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| GeneXpert CT/NG Assay | | | | | |
| **Purpose** | This procedure provides instructions for performing the Xpert CT/NG assay on the Cepheid GeneXpert system. | | | | |
| **Policy Statements** | This procedure applies to all technical staff performing testing on the GeneXpert. | | | | |
| **Principle and Clinical Significance** | This qualitative test is intended for use to detect *Chlamydia trachomatis* (CT) and *Neisseria gonorrhoeae* (NG) DNA to aid in the diagnosis of chlamydial and gonorrheal disease.[1]  *Chlamydia trachomatis* are gram-negative bacteria that exist as obligate intracellular parasites. It is the one of the most prevalent sexually transmitted infections (STI). The CT species is comprised of at least 15 disease causing serovars; serovars D through K are the major cause of genital chlamydial infections. If left untreated, CT can cause non-gonococcal urethritis, epididymitis, proctitis, cervicitis, and acute salpingitis. Untreated disease in women can result in pelvic inflammatory disease (PID) and infertility (40% and 20%, respectively).[1, 2]  *Neisseria gonorrhoeae* are gram-negative diplococcic that cause gonorrheal disease, which is the second most commonly reported bacterial STI. Males with a urethral NG infection typically produce symptoms that prompt them to seek treatment. Women often do not produce symptoms until further complications (e.g. PID) occur.[1, 2]  The GeneXpert Instrument System automates and integrates sample purification, nucleic acid amplification, and detection of the target sequences using real-time PCR. The system requires the use of single-use disposable cartridges that hold the PCR regents and host the PCR process.[1, 3]  The Xpert CT/NG Assay includes reagents for the 5’ exonuclease real-time PCR detection and differentiation of CT and NG. Reagents for the detection of a Sample Processing Control (SPC), a Sample Adequacy Control (SAC), and a Probe Check Control (PCC) are also included in the cartridge. The SPC is present to control for adequate processing of the target bacteria and to monitor the presence of inhibitors in the PCR reaction. The SAC reagents detect the presence of a single copy human gene and monitor whether the specimen contains human DNA. The PCC verifies reagent rehydration, PCR tube filling in the cartridge, probe integrity, and dye stability. The primers and probes in the Xpert CT/NG Assay detect chromosomal sequences in the bacteria. One target is detected for CT (CT1) and two different targets are detected for NG (NG2 and NG4). Both NG targets need to be positive for the Xpert CT/NG assay to give a positive NG result.[1, 3]  The Xpert CT/NG Assay is designed for use with genital and urine specimens (first-catch male and female urine, endocervical swab, and vaginal swab specimens) collected in specific Urine and Endocervical /Vaginal Specimen Collection kits designed to preserve patient specimens to allow transport to the laboratory for analysis with Xpert CT/NG Assay and GeneXpert System.[1, 4, 5]  The specimen is briefly mixed by inverting the collection tube several times and transferred to the sample chamber of the Xpert CT/NG cartridge using the supplied transfer pipette (filled to mark). The GeneXpert cartridge is loaded onto the GeneXpert System, which performs hands-of, automated sample processing, and real-time PCR for the detection of DNA in approximately 90 minutes.[1] | | | | |
| **Test Code** | **CGPCR** | | | | |
| **Sample** | 1. **Acceptable specimens:**  * First-catch male and female urine specimens in urine Xpert Collection Tube * Endocervical and vaginal swabs collected with the Xpert CT/NG Vaginal/Endocervical Specimen Collection Kit or Xpert Vaginal/Endocervical Collection Kit  1. **SDES codes/Specimen type:**  * **UR**- Urine, collect method not specified * **VOID**- Voided urine * **VAG**-Vagina * **CERV**-Cervix  1. **Specimen Collection and Transport:**  * Refer to *Lab Test Directory* on StarNet  1. **Specimen assessment:**  * Refer to the policy MCVI 2.1 *Specimen Rejection Criteria.*  1. **Specimen Storage**  * First-catch male and female urine:   + Stability, NEAT:     - Room temp (2-28°C): 24 hours     - 4°C: 8 days   + Stability, in Cepheid Sample Transport Reagent (7mL):     - Room temp (2-28°C): 3 days     - 2-15°C: 45 days * Transfer NEAT urines to Cepheid Sample Transport Reagent as soon as possible after collection. * Endocervical and vaginal swabs collected with the Xpert CT/NG Vaginal/Endocervical Specimen Collection Kit, and the Xpert Vaginal/Endocervical Collection Kit   + Stability in Cepheid Sample transport reagent:     - 2-30°C: 60 days * Specimens should be stored at 4-8°C (refrigerated) | | | | |
