen's cut-off level. The *i*Cup detected 100% of the GC/MS positive specimens within 25% of the screen's 1000ng cut-off.

The only GC/MS positive specimens not identified as positive by the *i*Cup were close to or below the *i*Cup 's 1000 ng/ml cutoff. Specimens whose results differed between GC/MS and the *i*Cup contained the following concentration levels in ng/ml: (781,862, 878, 898, 898, 1000, 1032, 1137, and 1204).

### **METHAMPHETAMINE (mAMP) DRUG SCREEN**

Accuracy: The performance of the the *i*Cup methamphetamine drug screen was compared to the laboratory initial screen and a leading commercially available amphetamine rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between –25% or +25% of the cut-off concentration of 1,000 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 500 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	145	0	145
Test Device	Negative	1	154	155
Total Re	sults	146	154	300

Relative Sensitivity: >99% Relative Specificity: >99% Accuracy: >99%

**Note:** The *i*Cup detected 100% of the GC/MS positive specimens at or above the screen's 1000ng cut-off.

The only GC/MS positive specimen not identified as positive by the <code>iCup</code> was below the <code>iCup</code> 's 1000 ng/ml cut-off. The discrepant specimen contained d-methamphetamine at a concentration of 954 ng/ml.

#### MARIJUANA (THC) DRUG SCREEN

Accuracy: The performance of the the *i*Cup amphetamine drug screen was compared to the laboratory initial screen and a leading commercially available marijuana rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between -25% or +25% of the cut-off concentration of 50 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 15 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	143	0	143
Test Device	Negative	7	150	157
Total Re	sults	150	150	300

Relative Sensitivity: 95% Relative Specificity: >99% Accuracy: 98%

**Note:** The FDA requires on-site test devices to be accurate within 25% of the drug screen's cut-off level. The *i*Cup detected all but two of the GC/MS positive specimens within +25% of the screen's 50ng cut-off.

The only GC/MS positive specimens not identified as positive by the *i*Cup were close to or below the *i*Cup 's 50 ng/ml cut-off. Specimens whose results differed between the GC/MS and the *i*Cup contained the following concentration levels in ng/ml: (15, 15, 16, 32, 51, 73, and 79)

### **COCAINE DRUG SCREEN**

Accuracy: The performance of the the <code>iCup</code> cocaine drug screen was compared to the laboratory initial screen and a leading commercially available amphetamine rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between -25% or +25% of the cut-off concentration of 300 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 150 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	136	0	136
Test Device	Negative	13	151	164
Total Results		149	151	300

Relative Sensitivity: 91% Relative Specificity: >99% Accuracy: 96%

**Note:** The FDA requires on-site test devices to be accurate within 25% of the drug screen's cut-off level. The *i*Cup detected all but one of the GC/MS positive specimens within 25% of the screen's 300ng cut-off.

The only GC/MS positive specimens not identified as positive by the *i*Cup were close to or below the *i*Cup 's 300 ng/ml cut-off. Specimens whose results differed between GC/MS and the *i*Cup contained the following concentration levels in ng/ml: (153, 158, 188, 202, 228, 233, 243, 254, 265, 270, 298, 358 and 381)

### **OPIATES-2000 (OPI) DRUG SCREEN**

Accuracy: The performance of the the <code>iCup</code> opiates drug screen was compared to the laboratory initial screen and a leading commercially available opiates rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between –25% or +25% of the cut-off concentration of 2,000 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 2000 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	134	16	150
Test Device	Negative	0	150	150
Total Results		134	166	300

Relative Sensitivity: >99% Relative Specificity: 90% Accuracy: 95%

**Note:** The *i*Cup correctly identified 100% of the specimens determined to be positive by GC/MS.

The specimens identified as positive by the <code>iCup</code> but below 2000 ng/ml by GC/MS were all at or above 1500 ng/ml. Therefore the <code>iCup</code> performed at >99% accuracy within +/-25% of the screens 2000 ng/ml cut-off.

#### **PCP DRUG SCREEN**

Accuracy: The performance of the the *i*Cup PCP drug screen was compared to the laboratory initial screen and a leading commercially available PCP rapid test. Testing was performed on 212 clinical specimens. At least ten percent of the specimens evaluated were between –25% or +25% of the cut-off concentration of 25 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 25 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
<i>i</i> Cup Test Device	Positive	50	5	55
l est Device	Negative	0	157	157
Total Res	sults	50	162	212

**Note:** The *i*Cup correctly identified 100% of the specimens determined to be positive by GC/MS.

The specimens identified as positive by the *i*Cup but below 25 ng/ml by GC/MS were all at or above 15 ng/ml. Therefore; the *i*Cup detected 5 positive specimens that were below the 25 ng/ml GC/MS cut-off, but there were no false-positive results.

