DST 906: Care and Treatment Plan:

Gonorrhea

Care and Treatment Plan: Gonorrhea (reportable)

Definition

Bacterial infection caused by the transmission of *Neisseria gonorrhoeae* (*N. gonorrhoeae* or GC) during sexual contact in which body fluids are exchanged.

Registered Nurses with **Reproductive health – sexually transmitted infection** Certified Practice designation (RN(C)) are authorized to manage, diagnose, and treat individuals with Gonorrhea.

Potential Causes

Bacterial:

• Neisseria gonorrhoeae

Predisposing Risk Factors

• Sexual contact where there is transmission through the exchange of body fluids

Typical Findings

Sexual History

- · Sexual contact with at least one partner
- May be asymptomatic
- Sexual contact with someone with confirmed positive laboratory test for STI
- Short incubation period (e.g., urethritis and discharge may develop within one week of sexual contact)

Physical Assessment

GC infection can be asymptomatic in all sites; however, urethral GC infections are often symptomatic, while pharyngeal, vaginal and rectal infections are often asymptomatic.

- Inflammation of the tissues around the eye including acute redness, purulent discharge and crusting (symptoms of
 conjunctivitis); can be caused from gonococcal infection in the eye; consult with or refer to physician or nurse
 practitioner (NP) for symptoms of conjunctivitis
- Sore throat (throat infection is most often asymptomatic)
- Urethral symptoms such as, discharge (usually purulent, may be copious), itch or awareness
- Painful (dysuria) or difficult urination
- Testicular pain and/or swelling (symptoms of epididymitis)
- Abnormal change in vaginal discharge
- Abnormal vaginal bleeding:
 - o Vagina with or without cervix: after intercourse or between menstrual period
 - Vagina after vaginoplasty: abnormal vaginal bleeding is not always STI-related as longer post-operative symptoms
 of bleeding could be indicative of hypergranulation; refer to the STI history and physical exam information listed
 above (page 3-5) for more information, and especially for clients experiencing pain, discharge, or bleeding in the
 first 3 to 4 month post-operative period
- Lower abdominal pain (symptom of pelvic inflammatory disease)
- Dyspareunia
- Inflammation of the rectum, rectal pain and anal discharge (symptoms of proctitis)

Diagnostic Tests

Full STI screening is recommended. See DST 900: Assessment and Diagnosistic Guideline: STI

GC C&S is indicated for all clients who are symptomatic and/or are a contact to gonorrhea.



Throat:

- For clients who are symptomatic and/or are a contact to GC, collect:
 - GC culture & sensitivity (C&S) swab
 - GC NAAT swab
- For asymptomatic clients who require screening only, collect:
 - GC NAAT swab, if indicated in sexual health history

Penile urethra (with or without phalloplasty or metoidioplasty with urethral lengthening):

- For clients who are symptomatic with urethral discharge and/or are a contact to GC, collect specimen from visible discharge at the meatus insertion into the urethra is not required:
 - GC C&S swab
 - Smear for typical intracellular diplococci (TID) and polymorphonuclear leukocytes (PMNs) (collect only if immediate microscopy is available)
 - GC NAAT urine: ideally the client should not have voided in the previous 1-2 hours; collect first void 10-20ml

Vagina:

With cervix:

- Asymptomatic: GC NAAT vaginal (preferred) or cervical swab
- Symptomatic or contact to GC: a full pelvic examination is recommended to collect GC C&S cervical (preferred) or vaginal swab (when unable to perform pelvic exam) and GC NAAT vaginal (preferred) or cervical swab

After total hysterectomy (no cervix):

- o Asymptomatic: GC NAAT urine (preferred) or vaginal swab
- o Symptomatic or contact to GC: GC C&S vaginal swab and GC NAAT urine specimen (preferred) or vaginal swab

After vaginoplasty:

- GC NAAT urine: ideally the client should not have voided in the previous 1-2 hours; collect first void 10-20ml
- Asymptomatic/symptomatic-GC NAAT urine

Rectum:

