premier platinum

TEST PROCEDURE

How to perform the test



1. Add 500 µL of Sample Diluent to a test tube.



2. Solid Stool: Using applicator stick, add small portion (5 - 6 mm diameter) of throughly mixed stool to tube and emulsify using the applicator stick.

Liquid Stool: Using Transfer pipette, add 100 µL of stool to tube. Rinse pipette several times.



Vortex for 15 seconds.



 Using the transfer pipette, add 100 uL of diluted stool to a test well (PAT).



5. Add 2 drops of Positive Control and 100 µL of Negative Control (Sample Diluent) reagents to designated wells.



6. Add 1 drop Enzyme Conjugate to all wells. Shake firmly for 30 seconds. Seal the plate and incubate at 19-27 C for 1 hour.



- 7. Wash 5 times with 1X Wash Buffer.
 - *See package insert for proper wash procedure



8. Add 2 drops of Substrate Solution I to all wells. Shake firmly for 30 seconds and incubate for 10 minutes at 19-27 C.



9. Add 2 drops of Stop Solution I to each well and firmly shake for 30 seconds.

Interpretation of Results

Negative: Colorless to Faint Yellow Positive: Definite Yellow Color

A faint yellow color must be evaluated spectrophotometrically. If a spectrophotometer is not available, the cutoff must be determined by another method.

Spectrophotometric Single Wavelength (450 NM):

Positive Control= OD 450 ≥ 0.640 Negative Control= OD 450 < 0.140 Positive Test: OD 450 ≥ 0.140 Negative Test: OD 450 < 0.140

Spectrophotometric Dual Wavelength (450/630 NM):

Positive Control= OD 450/630 ≥ 0.600 Negative Control= OD 450/630 < 0.100 Positive Test: OD 450/630 ≥ 0.100 Negative Test: OD 450/630 < 0.100

NOTE: This chart does not contain complete instructions for use. For further information please thoroughly read the package insert.



This illustration is representative of the current Package Insert at the time of publication. Please refer to the most current version of the Package Insert for complete instructions.

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