

If a conflict arises between a Clinical Payment and Coding Policy and any plan document under which a member is entitled to Covered Services, the plan document will govern. If a conflict arises between a CPCP and any provider contract pursuant to which a provider participates in and/or provides Covered Services to eligible member(s) and/or plans, the provider contract will govern. "Plan documents" include, but are not limited to, Certificates of Health Care Benefits, benefit booklets, Summary Plan Descriptions, and other coverage documents. Blue Cross and Blue Shield of New Mexico may use reasonable discretion interpreting and applying this policy to services being delivered in a particular case. BCBSNM has full and final discretionary authority for their interpretation and application to the extent provided under any applicable plan documents.

Providers are responsible for submission of accurate documentation of services performed. Providers are expected to submit claims for services rendered using valid code combinations from Health Insurance Portability and Accountability Act approved code sets. Claims should be coded appropriately according to industry standard coding guidelines including, but not limited to: Uniform Billing Editor, American Medical Association, Current Procedural Terminology, CPT® Assistant, Healthcare Common Procedure Coding System, ICD-10 CM and PCS, National Drug Codes, Diagnosis Related Group guidelines, Centers for Medicare and Medicaid Services National Correct Coding Initiative Policy Manual, CCI table edits and other CMS guidelines.

Claims are subject to the code edit protocols for services/procedures billed. Claim submissions are subject to claim review including but not limited to, any terms of benefit coverage, provider contract language, medical policies, clinical payment and coding policies as well as coding software logic. Upon request, the provider is urged to submit any additional documentation.

Diagnostic Testing of Common Sexually Transmitted Infections

Policy Number: CPCPLAB051

Version 1.0

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Description

The Plan has implemented certain lab management reimbursement criteria. Not all requirements apply to each product. Providers are urged to review Plan documents for eligible coverage for services rendered.

Reimbursement Information:

This policy is limited to testing for *C. trachomatis, N. gonorrhoeae, T. pallidum, T. vaginalis,* HSV, and HPV. The following conditions and/or tests are discussed in the corresponding policies:

- Human Immunodeficiency Virus CPCPLAB065
- Hepatitis B and C CPCPLAB015
- Pediatric Preventive Screening CPCPLAB016
- Cervical Cancer Screening CPCPLAB002
- Pathogen Panel Testing CPCPLAB045

For STI screening in pregnant individuals, please see CPCPLAB014 Prenatal Screening (Nongenetic).

- 1. Antibody testing for syphilis infection **may be reimbursable** in the following situations:
 - a. For any asymptomatic person in a high-risk category (See Notes 1 & 2), once a year assessment using either a "standard" or "reverse" algorithm that includes initial and confirmatory tests for any initial positive test such as:
 - i. Treponemal Ig test **AND**
 - ii. Nontreponemal test;
 - For diagnosis of any person presenting with signs and/or symptoms of a syphilis infection* (See Note 3);
 - Once every three months for HIV-positive men or men who have sex with men (MSM);
 - d. Treponemal Ig testing and nontreponemal testing (once prior to transplant) as a part of a pre-transplant assessment in both donors and recipients for an allogeneic hematopoietic stem cell transplantation (allo-HCT);
 - e. When a nontreponemal test is used as a test of cure (TOC) for a positive syphilis infection.
- For asymptomatic individuals NOT belonging to a high-risk category (See Notes 1 & 2) antibody screening for syphilis may be reimbursable only in the following situations:
 - a. As part of newborn screening;
 - b. As part of follow-up in a victim of sexual assault;
 - c. For sexually active individuals less than 18 years of age (annually).

