

Bassett Healthcare Network
Point of Care Testing
Stanbio QuPID Urine hCG Competency

Employee Information

Name and Title (print): _____

Employee #: _____ Location: _____ Date: _____

Skills Review and Evaluation

The operator has demonstrated understanding and proficiency in the following:

	Yes	No
Specimen Collection and Patient Testing - Two patient identifiers must be confirmed prior to testing. - Label test device with two patient identifiers and place on a clean, flat surface, dispense 2 full drops of urine sample into the sample well, set timer for 3 minutes and read results when timer ends at 3 minutes). - Result interpretation: positive results are indicated by a line in both the 'C' area and 'S' area; negative results are indicated by a line in the 'C' area only; invalids results are indicated by the absence of a line in the 'C' area, even when a line is present in the 'S' area). - Result is recorded via Enter/Edit Results function in Epic (POC7), or via MTE depending upon performing location. The POC Urine Pregnancy narrative comment automatically attaches to results in patient chart.		
Network Reference Range - Reference range (negative: <20 mIU/mL; positive: >20 mIU/mL) - If a specimen is too dilute and has low specific gravity, it may not contain representative levels of hCG and will produce a false negative result. - If pregnancy is suspected, the test may be repeated after 48 hours or by ordering a serum quantitative hCG test.		
Quality Control - External quality controls are required with each new test kit or set of liquid controls - When control results fail to fall within acceptable range repeat the test; contact Point of Care after two consecutive failed results. - Test devices contain two procedural controls (a positive control line that appears in the 'C' area <u>and</u> a clear test area background). Failure of either procedural control indicates an invalid result, and that the test must be repeated with a new test device.		
Supplies and Storage - Discard control solution: at expiration date when stored at 2-8°C; 8 weeks after opening or expiration date, whichever occurs first, when stored at 15-27°C. Label control solutions with open, expiration, and initials. - Discard test kit at expiration date printed on box. Label test kit with a StanBio Pregnancy QC label (print shop #5006). - Storage requirements for test kit (2-30°C) and control solutions (2-27°C).		
Cleaning - Decontaminate benchtops with Clorox Healthcare Bleach Germicidal Wipes. - Cleaning is required daily, and must be documented on the Point of Care Maintenance Record.		
Competency - Healthstream or written test annually. <u>And either</u> - Run two levels of QC each Healthstream assignment or written test. <u>or</u> - Unknown sample in April or October		

The above skills have been successfully demonstrated.

I agree with the assessor that I feel competent to perform the above skills.

Assessor Signature

Employee Signature

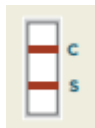
Skills Test

1. When stored at 2-8°C, control solutions are stable until the expiration date printed on the bottles.
 - a. True
 - b. False
2. Once opened, control solutions are stable for _____ weeks at room temperature.
 - a. 3
 - b. 4
 - c. 6
 - d. 8
3. Two levels (positive and negative) of external quality control are required:
 - a. Once per patient testing day
 - b. Once weekly
 - c. Once per new kit
 - d. No external quality control is required
4. When performing a test, _____ drop(s) of urine or control solution should be applied to the round sample well on the device.
 - a. 1
 - b. 2
 - c. 3
 - d. 4
5. Results can be read _____ minutes from sample application.
 - a. 3
 - b. 4
 - c. 5
 - d. 6
6. Control solutions must be labeled upon opening with:
 - a. Expiration
 - b. Open date, expiration, and initials
 - c. Open date, time, and initials
 - d. No label is required
7. The following is true regarding procedural (internal) control **EXCEPT**:
 - a. The negative procedural control is the absence of interfering substance in the background
 - b. The positive control line appears if the test was performed correctly
 - c. The absence of the positive control line is acceptable if a line develops in the sample region
 - d. The positive control line indicates reagent stability

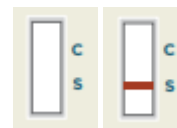
Indicate the correct test result based on the images below:



8. _____



9. _____



10. _____

Score: _____ (minimum 80% to pass, review incorrect answers)

Assessment of Quality Control Performance							
Lot #				Test Kit			
				Exp. Date			
(choice of one)							
Negative QC				Positive QC			
Lot #	Exp. Date	Result	Internal QC	Lot #	Exp. Date	Result	Internal QC

I attest that the above skills test and quality control assessment were successfully completed.

Assessor's Signature

Employee Signature