Relative Sensitivity: >99% Relative Specificity: 97% Accuracy: 98%

### **BARBITURATES (BAR) DRUG SCREEN**

Accuracy: The performance of the the <code>iCup</code> barbiturates drug screen was compared to the laboratory initial screen and a leading commercially available barbiturates rapid test. Testing was performed on 292 clinical specimens. At least ten percent of the specimens evaluated were between -25% or +25% of the cut-off concentration of 300 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 300 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	122	4	126
Test Device	Negative	10	156	166
Total Res	sults	132	160	292

Relative Sensitivity: 92% Relative Specificity: 98% Accuracy: 95%

**Note:** The FDA requires on-site test devices to be accurate within 25% of the drug screen's cut-off level. The *i*Cup's accuracy within 25% of the screen's 300ng cut-off was 97%

The specimens identified as positive by the *i*Cup but below 300 ng/ml by GC/MS were all at or above 218 ng/ml. Therefore; the *i*Cup detected 4 positive specimens that were below the 300 ng/ml GC/MS cut-off, but there were no false-positive results.

### **BENZODIAZEPINE (BZO) DRUG SCREEN**

Accuracy: The performance of the the *i*Cup benzodiazepines drug screen was compared to the laboratory initial screen and a leading commercially available benzodiazepines rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between –25% or +25% of the cut-off concentration of 300 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 300 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
<i>i</i> Cup Test Device	Positive	129	7	136
l est Device	Negative	5	159	164
Total Results		134	166	300

Relative Sensitivity: 96% Relative Specificity: 96% Accuracy: 96%

**Note:** The FDA requires on-site test devices to be accurate within 25% of the drug screen's cut-off level. The *i*Cup detected all but one of the GC/MS positive specimens within +25% of the screen's 3000ng cut-off.

The only GC/MS positive specimens not identified as positive by the *i*Cup were close to or below the *i*Cup 's 300 ng/ml cut-off. Specimens whose results differed between the GC/MS and the *i*Cup contained the following concentration levels in ng/ml: (323,326, 334, 335, and 540)

The iCup

detected 5 positive specimens that were below the 300 ng/ml GC/MS cut-off, but there were no false-positive results.

#### **METHADONE (MTD) DRUG SCREEN**

Accuracy: The performance of the the <code>iCup</code> methadone drug screen was compared to the laboratory initial screen and a leading commercially available methadone rapid test. Testing was performed on 300 clinical specimens. At least ten percent of the specimens evaluated were between -25% or +25% of the cut-off concentration of 300 ng/ml. Presumptive positive results were confirmed by GC/MS.

When compared to GC/MS at 200 ng/mL, the results are provided in the table below:

Method		GC/MS		Total
	Results	Positive	Negative	Results
iCup	Positive	132	0	132
Test Device	Negative	1	167	168
Total Results		133	167	300

**Note:** The *i*Cup detected >99% of the GC/MS positive specimens at +/- 25% of the screens 300 ng/ml cut-off.

The only GC/MS positive specimen not identified as positive by the *i*Cup contained methadone at a concentration of 534 ng/ml.

The *i*Cup did not detect any specimens as positive that did not confirm by GC/MS at 200 ng/ml.

Relative Sensitivity: >99% Relative Specificity: >99% Accuracy: >99%

### TRICYCLIC ANTIDEPRESSANT (TCA) DRUG SCREEN

Accuracy: The performance of the the *i*Cup TCA drug screen was compared to the laboratory initial screen and a leading commercially available TCA rapid test. Testing was performed on 200 clinical specimens. At least ten percent of the specimens evaluated were between –25% or +25% of the cut-off concentration of 1,000 ng/ml. Presumptive positive results were confirmed by HPLC.

When compared to HPLC at 300 ng/mL, the results are provided in the table below:

Method		HPLC		Total
	Results	Positive	Negative	Results
<i>i</i> Cup Test Device	Positive	60	0	60
l est Device	Negative	0	140	140
Total Results		60	140	200

**Note:** The *i*Cup correctly identified 100% of the specimens determined to be positive by GC/MS, and all specimens identified as positive by the *i*Cup were confirmed positive by HPLC at or above a 300 ng/ml concentration level.

There is no DHHS specified confirmation level for TCA

Relative Sensitivity: >99% Relative Specificity: >99% Accuracy: >99%



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