| **Special Safety Precautions** | **Microbiologists/virologists are subject to occupational risks associated with specimen handling. Refer to the safety policies located in the safety section of the *Microbiology* and *Virology Policy Manual*:**   1. ***Biohazard Containment*** 2. ***Safety in the Microbiology/Virology Laboratory***  * ***Biohazardous Spills*** | | | | |
| **Materials** | |  |  |  | | --- | --- | --- | | Reagents | Supplies | Equipment | | * 10% bleach * 70% ethanol | * CT/NG Vaginal/endocervical Specimen Collection Kits * CT/NG Specimen Urine Collection Kits * Xpert CT/NG Assay cartridges * Transfer pipettes * Simple racks * Cartridge transfer tray   Store kits at 2-28°C. Kits are stable until the expiration date printed on the outer box. | * Biosafety Hood * Cepheid GeneXpert Instrument and computer * Printer * 1000uL pipette | | | | | |
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| **Calibration** | Annual “Xpert Check Kit” calibration performed by Cepheid. | | | | |
| **Quality Control** | **Daily Quality Control:**  Once an Xpert cartridge has been loaded and before the sample processing steps begin, the software checks the optics, the readiness of the module’s mechanical components, and the ambient temperature of the module to assure proper performance of PCR, and the physical integrity of the cartridge.  Each test includes a Sample Processing Control (SPC), a Sample Adequacy Control (SAC), and a Probe Check Control (PCC).   * SPC: Ensures the sample was correctly processed. It contains DNA from *Bacillus globigii* and verifies the sample processing and target amplification. The SPC verifies that binding and elution of target DNA have occurred if the organisms are present and verifies that sample processing is adequate. Additionally, this control detects sample-associated inhibition of the real-time PCR assay. The SPC should be positive in an analyte negative sample and can be negative OR positive in an analyte positive sample. * SAC: Ensures that the sample contains human cells or human DNA. The SAC signal is only to be considered in an analyte negative sample. A negative SAC indicates that no human cells are present in the sample due to insufficient mixing of the sample or because of an inadequately taken sample. * PCC: Performs a check on the amplification portion of the assay. Before the PCR reaction starts, the GeneXpert instrument measures the fluorescence signal from the probes to monitor bead rehydration, reaction-tube filling, probe integrity, and dye stability. Therefore, it controls for missing or incompletely hydrated beads of enzyme and target specific reagent. It also controls for the generated fluorescence which must meet internal acceptance criteria.   **External Quality Control:**   * Perform QC using external positive and negative controls every 30 days. Record results in the GeneXpert assay binder on the Log. * See IQCP document. * See Quality Control Procedure.   **New Lot/Shipment Quality control:**   * Perform QC using external positive and negative controls with each new lot or shipment before putting into service. Record results in the GeneXpert assay binder on the Log. * See Quality Control Procedure   **Engineering control:**   * Perform wipe testing every 30 days to monitor for contamination. * See Quality Control Procedure.   **NOTE:** External quality control may be performed on an as needed basis if certain circumstances arise. Examples include:   * Drift in results (e.g., increasing/decreasing positivity rates) * Potential contamination (negative control) * After Xpert check or drastic system maintenance | | | | |