- For clients who are symptomatic and/or are a contact to GC, collect:
 - o GC C&S swab
 - GC NAAT swab
- For asymptomatic clients who require screening only, collect:
 - GC NAAT swab, if indicated in sexual health history

Notes:

- 1) If urethral swabs are indicated (e.g., for symptomatic clients), the urine specimen is collected after the urethral swab.
- 2) GC NAAT urine specimens may be collected as the only diagnostic test in agencies or circumstances where:
 - o GC C&S is unavailable, and the client is symptomatic
 - o client is asymptomatic
 - o if client declines recommended testing of cervical or vaginal sites
- 3) In general, self-collected vaginal swabs are indicated when a full or partial pelvic examination is not required or appropriate. Clinician-collected vaginal swabs are generally done when a partial or full pelvic examination is required or requested by the client.
- 4) Recent data show that NAAT vaginal swabs for *C. trachomatis* and *N. gonorrhoeae* identify as many or more infections over cervical, urethral, or urine specimens.



Clinical Evaluation/Clinical Judgement

Treat all clients with confirmed gonorrhea by positive laboratory result.

If providing treatment for a client with confirmed positive cervical, vaginal or urine laboratory test for *Neisseria gonorrhoeae*; assess for signs of pelvic inflammatory disease (PID) through symptoms inquiry and/or physical assessment (bimanual exam), if indicated.

Treat all persons identified as a sexual contact within the past 60 days to a confirmed gonorrhea or case. If there are no sexual contacts in the previous 60 days, then follow-up should occur for the last sexual contact.

Management and Interventions

Goals of Treatment

- · Treat infection
- Prevent complications
- Prevent the spread of infection

Treatment of Choice

Treatment	Notes
First Choice	General:
Cefixime 800mg PO in a single dose and Azithromycin 1gm PO in a single dose OR Ceftriaxone 250mg IM in a single dose and	 Treatment covers both gonorrhea and chlamydia. Canadian Guidelines for STI (CGSTI, PHAC, 2013) recommend ceftriaxone IM and azithromycin PO for the treatment of uncomplicated anogenital and pharyngeal infection; however, BC surveillance patterns of GC resistance suggest that both cefixime and ceftriaxone are appropriate choices for the treatment of GC. Future GC Treatment regimens will continue to reflect national recommendations in association with local GC antimicrobial resistance trends (AMR) trends. Retreatment is indicated if the client has missed 2 consecutive doses of
Azithromycin 1gm PO in a single dose Second Choice Cefixime 800mg PO in a single dose and Doxycycline 100mg PO BID for 7 days	 doxycycline or has not completed a full 5 days of treatment. 5. Consult a physician or NP if client is unable to use cefixime, ceftriaxone, or azithromycin. 6. See BCCDC <u>STI Medication Handouts</u> for further medication reconciliation and client information. 7. See <i>Monitoring and Follow-up</i> section for test-of-cure (TOC) requirements.
OR Ceftriaxone 250mg IM in a single dose and Doxycycline 100mg PO BID for 7 days	 Allergy and Administration: DO NOT USE ceftriaxone or cefixime if history of allergy or anaphylaxis to cephalosporins. Consult with or refer to a physician or NP if history of anaphylaxis or immediate reaction to penicillin. DO NOT USE azithromycin if history of allergy to macrolides. DO NOT USE doxycycline if pregnant and/or allergic to doxycycline or other tetracyclines. If an azithromycin or doxycycline allergy or contraindication exists, consult with/refer to a physician or NP for alternate treatment.