- 3. Polymerase chain reaction (PCR) and nucleic acid amplification testing (NAAT) for syphilis **is not reimbursable**.
- 4. Qualitative nucleic acid amplification tests (NAATs) for chlamydia **may be reimbursable** in the following situations:
 - a. Once a year assessment for any asymptomatic person in a high-risk category (**See Notes 1 & 4**);
 - b. For diagnosis of any person presenting with signs and/or symptoms of a chlamydial infection (**See Note 5**);
 - c. For diagnosis of any person with suspected lymphogranuloma venereum (LGV);
 - d. As test of cure of treatment at least three months after initial chlamydial diagnosis.
- For asymptomatic individuals NOT belonging to a high-risk category (See Note 1 & 4), NAAT screening for chlamydia may be reimbursable only in the following situations:
 - a. As part of newborn screening;
 - b. As part of follow-up of victim of sexual assault;
 - c. For sexually active individuals less than 18 years of age (annually).
- 6. Serology testing for chlamydia or lymphogranuloma venereum (LGV) **is not reimbursable.**
- 7. Qualitative nucleic acid amplification tests (NAATs) for gonorrhea **may be reimbursable** in the following situations:
 - a. Once a year assessment for any asymptomatic person in a high-risk category (**See Note 1 & 4**);
 - b. For diagnosis of any person presenting with signs and/or symptoms of a gonorrheal infection (**See Note 6**);
 - c. As test of cure for treatment.
- 8. For an individual that does not respond to initial treatment, culture testing for *N. gonorrhoeae* to determine antimicrobial susceptibility **may be reimbursable**.
- For asymptomatic individuals NOT belonging to a high-risk category (See Note 1 & 4), NAAT screening for gonorrhea may be reimbursable only in the following situations:
 - a. As part of newborn screening;
 - b. As part of follow-up of victim of sexual assault;
 - c. For sexually active individuals less than 18 years of age (annually).
- 10. Qualitative NAAT for *T. vaginalis* **may be reimbursable** in the following situations:
 - a. For symptomatic individuals (See Note 7);
 - b. Follow-up testing a minimum of three months after initial trichomoniasis diagnosis

- c. Annual screening for asymptomatic individuals belonging to a high-risk group (See **Note 8**)
- d. Annual screening for asymptomatic individuals who have an HIV infection
- e. As part of follow-up in a victim of sexual assault.
- 11. Rapid identification of *Trichomonas* by enzyme immunoassay **is not reimbursable.**
- 12. For symptomatic individuals (**See Note 8**), testing for *Mycoplasma genitalium* using qualitative NAAT **may be reimbursable**.
- 13. For asymptomatic individuals (**See Note 9**), screening for *M. genitalium* using NAAT **is not reimbursable.**
- 14. When an individual meets the conditions described above, multitarget PCR testing (targets limited to *C. trachomatis, N. gonorrhoeae, T. vaginalis,* and *M genitalium*) **may be reimbursable.**
- 15. For individuals with active genital ulcers or mucocutaneous lesions, qualitative NAAT for herpes simplex virus (HSV-1) or herpes simplex virus-2 (HSV-2) **may be reimbursable**.
- 16. Immunoassay testing for herpes simplex virus-1 (HSV-1), and/or herpes simplex (non-specific type test) **is not reimbursable**.
- 17. Type-specific serologic testing for herpes simplex virus-2 (HSV-2) using a glycoprotein G2 (gG2) **may be reimbursable** in the following situations:
 - a. Recurrent or atypical genital symptoms or lesions in individuals with a negative herpes simplex virus PCR or culture result;
 - b. For the clinical diagnosis of genital herpes in individuals with a negative PCR or culture result or without laboratory confirmation;
 - c. When an individual's partner has genital herpes.
- 18. Screening for herpes simplex virus-1 or herpes simplex virus-2 (HSV-1 and HSV-2) in asymptomatic individuals **is not reimbursable**.
- 19. In the diagnosis and/or assessment of cancer or cancer therapy (immunohistochemistry testing for p16 or NAAT testing for high-risk human papillomavirus [HR-HPV]), testing for HR-HPV **may be reimbursable**.
- 20. Testing for HPV **is not reimbursable** in the following situations:
 - To screen for oncogenic high-risk types, such as HPV-16 and HPV-18, as part of a general sexually transmitted disease (STD) or sexually transmitted infection (STI) screening process or panel for asymptomatic patients;
 - b. As part of diagnosis of anogenital warts;
 - c. Testing for low-risk types of HPV;

- d. In the general population either as part of a panel of tests or as an individual NAAT to determine HPV status.
- 21. Prior to beginning a Preexposure prophylaxis (PrEP) regimen, triple panel testing (hepatitis B surface antigen [HBsAg], hepatitis B surface antibody [anti-HBs], total antibody to hepatis B core antigen [anti-HBc]) to screen for hepatitis B **may be reimbursable**:
- 22. Prior to beginning or while an individual is undergoing a preexposure prophylaxis (PrEP) regimen for HIV prevention, the following screens/tests for additional STIs **may be reimbursable**:
 - a. Qualitative NAAT screening for gonorrhea and chlamydia:
 - i. Once every three months for MSM
 - ii. Once every six months for sexually active individuals
 - b. Blood test to screen for syphilis:
 - i. Once every three months for MSM
 - ii. Once every six months for sexually active individuals.
- 23. Nucleic acid testing to determine antimicrobial susceptibility in *N. gonorrhoeae* or macrolide resistance in *M. genitalium* **is not reimbursable.**
- 24. Direct probe detection and/or quantitative NAAT for the following microorganisms is not reimbursable:
 - a. Chlamydia trachomatis
 - b. Neisseria gonorrhoeae
 - c. Herpes Simplex Virus-1
 - d. Herpes Simplex Virus-2
 - e. Human Papillomavirus
 - f. Treponema pallidum