| **Procedure** | **Cartridge preparation:**   1. Clean hood with10% bleach (made daily) followed by 70% ethanol. 2. Change gloves. 3. Obtain a CT/NG Assay cartridge, transfer pipette, and sample transport tube to be tested. 4. Label the side of the cartridge with a bar-coded foot-label. 5. Open the cartridge lid. 6. Gently invert the transport tube 3 to 4 times. 7. Open the transport tube lid and draw up specimen in the transfer pipette until the upper fill mark is reached. 8. Insert the pipette to the bottom of the well in the cartridge and empty the pipette’s content into the cartridge.      1. Close the cartridge lid, and set onto the transfer tray. 2. Change gloves and proceed to prepare additional samples or start the test.   NOTES:  -Hood surfaces must be cleaned between samples with 10% bleach followed with 70% ethanol if there were any splashes, spills, or uncertainty of cleanliness.  -\*\*Start the test within 30 minutes of adding the sample to the cartridge  **Starting the test:**   1. Ensure clean gloves are on before stepping to the computer work space. 2. If instrument and computer are turned off: start up the instrument by flipping the power switch located in the back of the instrument. Turn on the computer next. 3. Log onto the appropriate Windows account:    1. User: .\Cepheid-Admin    2. Password: cphd 4. The GeneXpert software will launch automatically. If it doesn’t double-click the GeneXpert Dx software shortcut icon on the desktop. 5. Log onto the software.    1. User: First 6 letters of your first and last name (combined)    2. Password: First 6 letters of your first and last name (combined) 6. In the GeneXpert System window, click **Create Test.** 7. Navigate to the **Sample ID** box. Scan or type in the sample ID. 8. Scan the barcode on the cartridge.   NOTE: if the barcode on the cartridge does not scan, then repeat the test with a new cartridge.   1. Select Xpert CT-NG from the **Select Assay MENU.** 2. Select the appropriate test type for samples or controls. 3. Enter additional information in the “notes” field (day of QC, collect date, etc.) if needed.      1. Click **Start Test**. 2. Enter your username and password, if requested. 3. Open the instrument module door with the blinking green light.   NOTE: when setting up for testing you may opt to use any available module.   1. With the barcode facing towards you, set the cartridge into the module and close the door. 2. Wait for the test to start and the light to stop blinking. The test will run for 90 minutes. 3. Turn printer on. 4. Remove the cartridge when testing is finished (the light will be off and the system will release the door lock). 5. Dispose of used cartridges into bio-bags and place into biohazard sharps bins. 6. Clean any equipment used (pipettes, racks, transfer tray, etc.), hood, and counters (including keyboard, scanner, and mouse) at the end of the day.   NOTE: Sample processing, testing, and cleaning should follow a unidirectional work-flow to avoid contamination. | | | | |
| **Interpretation/ Results** | 1. Click on **View Results** on the top drop-down menu bar and select **View Test**. 2. Select the result you would like to review: Click **OK**. 3. Review result interpretations and amplification curves for exponential growth.    1. NOTE: SAC and SPC do not need to pass for a positive result to be valid.    2. NOTE: SAC and SPC do need to pass for a negative result to be valid.      1. Click on the **Errors** tab to ensure no errors occurred during testing. (Section 9.18.2 in Operator Manual provides error code descriptions)   **Reasons to retest:**   1. An INVALID result. This may indicate:    1. The sample was inadequate.    2. The sample was not properly processed.    3. PCR was inhibited. 2. An ERROR result. This may indicate:    1. The reaction tube was filled improperly.    2. A reagent probe integrity problem was detected.    3. The maximum pressure limit was exceeded.    4. A valve positioning error was detected. 3. NO RESULT:    1. This result indicated that insufficient data were collected (e.g. test stopped while in progress or power failure occurred).   NOTE: Record any failures on the “GeneXpert Service and Error Log” log. | | | | |