Gonorrhea



Third Choice

Azithromycin 2gm PO in a single dose

- 5. Azithromycin and doxycycline are sometimes associated with gastrointestinal adverse effects. Taking medication with food and plenty of water may minimize adverse effects.
- 6. The preferred diluent for ceftriaxone IM is 0.9 ml lidocaine 1%.
- 7. DO NOT USE lidocaine if history of allergy to lidocaine or other local anesthetics. Use cefixime PO as alternate treatment.
- 8. For <u>IM injections of ceftriaxone</u> the ventrogluteal site is preferred.
- Advise the client to remain in the clinic for at least 15 minutes-post IM injection in case of anaphylactic reaction to treatment. Provide anaphylaxis treatment as required, using BCCDC <u>CDC Manual- Chapter 2: Immunization Part 3: Management of Anaphylaxis in a Non-Hospital Setting</u>, November 2016.
- 10. If serious allergic reaction develops including difficulty breathing and/or severe itchiness, have the client inform clinic staff immediately. If symptoms develop after leaving the clinic, advise the client to seek immediate emergency care.
- 11. Advise client they may experience pain redness and swelling at the injection site. If any of these effects persist or worsen advise to contact health care provider.
- 12. Recent data has emerged regarding azithromycin and QT prolongation. Although rare, it is more significant in older populations, those with preexisting heart conditions, arrhythmias, or electrolyte disturbances.

It is unclear how significant these findings are in young to mid-age healthy adults consuming a one-time dose of azithromycin; however, please use the following precautions:

Consult with or refer to an NP or physician if the client:

- Has a history of congenital or documented QT prolongation.
- Has a history of electrolyte disturbance in particular hypokalemia, hypomagnesaemia.
- Has clinically relevant bradycardia, cardiac arrhythmia, or cardiac insufficiency.
- o Is on any of the following medications:
 - Antipsychotics: pimozide (Orap®), ziprasidone (Zeldox®)
 - Cardiac: dronedarone (Multaq®)
 - Migraine: dihydroergotamine (Migranal®), ergotamine (Cafergot®)

Pregnant or Breast-/Chest-Feeding Adults

For all pregnant or breast-/chest-feeding clients, consult with or refer to a physician or NP. Test-of-cure (TOC) is recommended for pregnant and/or breast-/chest-feeding clients and should be performed at 3-4 weeks after completion of treatment

Partner Counselling and Referral

People who have confirmed laboratory tests positive for *Neisseria gonorrhoeae* are offered partner counselling and referral to identify all the people who may have been exposed through sexual contact in the previous 60 days. If no sexual partner in the previous 60 days then follow up should occur for the last sexual contact (see *DST 901: Care and Treatment Plan: Treatment of STI Contacts*).



Monitoring and Follow-up

Repeat testing at 6 months is recommended due to potential high risk of re-infection.

Perform TOC in the following situations:

- If symptoms persist
- For all pregnant and/or breast-/chest-feeding clients
- · For all pharyngeal infections
- If treatment was other than the recommended first choice
- For clients who received antibiotics linked to a case who had treatment failure or demonstrated resistance to the same antibiotic

TOC using GC culture (C&S) should be performed a minimum 3-7 days post-treatment completion. TOC using GC NAAT (if culture is not available), should be performed 3-4 weeks post-treatment completion.

Potential Complications

- · Epididymitis
- Infertility
- Sexually-acquired reactive arthritis
- · Disseminated gonococcal infection (DGI)
- Pelvic inflammatory disease (PID)
- Ectopic pregnancy
- Chronic pelvic pain

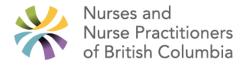
Client Education

Counsel client regarding:

- Abstaining from sexual activity during the 7-day course of treatment or for 7 days post single-dose therapy for clients and their contacts.
- Informing last sexual contact AND any sexual contacts within the last 60 days that they require testing and treatment.
- Methods of partner notification.
- The appropriate use of medications (dosage, side effects, and need for re-treatment if dosage not completed, or symptoms do not resolve).
- Harm reduction (e.g., condom use significantly reduces the risk of transmission).
- cleaning sex toys between use and using condoms if sharing sex toys
- The benefits of routine STI screening.
- The potential complications from untreated gonorrhea.
- Co-infection risk for HIV when another STI is present.
- The asymptomatic nature of STI.
- Repeating STI screening which includes testing for Neisseria gonorrhoeae in 6 months' time as re-infection rate is high.
- The importance of revisiting a health care provider if symptoms persist.

Consultation and/or Referral

Consult with or refer to a physician or NP for all clients who are pregnant or breast-/chest-feeding. Consult with or refer to physician or NP for allergy/contraindications to treatment outlined in this DST.