NOTE 1: For sexually active children and adolescents under the age of 18, risk factors for chlamydia, gonorrhea and/or syphilis infection as defined by the CDC include: (4)

- Initiating sex early in adolescence;
- Living in detention facilities;
- Receiving services at STD clinics;
- Being involved in commercial sex exploitation or exchanging sex for drugs, money, food, or housing;
- Having multiple sex partners;
- Having sequential sex partners of limited duration or concurrent partnerships;
- Failing to use barrier protection consistently and correctly;
- Having lower socioeconomic status, and facing numerous obstacles to accessing healthcare;
- At risk individuals also include:
 - Males who have sex with males (YMSM);
 - Transgender youths;

o Youths with disabilities, substance abuse, or mental health disorders.

NOTE 2: High-risk for Syphilis (5,6):

- Sexually active men who have sex with men (MSM);
- Sexually active individuals with an HIV-positive status;
- Having a sexual partner recently diagnosed with an STI;
- · Exchanging sex for money or drugs;
- Individuals in adult correctional facilities;
- During pregnancy when the following risk factors are present:
 - o Sexually active HIV-positive status;
 - Sexually active with multiple partners;
 - Sexually active in conjunction with drug use or transactional sex;
 - Late-entry to prenatal care (i.e., first visit during the second trimester or later) or no prenatal care;
 - o Methamphetamine or heroin use;
 - o Incarceration of the woman or her partner;
 - Unstable housing or homelessness.

NOTE 3: Signs and Symptoms of a Syphilis Infection (5,7)

- Chancre;
- Skin rash and/or mucous membrane lesions in mouth, vagina, anus, hands, and feet;
- Condyloma lata;
- Secondary symptomology can include fever, fatigue, sore throat, swollen lymph nodes, weight loss, muscle aches, headache, and hair loss;
- Signs and symptoms of neurosyphilis can include severe headache, trouble with muscle movements, muscle weakness or paralysis (not being able to move certain parts of the body), numbness, and changes in mental status (trouble focusing, confusion, personality change) and/or dementia (problems with memory, thinking, and/or making decisions).
- Signs and symptoms of ocular syphilis can include eye pain or redness, floating spots in the field of vision ("floaters"), sensitivity to light, and changes in vision (blurry vision or even blindness).
- Signs and symptoms of otosyphilis may include hearing loss, ringing, buzzing, roaring, or hissing in the ears ("tinnitus"), balance difficulties, and dizziness or vertigo.
- Signs and symptoms of late/tertiary syphilis include inflammatory lesions of the cardiovascular system (e.g., aortitis, coronary vessel disease), skin (e.g., gummatous lesions), and bone (e.g., osteitis).

NOTE 4: High-risk for Chlamydia and/or Gonorrhea (8-11):

- Sexually active men who have sex with men (MSM);
- Sexually active individuals with an HIV-positive status;
- Sexually active individuals with a cervix who are under the age of 25;
- Individuals with a cervix who are 25 years of age or older and who have multiple sexual partners, multiple sexual partners, or a sex partner with concurrent partners; practice inconsistent condom use when not in a mutually monogamous relationship;

- Having a sexual partner recently diagnosed with an STI;
- Previous or concurrent STI;
- · Exchanging sex for money or drugs;

NOTE 5: Signs and Symptoms of a Chlamydia Infection (8,11):

- Genital symptoms, including "discharge, burning during urination, unusual sores, or rash;"
- Pelvic Inflammatory Disease, including "symptoms of abdominal and/or pelvic pain, along with signs of cervical motion tenderness, and uterine or adnexal tenderness on examination;"
- Urethritis;
- Pyuria;
- Dysuria;
- Increase in frequency in urination;
- Epididymitis (with or without symptomatic urethritis) in men;
- Proctitis;
- Sexually acquired chlamydial conjunctivitis.

NOTE 6: Signs and Symptoms of Gonorrhea (10):

- Dysuria;
- Urethral infection;
- Urethral or vaginal discharge;
- Epididymitis (Testicular or scrotal pain);
- Rectal infection symptoms include anal itching, discharge, rectal bleeding, and painful bowel movements.