| **Result Reporting** | 1. Ensure that the printer is turned on.    1. Reports will print automatically. 2. Results will automatically transmit to the LIS. 3. Log into Sunquest to release results. 4. Select Result Entry from Menu options 5. In the Configuration field select CGX from the dropdown box.      1. Click on the  button located in the lower right corner to populate the transmitted results. 2. Review messages located on the top and results. Compare results to the GeneXpert report. | | | | |
|  | 1. Check the release box. 2. Click  button located on the lower left corner. Click  when the “Verify Release Destination” window opens. 3. Call a completed worksheet for CGPCR, check results, and staple to Genexpert Report. Place in the GeneXpert CT/NG result binder. 4. Store samples in fridge:    1. Mark positive samples on side of caps (1 line: CT, 2 lines: NG, 3 lines: dual positive). 5. Discard old samples after 2 weeks. | | | | |
| **Critical Results** | 1. The code SURE (Semi-urgent result) will automatically append if the result is Positive AND the patient is < 12 y/o. These results must be called to the provider. Add the code CAL, press tab, enter semi-colon, record who the result was relayed to and the time/date. | | | | |
| **Invalid Results** | 1. IF an invalid result is repeated AND a **valid** result is obtained, select and only release the valid result interpretation in the LIS. 2. IF an invalid result is repeated AND an **invalid** result is obtained, select only one of the invalid results to verify. The provider must be notified of these results.   The CGPCR result will be reported as **unresolved** (UNRE) and the following code SIA will automatically append: “This sample is inhibitory to amplification and the results are inconclusive. Consider repeat collection if clinically indicated.”  Add the code CAL, press tab, enter semi-colon record who the result was relayed to and the date/time. | | | | |
| **Correcting Results** | 1. Open Result Entry, select the Manual resulting mode (top left corner), from the configuration drop down select CGPCR. Click  in the lower right corner.      1. Enter the Specimen ID, enter Tab and click Yes to modify the result. | | | | |
|  | 1. Change the incorrect result. The corrected result comment will automatically append. Add the CAL comment, press tab, enter a semi-colon and record who was called and the time/date.      1. Click . Click  when the “Verify Release Destination” window opens. | | | | |
| **Limitations** | • The Xpert CT/NG Assay has only been validated with the following specimen types, collected with the Cepheid Xpert CT/NG Vaginal/Endocervical, Xpert Vaginal/Endocervical, Xpert CT/NG Urine or Xpert Urine Specimen Collection Kits:  • Endocervical swabs  • Patient-collected vaginal swabs  • Male and female urine   * Because the detection of CT and NG is dependent on the DNA present in the sample, reliable results are dependent on proper sample collection, handling and storage. * With endocervical and patient-collected vaginal specimens, assay interference may be observed in the presence of: blood (>1% v/v) or mucin (>0.8% w/v). * With urine specimens, assay interference may be observed in the presence of: blood (>0.3% v/v), mucin (>0.2% w/v), bilirubin (>0.2 mg/mL), or Vagisil feminine powder (>0.2% w/v). * The effects of other potential variables such as vaginal discharge, use of tampons, douching, and specimen collection variables have not been determined. * A negative test result does not exclude the possibility of infection. * The Xpert CT/NG Assay should not be used for the evaluation of suspected sexual abuse or for other medico-legal indications. Additional testing is recommended in any circumstance when false positive or false negative results could lead to adverse medical, social, or psychological consequences. * The Xpert CT/NG Assay provides qualitative results. No correlation can be drawn between the magnitude of the Ct value and the number of cells in an infected sample. * The predictive value of an assay depends on the prevalence of the disease in any particular population. Positive results in a low prevalence population should be evaluated carefully. * The Xpert CT/NG Assay should not be used to monitor