



## *Manufacturer's Information*

# **Test No. 15-502: Drug Analysis**

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Each sample pack consisted of two items. Item 1 contained 500 mg of powder consisting of approximately 70% Carisoprodol; the remainder of the sample consisted of tablet binders/fillers. Item 2 contained 200 mg of powder consisting of acetaminophen. Carisoprodol is a Schedule IV controlled substance in the United States.

### SAMPLE PREPARATION-

Tablets containing Carisoprodol were used for Item 1. The tablets were ground into powder and sieved before weighing each sample. The acetaminophen in Item 2 was a pure powder.

ITEM 1 (PREPARATION): Approximately 500 mg of the powder containing Carisoprodol was weighed out and deposited into a glassine bag which was folded and secured with a label. The folded glassine bag was placed into a small zip top bag. The small zip top bag was heat sealed and then placed into a pre-labeled 5 1/2 inch coin envelope.

ITEM 2 (PREPARATION): Approximately 200 mg of the acetaminophen powder was weighed out and deposited into a glassine bag which was folded and secured with a label. The folded glassine bag was placed into a small zip top bag. The small zip top bag was heat sealed and then placed into a pre-labeled 5 1/2 inch coin envelope.

SAMPLE PACK ASSEMBLY: One of each of the Item 1 and Item 2 envelopes was placed into a larger pre-labeled sample pack envelope.

VERIFICATION: The above results were confirmed by predistribution laboratories, who used the following combined list of techniques: Color Tests, FTIR, GC, and GC/MS.