NOTE 7: Signs and Symptoms of Trichomoniasis (12):

- Vaginal or penile discharge;
- Itching, irritation, burning sensation, or soreness of the genitalia;
- Discomfort or burning sensation during/after urination and/or ejaculation;
- Urethritis;
- Epididymitis;
- Prostatitis.

Note 8: <u>High-risk for Trichomoniasis:</u> (13)

- Receiving care in high-prevalence settings (e.g., STI clinics, correctional facilities)
- Having multiple sexual partners
- Exchanging sex for money or drugs
- Having a previous or concurrent STI
- Drug misuse
- History of incarceration
- Sexually active individuals with an HIV-positive status

NOTE 9: Signs and Symptoms of *M. genitalium* Infection (14):

• When present, typical symptoms of *M.gen*-urethritis in men include dysuria, urethral pruritus, and purulent or mucopurulent urethral discharge.

- When present, typical symptoms of *M.gen* cervicitis in women include vaginal discharge, vaginal itching, dysuria, and pelvic discomfort.
- When present, typical symptoms of PID due to M.gen include mild to severe pelvic pain, abdominal pain, abnormal vaginal discharge, and/or bleeding.

Procedure Codes

The following is not an all-encompassing code list. The inclusion of a code does not guarantee it is a covered service or eligible for reimbursement.

Codes

86592, 86593, 86631, 86632, 86694, 86695, 86696, 86704, 86706, 86780, 87081, 87110, 87181, 87340, 87490, 87491, 87492, 87528, 87529, 87530, 87563, 87590, 87591, 87592, 87623, 87624, 87625, 87626, 87661, 87797, 87798, 87799, 87800, 87808, 88341, 88342, 88344, 0064U, 0065U, 0096U, 0210U, 0402U, 0455U, 0463U, 0483U, 0484U, G0499,

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Policy Update History:

Approval Date	Effective Date; Summary of Changes
09/26/2025	01/03/2026 Document updated with literature review. The
	following changes were made to Reimbursement Information:
	For clarity added "qualitative" to "qualitative NAAT' in #4, #7,
	#12, #15; added "NAAT" to #5 and #9; revised #10 to state
	"Qualitative NAAT for T. vaginalis may be reimbursable in the
	following situations: a) For symptomatic individuals (see Note
	5); b) Follow-up testing a minimum of three months after initial
	trichomoniasis diagnosis; c) Annual screening for
	asymptomatic individuals belong to a high-risk group (see
	Note 6); d) Annual screening for asymptomatic individuals who

have an HIV infection; e) As part of follow-up in a victim of sexual assault." Revised #21 to state: "Prior to beginning a Preexposure prophylaxis (PrEP) regimen, triple panel testing (hepatis B surface antibody (anti-HBc)), total antibody to hepatitis B core antigen [anti-HBc)] to screen for hepatitis B may be reimbursable." Revised #22 to state: "Prior to beginning or while an individual is undergoing a preexposure prophylaxis (PrEP) regimen for HIV prevention, the following screens/tests for additional STIs may be reimbursable: a) Qualitative NAAT screening for gonorrhea and chlamydia: i) Once every three months for MSM. ii) Once every six months for sexually active individuals. b) Blood testing to screen for syphilis: i) Once every three months for MSM. ii) Once every six months for sexually active individuals." Revised leading statement in #24 to state: "Direct probe detection and/or quantitative NAAT". Added codes 87626, 87800; removed codes 82565, 82575, 84702, 84703, 86701, 86702, 86703, 86705, 86803, 86804, 87660, 0500T, G0432, G0433, G0435, G0472, G0475, S3645. References revised. 10/30/2024 10/15/2025; Document updated with literature review. The following changes were made to Reimbursement Information: Added 1.d. Treponemal Ig testing and nontreponemal testing (once prior to transplant) as a part of a pre-transplant assessment in both donors and recipients for an allogeneic hematopoietic stem cell transplantation (allo-HCT); added 12: For symptomatic individuals (See Note 8), screening for M. genitalium using NAAT is not reimbursable; added 1. vaginalis, and M genitalium of a syphilis infection to Note 3; added Note 8 for Signs and Symptoms of M. genitalium infection. Added codes 87563, 0402U, 0455U, 0463U, 0483U, 0483U, 0484U; removed 0167U, 0353U, 0354U. References revised. 11/01/2023: Document updated with literature review. Reimbursement information revised for clarity. Added #10: When an individual meets the conditions described above for both chlamydia and gonorrhea, multitarget PCR testing		
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