therapeutic success as residual target nucleic acid may persist for up to three weeks. * Xpert CT/NG Assay performance has not been evaluated in patients less than 14 years of age. During validation testing a total of eight patients under the age of 14 had samples submitted for testing (6 urine, 2 vaginal). All sample results were negative and in agreement with the comparator method, with the exception of one that was invalid. Due to a low frequency of testing this population and availability of resources, a more thorough evaluation was not possible. * Xpert CT/NG Assay performance has not been evaluated in pregnant women, or in patients with a history of hysterectomy. * The Xpert CT/NG Assay has not been validated for use with vaginal swab specimens collected by patients at home. The patient-collected vaginal swab specimen application is limited to healthcare facilities where support/counseling is available to explain procedures and precautions. * The Xpert CT/NG Assay has not been evaluated with patients who are currently being treated with antimicrobial agents active against CT or NG. * Results from the Xpert CT/NG Assay should be interpreted in conjunction with other laboratory and clinical data available to the provider.[1] * The NG4 target appears to have some homology to sequences in some *N. mucosa* and *N. subflava* strains.[6] | | | | |
| **Method Performance Specifications** | According to the manufacturer (per the package insert):   |  |  |  | | --- | --- | --- | | **Female Swabs** | **CT** | **NG** | | Vaginal: Sensitivity | 99.5% | 100% | | Vaginal: Specificity | 99.1% | 99.9% | | Endocervical: Sensitivity | 96.0% | 100% | | Endocervical: Specificity | 99.6% | >99.9% | |  |  |  | | **Urine** |  |  | | Female: Sensitivity | 98.1% | 94.4% | | Female: Specificity | 99.8% | >99.9% | | Male: Sensitivity | 98.5% | 98.3% | | Male: Specificity | 99.8% | 99.9% |   The LODs were determined by using two serovars of CT and two strains of NG. Testing was performed on patient collected vaginal swabs and male urines.   |  |  |  |  |  | | --- | --- | --- | --- | --- | |  | **CT Serovar (eb/mL) D** | **CT Serovar (eb/mL) H** | **NG Strain (cfu/mL) #19424** | **NG Strain (cfu/mL) #49226** | | **Vaginal swab** | 84 | 161 | 1.5 | 1.6 | | **Urine** | 75 | 134 | 2.7 | 1.2 | | | | | |
| **References** | 1. **Xpert CT/NG Assay Package Insert, Rev. D**. In. Sunnyvale, CA: Cepheid Inc.; 2016.  2. **Sexually Trasmitted Diseases**. In. Atlanta, GA: Centers for Disease Control and Prevention.  3. **GeneXpert Dx System Operator Manual: Software Version 4.8, 3010045, Rev. K**. In. Sunnyvale, CA: Cephied Inc.; 2016.  4. **Xpert Urine Specimen Collection Kit Package Insert, Rev. 01**. In. Sunnyvale, CA: Cepheid Inc.; 2016.  5. **Xpert Vaginal/Endocervical Specimen Collection Kit Package Insert, Rev. 00**. In. Sunnyvale, CA: Cepheid Inc.; 2015.  6. Tabrizi SN, Unemo M, Golparian D, Twin J, Limnios AE, Lahra M, et al. Analytical evaluation of GeneXpert CT/NG, the first genetic point-of-care assay for simultaneous detection of Neisseria gonorrhoeae and Chlamydia trachomatis. *Journal of clinical microbiology* 2013; 51(6):1945-1947. | | | | |
| **Alternate Methods** | 1. Hologic Chlamydia trachomatis Amplified RNA Assay (Mayo Medical Laboratories, collected in Aptima Collection Kit) 2. Chlamydia trachomatis culture 3. Chlamydia trachomatis direct FA 4. Hologic Neisseria gonorrhoeae Amplified RNA assay (Mayo Medical Laboratories, collected in Aptima Collection Kit) 5. GC culture 6. Genital culture | | | | |
| **Proficiency Testing** | CAP materials: 3 shipments a year with 5 samples. | | | | |
| **Training Plan/ Competency Assessment** | **Training Plan** | | | **Initial Competency Assessment** | |
| 1. Employee must read the procedure. 2. Employee will demonstrate the ability to perform procedure, record results, and document corrective action after instruction by the trainer. | | | 1. Direct observation | |
| **Historical Record** |  |  |  | |  |
|  | **Version** | **Written/Revised by:** | **Effective Date:** | | **Summary of Revisions** |
| 1 | Julie Laramie/Helen Stefan | 04.16.2018 | | Initial Version |
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