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Documentation

- Neisseria gonorrhoeae is reportable
- Complete H208 form as per reporting procedures
- As per agency policy

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References

More recent editions of any of the items in the reference list may have been published since the DST was published. If you have a newer version, please use it.

Barbee, L.A., Kerani, R.P., Dombrowski, J.C., Soge, O.O. & Golden, M.R. (2013). *A* retrospective comparative study of 2-drug oral and intramuscular cephalosporin treatment regimens for pharyngeal gonorrhea. *Clinical Infectious Diseases*, 56(11), pp.1539 1545.

British Columbia Centre for Disease Control (BCCDC). (2014). <u>British Columbia treatment guidelines: Sexually transmitted infections in adolescent and adults</u>. STI/HIV Prevention and Control Division, BCCDC.

BCCDC. (2009). BCCDC Communicable disease manual Chapter 2: Immunization program. Vancouver: BCCDC.

BCCDC. (2013). BCCDC's videos on injection landmarking and technique.

BCCDC Public Health Laboratory (BCCDC PHL). (2016). Laboratory trends. Vancouver, BC.

Centers for Disease Control and Prevention (CDC). (2015). Sexually transmitted diseases treatment guidelines. Atlanta, GA.

Hottes, T.S., Lester, R.T., Hoang, L., McKay, R., Gilbert, M., Patrick, D.M., Wong, T. Martin, R. & Ogilvie, G. (2013). Cephalosporin and azithromycin susceptibility in Neissieria gonorrheoeae isolates by site of infection, British Columbia, 2006 to 2011. *Sexually Transmitted Diseases*, 40(1), pp.46-51.

Lunny, C., Taylor, D., Hoang, L., Wong, T., Gilbert, M., Lester, R. & Greub, G. (2014). <u>Self-collected versus clinician-collected sampling for chlamydia and gonorrhoea screening:</u> A systemic review and meta-analysis. *PLoS ONE*, 10(7), pp.1-23.

Papp, J. et al. (2014). Recommendations for the laboratory-based detection of <u>Chlamydia trachomatis</u> and <u>Neisseria</u> gonorrhoeae. Morbidity and Mortality Weekly Report, 63(1).

Pattman, R., Snow, M., Handy, P., Sankar, K.N. & Elawad, B. (2005). *Oxford Handbook of Genitourinary Medicine, HIV, and AIDS*, 1st Edition, Copyright (c) 2005 Oxford University Press.

Public Health Agency of Canada (PHAC). (2013). <u>Gonococcal infections</u>. In: *Canadian guidelines on sexually transmitted infections*.

PHAC. (2017). 2016 Updates Summary. In: Canadian guidelines on sexually transmitted infections.

PHAC. (2017). <u>Treatment of N. gonorrhoeae in response to the discontinuation of spectinomycin: Alternative treatment guidance statement</u>. In: *Canadian guidelines on sexually transmitted infections*.

Shafer, M., Moncada, J., Boyer, C.B., Betsinger, K., Flinn, S.D. & Schachter, J. (2003). Comparing first-void urine specimens, self-collected vaginal swabs, and endocervical specimens to detect *Chlamydia trachomatis* and *Neisseria gonorrheoeae* by a nucleic acid amplification test. *Journal of Clinical Microbiology*, 41(9), pp 4395-4399.

Schacter, J., Moncada, J., Liska, S., Shayevich, C. & Klausner, J.D. (2008). Nucleic acid amplification tests in the diagnosis of chlamydial and gonococcal infections of the oropharynx and rectum in men who have sex with men. *Sexually Transmitted Diseases*, 35(7), pp.637-642.

Stewart, C.M., Shoeman, S.A., Booth, A.R., Smith, S.D., Wilcox, M.H. & Wilson, J.D. (2012). <u>Assessment of self-taken swabs</u> versus clinician taken swab cultures for diagnosing gonorrhoea in women: single centre, diagnostic accuracy study. *BMJ*, 34, e8107.