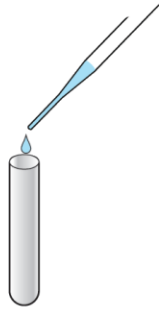


premier®

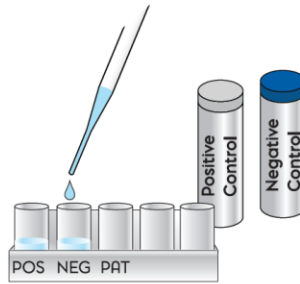
H. pylori

TEST PROCEDURE

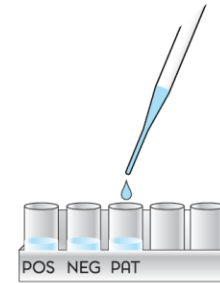
How to perform the test



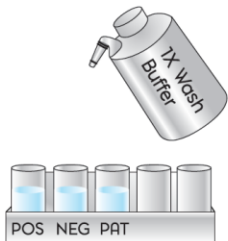
1. Dilute each specimen, positive and negative control 1:50 by mixing 10 µL of each with 0.5 mL Sample Diluent.



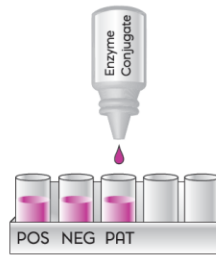
2. Detach microwells needed, place in holder and add 100 µL of each diluted control.



3. Add 100 µL diluted patient specimen to the appropriate well. Incubate at room temperature for 20 minutes.



4. Wash 3 times with 1X Wash Buffer. *See package insert for proper wash procedure.



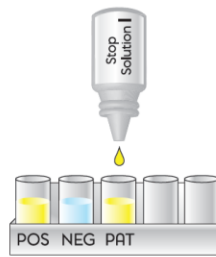
5. Add 2 drops of Enzyme Conjugate to all wells. Incubate at room temperature for 20 minutes.



6. Repeat wash procedure, as in Step 4.



7. Add 2 drops of Substrate I to all wells. Incubate at room temperature for 10 minutes.



8. Add 2 drops of Stop Solution I to all wells and shake firmly for 30 seconds.

9. Read results visually or spectrophotometrically.

Visual

Negative = colorless

Positive = definite yellow color

Spectrophotometric

Single Wavelength (450 nm)

Negative = $OD_{450} < 0.120$

Positive = $OD_{450} \geq 0.120$

Spectrophotometric

Dual Wavelength (450/630 nm)

Negative = $OD_{450/630} < 0.070$

Positive = $OD_{450/630} \geq 0.070$



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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