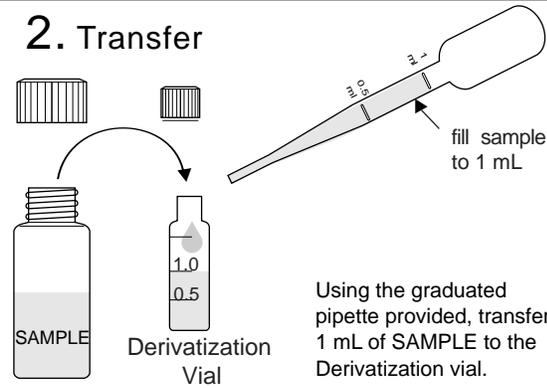


1. Collect Sample/Extract



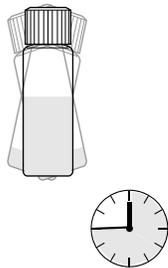
Collect diluted sample/extract.

2. Transfer



Using the graduated pipette provided, transfer 1 mL of SAMPLE to the Derivatization vial.

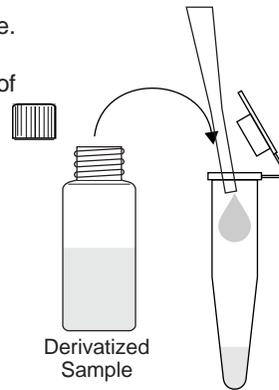
3. Mix & Incubate



Mix for 30 seconds. Incubate in a 45°C heat block for 45 minutes.

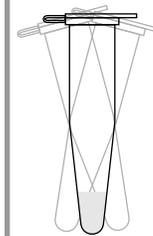
4. Transfer

Squeeze the upper bulb to draw sample into the pipette. The required amount of sample will fit into the stem of the pipette. Any excess will flow into the overflow bulb. Squeeze the upper bulb again to dispense the required amount of sample into conical tube.



(The conical flip-top tube contains dried reagents.)

5. Shake and incubate



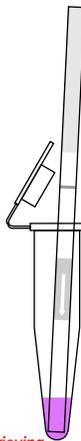
Close the conical flip-top tube and shake for 30 seconds. Check tube to ensure dried reagent dissolves completely (will turn the sample purple).



Incubate for 10 minutes.

6. Test

Insert test strip into conical flip-top tube with arrow pointing down (sample pad down). Incubate for 10 minutes.

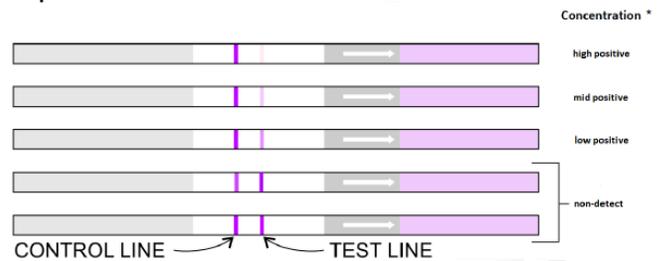


7. Dry

Remove test strip. Lay flat and allow to continue developing for 10 minutes.



8. Interpret



INTERPRET TEST

CONTROL LINE	TEST LINE	INTERPRETATION
NO CONTROL LINE PRESENT	NO TEST LINE PRESENT	INVALID RESULT
CONTROL LINE PRESENT	VERY FAINT TEST LINE OR NO TEST LINE PRESENT	HIGH CONCENTRATION
CONTROL LINE PRESENT	MODERATE INTENSITY TEST LINE PRESENT	LOW TO MODERATE CONCENTRATION

*See appropriate technical bulletin for actual sample concentration ranges in various matrices.